

Isabela Quintela Matos

**Solvation Free Energy Calculations of  
Asphaltene-Like Molecules Using The SAFT- $\gamma$   
Mie Force Field**

Rio de Janeiro

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Master's dissertation presented to Engenharia de Processos Químicos e Bioquímicos graduate program, Escola de Química, Universidade Federal do Rio de Janeiro, as required for obtaining a Master's degree in Chemical Engineering.

Universidade Federal do Rio de Janeiro

Escola de Química

Engenharia de Processos Químicos e Bioquímicos Graduate Program

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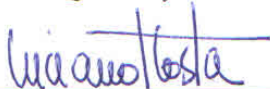
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# Abstract

Asphaltenes are a relevant fraction of crude oil to the oil industry since they can precipitate during the oil processing. Hence, we, at this dissertation, studied the solvation free energies of molecules mimicking asphaltenes in different solvents with a coarse-grained model known as SAFT- $\gamma$  Mie force field. We obtained solvation free energies by carrying out molecular dynamics simulations using the expanded ensemble method. The output of these simulations was then used to estimate the solvation free energies. For this, we employed the MBAR method. The results with solvents other than water had low absolute deviations from experimental data. In turn, hydration free energy calculations required a binary interaction parameter estimated with output data from molecular dynamics in order to obtain accurate free energy differences. These results indicated problems on the description of the water molecule by the SAFT- $\gamma$  Mie force field, but, generally, proved that this coarse-grained model could represent the solvation free energies of the studied solute-solvent pairs.

**Keywords:** Solvation free energy. Asphaltenes. SAFT- $\gamma$  Mie force field.

# Resumo

Os asfaltenos são uma fração relevante do petróleo, uma vez que podem precipitar durante o processamento e aumentar consideravelmente o custo de produção. Sendo assim, nesta dissertação, estudamos as energias livres de solvatação de moléculas do tipo asfaltenos em diferentes solventes com o campo de força coarse-grained SAFT- $\gamma$  Mie. Obtivemos energias livres de solvatação realizando simulações de dinâmica molecular com o método ensemble expandido. Os resultados dessas simulações foram então usados para estimar as energias livres de solvatação com o método MBAR. As energias livres de solvatação em solventes não aquosos apresentaram baixos desvios absolutos em relação aos dados experimentais. Por sua vez, os cálculos de energia livre de hidratação exigiram um parâmetro de interação binária estimado com dados obtidos com dinâmica molecular. Estes resultados indicaram problemas na descrição da molécula de água pelo campo de força SAFT- $\gamma$  Mie, mas, em geral, provaram que este modelo coarse-grained consegue representar as energias livres de solvatação dos pares de solutos-solventes estudados.

**Palavras-chave:** Energia livre de solvatação. Asfaltenos. Campo de força SAFT- $\gamma$  Mie.

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# List of symbols

$T$	Temperature
$P$	Pressure
$V$	Volume
$t$	Time
$p$	Momentum
$r$	Coordinates
$U, u$	Potential energy
$u$	Potential energy at reduced units
$m$	Mass
$v$	Velocity
$P$	Pressure
$\mathcal{H}$	Hamiltonian
$q$	Generalized coordinates
$K$	Kinetic energy
$N$	Number of atoms/molecules
$h$	Planck constant
$S$	Entropy
$\kappa_b$	Boltzmann constant
$\beta$	$1/\kappa_b T$
$\mu$	Chemical potential
$A$	Helmholtz free energy
$G$	Gibbs free energy
$f$	Free energy at reduce units

$F$	Forces
$\epsilon$	Depth of the potential well
$\sigma$	Distance corresponding to a zero intermolecular potential
$\lambda_r$	Repulsive exponent
$\lambda_a$	Attractive exponent
$x_i$	Molar fraction
$w_i$	Weight fraction
$\rho$	Density
$\lambda$	Coupling parameter
$\eta$	Arbitrary weight
$k_{ij}$	Binary interaction parameter

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# 1 Introduction

Solvation free energy calculations with molecular dynamics (MD) have a variety of applications ranging from drug design in the pharmaceutical industry to the development of separation technologies in the chemical industry. Solvation free energy is, more specifically, the difference in free energy related to the process of transferring a solute from an ideal gas phase into a liquid solution (SHIRTS *et al.*, 2003). Through the study of the solvation phenomenon, it is possible to obtain information about the behavior of the solvent in different thermodynamic conditions and the influence of the solute's molecular geometry. It is also possible to calculate other important properties with the solvation free energy, namely the activity coefficient at infinite dilution, Henry constant, and partition coefficients. Additionally, solvation free energy calculations can be part of the methodology of calculating solubility from molecular dynamics.

The solvation free energy calculations described above are intrinsically complex due to the many competing forces interfering in the behavior of the solute-solvent interaction. Also, free energy simulations are susceptible to sampling problems in low energy regions, and simulation results need to be correctly post-processed in order to yield free energy differences with small uncertainties. Another influencing factor in the output of these calculations is the choice of force field used to model the solvent and solute molecules. Force field is the name given to the group of parameters and equations used to represent the potential energy function of a system in molecular simulations. They have different levels of description, such as quantum mechanics, atomistic, and coarse-grained. The quantum mechanics approach describes the motion of electrons and requires for the solution of the Schrödinger equation during the simulation. In the atomistic description, only the atomic motions are represented, and this is done by solving Newton's equations of motion. Finally, in the coarse-grained description, atoms are grouped into pseudo-atoms or beads, and the equations of motion are solved for them.

These coarse-grained models are generally able to reproduce experimental free energy differences since the effects of reducing degrees of freedom in the entropy are counterbalanced by the reduction of enthalpic terms (KMIECIK *et al.*, 2016). This fact makes these models a viable option to decrease the computational time of solvation free energy calculations. Additionally, deficiencies in the description of small molecules by coarse-grained models can be revealed by free energy calculations (MOBLEY *et al.*, 2007; SHIRTS *et al.*, 2003). Hence, we, in this study, assess the performance and shortcomings of the SAFT- $\gamma$  Mie coarse-grained force field (AVENDAÑO *et al.*, 2011) with free energy calculations of a variety of solute-solvent pairs. We choose this coarse-grained force

field because it uses, unlike the majority of the force fields, the Mie potential (MIE, 1903) and because its method of obtaining parameters is more straightforward than those of other coarse-grained models. It was initially parameterized with pure component equilibrium and interfacial tension data (AVENDAÑO *et al.*, 2011), and this strategy has provided satisfactory results. Examples include the prediction of phase equilibrium of aromatic compounds (MÜLLER; MEJÍA, 2017), alkanes, light gases (HERDES *et al.*, 2015), and water (LOBANOVA *et al.*, 2015), thermodynamic properties of carbon dioxide and methane (AIMOLI *et al.*, 2014a), multiphase equilibrium of mixtures of water, carbon dioxide, and n-alkanes (LOBANOVA *et al.*, 2016), and water/oil interfacial tension (HERDES *et al.*, 2017).

We selected the solvents and solutes in our free energy calculations with the intention of testing the force field with standard sets used as a benchmark in solvation free energy calculations and with polycyclic aromatic substances used as models to asphaltenes. Asphaltenes are complicated to characterize by determining their composition on a molecular basis, but the literature broadly accepts that they can be described as a fraction of crude oil soluble in toluene and insoluble in n-alkanes (pentane, hexane, heptane) (SJÖBLOM *et al.*, 2003). They have motivated many studies with interest in developing models for their structure and behavior due to all the problems they can cause during their transportation and refining such as precipitation during the oil processing (SJÖBLOM *et al.*, 2015). This precipitation issue is a recurrent problem due to the growing market of the production of crude oil in deep waters, whose conditions are favorable to precipitation (BUENROSTRO-GONZALEZ *et al.*, 2004). As an example, asphaltene precipitation due to pressure drop can clog oil production equipment and cause a growth in the cost of production (JOSHI *et al.*, 2001). All these factors make the understanding of the behavior of asphaltenes in different chemical and physical environments relevant to the oil industry.

As said in the previous paragraph, asphaltene characterization still faces some issues. Hence, we choose to use polycyclic aromatic hydrocarbons (PAHs), which have well-defined characteristics, to initially test the efficiency of the SAFT- $\gamma$  Mie force field in describing the solvation phenomenon. PAHs are a group of organic compounds that have fused rings, carbon and hydrogen in their structure (RAVINDRA *et al.*, 2008), which is a characteristic shared with asphaltenes. The ones utilized in this work were phenanthrene, anthracene, and pyrene since they share similarities with asphaltenes regarding their solubility. In this context, we selected compounds that are used to characterize asphaltenes (toluene, hexane) as solvents in our free energy calculations. We also tested the anti-solvent/solvent effect of carbon dioxide due to its influence in asphaltene precipitation during the oil processing (SOROUSH *et al.*, 2014). Hence, the success of the description of solvation free energies of small asphaltene-like compounds by this force field can then open up the possibility of obtaining satisfactory results for



more complex asphaltene models with a force field with a low computational cost.

## 1.1 Objectives

The general objective of this work is to acquire reasonable estimates of solvation free energies of PAHs, which can represent asphaltenes, using molecular dynamics simulations with the SAFT- $\gamma$  Mie force field. By doing that, we intend to provide information about the solvation phenomenon of an important compound for the oil industry.

The specific objectives are to :

- Estimate the parameters of the SAFT- $\gamma$  Mie force field for phenanthrene using two available strategies for the parameterization of aromatic molecules with this force field.
- Estimate the binary interaction parameter required by the SAFT- $\gamma$  Mie force field for aqueous mixtures.
- Optimize the parameters necessary to carry out expanded ensemble simulations.
- Carry out expanded ensemble simulations to obtain the potential energies.
- Estimate solvation free energies with the Multistate Bennett Acceptance Ratio (MBAR) method using the potential energies obtained with molecular dynamics simulations.
- Calculate the partition coefficients for some systems studied here using the estimated solvation free energies.

## 2 Literature Review

### 2.1 Molecular Simulations of Asphaltene-Like Molecules

Asphaltenes, unlike other petroleum fractions, are not defined on a molecular basis. The most accepted definition is that they are a fraction of crude oil insoluble in n-alkanes (pentane, hexane, and heptane) and soluble in toluene (SJÖBLOM *et al.*, 2003). Due to uncertainties related to their structures, much work has been done to develop model compounds that have a well-defined structure and can represent asphaltenes. The two categories of models presented in the literature are the archipelago and continental models. In the archipelago model, asphaltenes consist of polyaromatic parts linked together by aliphatic or naphthenic moieties and, in the continental model, they consist of a single polyaromatic ring with linked aliphatic or naphthenic chains (MULLINS, 2010; MURGICH, 2003). The choice of the model's structure is highly essential since some structures/arrangements can cause the occurrence of high-energy regions during the simulation (LI; GREENFIELD, 2011).

In order to evaluate the strengths and shortcomings of asphaltene models, articles have been published about the calculations of their properties. There is an agreement in studies of these models with molecular simulation, such as the influence of the model in the packing tendency of the molecule (GREENFIELD, 2011). Kuznicki *et al.* (2009) utilized molecular dynamics in the study of the nanoaggregation of four types of model asphaltene molecules (continental, Violanthrone-79, anionic continental, and thiophenic anionic continental) in binary mixtures of toluene and water. The authors observed that, in thin films of toluene trapped between two aqueous phases, both interface-bound and core-bound asphaltenes have similar diffusion behavior. Headen *et al.* (2017) reported molecular dynamics simulations of four model asphaltenes, including archipelago and continental model. They alleged that there is no formation of nanoaggregates and that the distribution of asphaltene clusters is continuous for mixtures of asphaltenes in heptane.

Molecular dynamics simulations were also utilized in the study of Ervik *et al.* (2016a) to obtain the correct interfacial orientation of asphaltenes using a coarse-grained model of the interface and the asphaltene molecules. Also using a coarse-grained force field, Jover *et al.* (2015) carried out molecular simulations with a continental asphaltene model. The results reproduced experimental data of aggregation of asphaltene molecules in n-heptane and solubility in toluene. Gao *et al.* (2014) performed a molecular dynamics study with the GROMOS 45a3 force field (SCHULER *et al.*, 2001) to identify the influence of the structural characteristics of different asphaltene molecules.

Mikami *et al.* (2013) employed the archipelago model in their work to investigate the interfacial behavior of asphaltene molecules at the oil-water interface using molecular dynamics simulations with the OPLS-AA force field (JORGENSEN; TIRADO-RIVES, 1988). They found that asphaltenes are preferably distributed in the oil phase in the case of pure toluene and at the oil-water interface in the case of pure heptane. They also discovered an oscillatory behavior of asphaltene molecules at the oil-water interface when using the archipelago model. Teklebrhan *et al.* (2014) used a perylene-based model to study molecular association and interaction as well as the adsorption properties of the perylene molecule at the water/toluene or water/heptane interface.

## 2.2 Coarse-Grained (CG) Force Fields

Molecular simulations, as those described in the previous section, can be carried out at different levels of description. The detailed atomistic level or *ab initio* level is described by the laws of quantum mechanics. The system consists of a set of subatomic particles in which the Schrödinger's equation is solved for all of them. The next level is the atomistic description. In this description, the equations of classical mechanics are solved for the atoms in the system. Force fields at this level are based on van der Waals interactions, which may include neutral or Coulombic charged sites. The polarizability effect may also be accounted by some force fields. Contributions due to intramolecular interactions such as bond-stretching, angle-bending, and torsion are also considered in these kinds of force fields. When the simulation scale needs to be increased, and the atomistic simulations become too computationally expensive such as in the study of biological systems, the coarse-grained (CG) description can be an alternative to decrease the computational cost. This description considers that the system is made up of pseudo-atoms or beads representing multiple atoms or even an entire molecule.

There is an evident loss of information in grouping atoms; hence it is necessary to guarantee that the process of eliminating unnecessary or unimportant information ('coarse-graining') does not affect the system's physical behavior. Ideally, coarse-grained models need to have representability, robustness, transferability, and computational efficiency. Representability means that we can use a model at a state point other than the one in which it was parameterized. The other characteristic, robustness, is related to the model's ability to enable reliable predictions for various structural, thermodynamic, or transport properties. Finally, a transferable model is one in which the representation of atomic or chemical moieties have the same behavior in different molecules- e. g., a pseudo-atom representing  $CH_2$  should have the same characteristics both in an alkene molecule and in a polyethylene molecule (MÜLLER; JACKSON, 2014). To achieve the cited goals, coarse-grained force fields are usually developed by mapping the atomistic model to define the pseudo-atoms, which are generally formed by similar functional

groups.

The level of coarse-graining also needs to be established. The decision of the number of atoms per bead should be careful. There is a compromise between accuracy and computational efficiency since the simulations become faster when we group more atoms per bead, but the results can lose accuracy. After the mapping, CG force fields need to be parametrized. There are two different approaches, bottom-up and top-down, to link the simulations on the coarse-grained scale to another scale, schematically represented in Figure 2.2.1. The bottom-up approach uses information of a more detailed scale such as the quantum mechanics description or the atomistic description to obtain information necessary to the parametrization. This method highly depends on the quality of the more detailed model to succeed. In the top-down methodology, one obtains parameters from larger scales, namely experimental thermodynamic or transport properties.

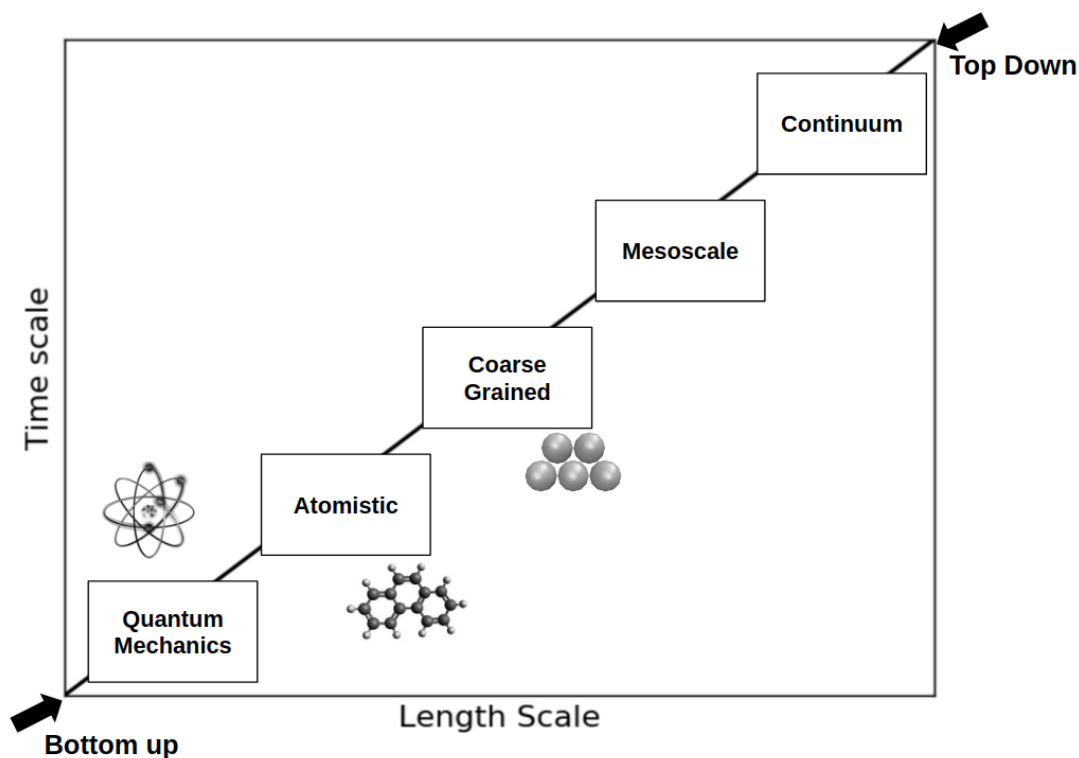


Figure 2.2.1 – Schematic representation of the two approaches of coarse-graining.

One of the first applications of coarse-grained models was in the study of protein folding (LEVITT; WARSHAW, 1975; LEVITT, 1976). These earlier protein CG models were based on known molecular structure, and they did not represent the properties of proteins correctly, but they gave an initial contribution to the knowledge of physicochemical forces associated with protein folding and protein interactions (KOGA; TAKADA, 2001). More recently, models were developed focusing on retaining the protein's chemical specificity. The Bereau and Deresno model (BEREAU; DESERNO, 2009)

represents a single amino acid residue with a maximum of four beads, and it was used in studies of protein folding and aggregation. However, this model still needs tuning to improve protein stability (BEREAU *et al.*, 2010). The OPEP (Optimized Potential for Efficient Protein Structure Prediction) model (STERPONE *et al.*, 2014; STERPONE *et al.*, 2015) represents a single amino acid with a maximum of six beads. It was used to investigate a variety of phenomena, ranging from protein folding to the modeling of DNA-RNA complexes (BARDUCCI *et al.*, 2011; CHEBARO *et al.*, 2009; STERPONE *et al.*, 2014). Other CG protein models used in the literature are the SCORPION (solvated coarse-grained protein interaction) (BASDEVANT *et al.*, 2013), the UNRES (United Residue) (ADAM *et al.*, 2014) and the MARTINI model (LARS *et al.*, 2013). The latter one is the most popular model for CG modeling of membrane proteins (MARRINK; TIELEMAN, 2013). Although these models are widely used, the existing coarse-grained force fields utilized to model proteins still need improvement, since the efficient use of coarse-grained models for proteins usually requires rigorous reconstruction of the atom-level representation, and this has only been possible for some moderate resolution coarse-grained models (KMIECIK *et al.*, 2016).

The MARTINI force field is also extensively used as a CG model for water. This force field represents four water molecules as one bead using a shifted Lennard-Jones potential. Despite its extensive use, the MARTINI water model does not correctly reproduce properties such as interfacial tension and compressibility (HE *et al.*, 2010). Besides, it can freeze at room temperature (WINGER *et al.*, 2009; MARRINK *et al.*, 2007), which requires the use of anti-freezing agents during the simulations. This behavior can be explained by the high level of coarse-graining (4:1), the lack of explicit charges, and the use of a 12-6 potential. Shinoda *et al.* (2007) used different forms of the Mie potential to build a versatile and transferable coarse-grained model for surfactant/water systems using density, interfacial tension, and hydration free energies (solvation free energies in water) data. In this model, three water molecules are represented by one pseudo-atom, and the water cross interactions are represented by a 12-4 Mie potential, and the surfactant (alkanes, oxyethylenes, ethylene glycols, ethers, and alcohols) interactions are represented by a 9-6 Mie potential. Chiu *et al.* (2010) used another type of intermolecular potential, the Morse Potential, which is softer than the LJ potential, to improve the MARTINI model.

Outside of the MARTINI framework, He *et al.* (2010) studied different levels of coarse-graining for water ranging from one to four molecules per bead using different Mie and Morse potentials. Other investigations also assessed the use of soft-core potentials to study aqueous solutions of surfactants (SHINODA *et al.*, 2007), as well as ionic liquids (BHARGAVA; KLEIN, 2009), lipids (SHINODA *et al.*, 2010), and membranes (PANTANO; KLEIN, 2009). Riniker and Gunsteren (2011) also developed a coarse-grained water model for the GROMOS force field using a systematic analysis of

different properties of small water clusters. According to their analysis, the best representation of water is achieved when one bead represents five water molecules. In this model, the potential energy is calculated using the Lennard-Jones potential, and electrostatic interactions are explicitly considered. Orsi and Essex (2011) proposed the ELBA coarse-grained model for molecular dynamics simulations of lipid membranes. In this model, electrostatics are modeled explicitly by point charges, and one water molecule is represented by a single Lennard-Jones bead embedded with a point dipole. Genheden (2016) expanded the ELBA force field to model 1-hexanol, 1-nonanol, n-hexane, and n-nonane by representing three carbons with a single bead. Another CG model for water based on the Mie Potential is the SAFT- $\gamma$  Mie force field (LOBANOVA *et al.*, 2015). In this strategy, there are two different models: CGW1-vle and CGW1-ift. Both of them represent one water molecule as one bead, and the Mie Potential has a repulsive and an attractive exponent equal to 8 and 6, respectively. The CGW1-vle model was parameterized using saturated-liquid density and vapor pressure data and should be used for simulations of aqueous systems in fluid-phase equilibrium at high temperatures and pressures. This model still suffers from premature freezing with a triple point at 343 K. The other model, CGW1-ift, was parameterized using saturated-liquid density and interfacial tension data. Hence, it is best suited for interfacial property calculations. Both models had temperature-dependent size and energy parameters and performed well for these properties over the entire liquid temperature range. The SAFT- $\gamma$  Mie force field has also been applied to other compounds with satisfactory results. Müller and Mejía (2017) parameterized the force field for aromatic compounds and tested it with simulations of fluid phase equilibrium. Herdes *et al.* (2015) carried out simulations of alkanes and light gases. Lobanova *et al.* (2016) tested the force field with binary and ternary mixtures of water and carbon dioxide. There are also articles with the SAFT- $\gamma$  Mie being used for computing thermodynamic and transport properties of carbon dioxide and methane (AIMOLI *et al.*, 2014a; AIMOLI *et al.*, 2014b) and water/oil interfacial tension (HERDES *et al.*, 2017).

## 2.3 Solvation Free Energies

Solvation free energy calculations with molecular dynamics can be used to evaluate the quality of a coarse-grained force field, such as the models described in the preceding section, since these estimations can reveal deficiencies in a force field. Besides this application, solvation free energies are used to obtain information about the behavior of the solvent in different thermodynamic conditions and to assess the influence of the solute's molecular geometry on the solvation phenomenon. Due to their range of application and inherent complexity, free energy calculations were the subject of a variety of studies in the last decades interested in improving the free energy

simulations and their post-processing methods (SHIRTS; CHODERA, 2008; PALIWAL; SHIRTS, 2011; SHIRTS; PANDE, 2005; YTREBERG *et al.*, 2006).

Recent articles (MOBLEY; GUTHRIE, 2014; MATOS *et al.*, 2017) made available a large database of hydration free energies of small molecules using the GAFF force field for the solutes and the TIP3P model for water. Beckstein *et al.* (2014) also calculated the hydration free energies for fifty-two compounds with the OPLS-AA force field. They obtained an overall root mean square deviation from the experimental data of 1.75 kcal/mol and concluded that the reproducibility of the Lennard-Jones parameters is the main constraint of the precision of their results. Izairi and Kamberaj (2017) also studied hydration free energies but with the intention of comparing the polar and nonpolar contributions. Garrido *et al.* (2009, 2011) calculated the free energy of solvation of large alkanes in 1-octanol and water with three different force fields (TraPPE, GROMOS, and OPLS-AA/TraPPE). These authors also estimated the solvation free energy of propane and benzene in non-aqueous solvents like n-hexadecane, n-hexane, ethylbenzene, and acetone with the united atoms TraPPE force field (TraPPE-UA) and the all-atoms TraPPE force field (TraPPE-AA). Roy *et al.* (2017) addressed the choice of the Lennard-Jones parameters for predicting solvation free energy of different solutes in 1-octanol. They calculated the solvation free energy of a set of 205 small organic molecules in 1-octanol and found that the force field parametrization of n-octanol proposed by Kobryn and Kovalenko (2008) provided the best agreement to the experimental data. Gonçalves and Stassen (2005) calculated the free energy of solvation using the polarizable continuum model coupled to molecular dynamics simulation with the GROMOS force field. These calculations were done with a representative set of solutes and with the solvents tetrachloride, chloroform, and benzene. Using the GAFF and the polarizable AMOEBA force fields, Mohamed *et al.* (2016) evaluated the solvation free energy of small molecules in toluene, chloroform, and acetonitrile, and obtained a mean unsigned error of 1.22 kcal/mol for AMOEBA and 0.66 kcal/mol for GAFF. To define the role of water as the solvent in the docking structure determination of proteins, Matubayasi (2017) developed a method to compute the solvation free energy of proteins while using OPLS-AA force field for the solutes and the TIP3P model for water. Genheden (2016) expanded the ELBA force field to calculate solvation free energies of more than 150 solutes taken from the Minnesota solvation database in polar (water, hexanol, octanol, and nonanol) and apolar (hexane, octane, and nonane) solvents. He obtained mean absolute deviations of 1 kcal/mol for water and 1.5 kcal/mol for hexane. In this model, three carbons are represented by a single bead and water is also represented by a single bead.

Despite the variety of data using the intramolecular Lennard-Jones potential, there are not many studies with the Mie Potential in free energy calculations. That is why we, in the present study, try to provide information about these predictions



with the SAFT- $\gamma$  Mie coarse-grained force field. As said before, the output of these calculations are highly dependent on the force field, and some coarse-grained models produced satisfactory results for these simulations. In addition, the use of a coarse-grained model can decrease the simulation time, as explained in Section 2.2. Therefore, the consideration of other coarse-grained approaches with similar performances to the all-atoms force fields can help increase the scale of solvation free energy calculations.

## 2.4 Solvation Free Energy Calculation Methods

Solvation free energy calculations account for the difference in free energy related to transferring the solute from the ideal gas phase to the liquid solvent phase. To do that, we gradually insert the solute in the solvent. This process is mathematically carried out by using a coupling parameter ( $\lambda$ ) on the total potential energy function, where  $\lambda$  represents the intermediate states in the transition from the solute in the ideal gas phase ( $\lambda = 0$ ) to the solute entirely inserted in the solvent phase ( $\lambda = 1$ ). Hence, during the solvation free energy simulations, we obtain total potential energies corresponding to these coupling parameters [ $U(\lambda)$ ]. After the simulations, these potential energies need to be post-processed and analyzed so as to calculate the solvation free energies efficiently. Therefore, in this section, we present a quick review of the methods available in the literature to perform the estimation of the solvation free energies. Since these calculations can have slow convergence, many scientific groups in the last decades focused on developing analysis methods to estimate these free energies. Almost all of these methods rely on the following approaches: thermodynamic integration, histograms, and free energy perturbation (FEP) based methods.

### 2.4.1 Thermodynamic integration

The thermodynamic integration method (KIRKWOOD, 1935) uses equilibrium averages to evaluate the energy derivative with respect to the coupling parameter ( $\lambda$ ):

$$\frac{\partial(G/ \kappa_b T)}{\partial \lambda} = \left\langle \frac{\partial \mathcal{H}}{\partial \lambda} \right\rangle_{N,P,T}. \quad (2.1)$$

In Eq. (2.1),  $\kappa_b$  is the Boltzmann constant,  $G$  is the Gibbs free energy and  $\mathcal{H}$  is the Hamiltonian of the system. The derivative is obtained through an analytical expression for every configuration data among the states retrieved from the molecular dynamics simulations. Some examples of methods for obtaining these expressions are the trapezoidal rule or natural cubic spline (PALIWAL; SHIRTS, 2011). There are also more complex schemes that are usually system specific, such as those found in Jorge *et al.* (2010) and Shyu and Ytreberg (2010). With the MD simulations for each coupling



parameter  $\lambda_k$  carried out, the average over the derivative at each state is computed and the free energy can be estimated upon integration over all  $\lambda$ -points:

$$\Delta G \approx \int_0^1 \left\langle \frac{\partial \mathcal{H}}{\partial \lambda_k} \right\rangle d\lambda. \quad (2.2)$$

The thermodynamic integration method has as a disadvantage its sensibility to the choice of intermediate states and the necessity of a higher number of intermediate states when compared to the other methods discussed here (RUITER *et al.*, 2013).

## 2.4.2 Histograms

Histograms are used to compute probability distributions. Usually, every histogram count is treated as the number of visits to a specific state. The standard practice when using histograms is to use the weighted histogram analysis method (WHAM) developed by Ferrenberg and Swendsen (1989) and generalized by Kumar *et al.* (1992) (CHIPOT; POHORILLE, 2007). It puts together different histograms by minimizing the statistical error in the computed density of states and entropy function. This method describes the total probability distribution as a weighted unbiased sum of probability distributions from biased simulations. It was developed to avoid problems related to data loss and high uncertainties (ROUX, 1995). The probability distribution dependent on the potential energy ( $U$ ) and temperature ( $T$ ) for the WHAM is

$$\begin{aligned} \tilde{\varrho}_r^*(U, T) &= \frac{\sum_i f_i(U) \exp(-\beta U)}{\sum_i f_{tot,i} \exp(\beta_i \tilde{A}_i - \beta_i U)}, \\ \exp(-\beta_i \tilde{A}_i) &= \sum_U \tilde{\varrho}_r^*(U, T), \text{ and} \\ \tilde{\varrho}_r(U, T) &= \frac{\tilde{\varrho}_r^*(U, T)}{\sum_U \tilde{\varrho}_r^*(U, T)}, \end{aligned} \quad (2.3)$$

where  $\beta = 1/\kappa_b T$ ,  $\tilde{A}_i$  gives the free energy for run  $i$ ,  $f_i(U)$  is the number of counts of energy  $U$  for run  $i$ , and  $f_{tot,i}$  is the total number of counts in run  $i$ . Eq. (2.3) is solved self consistently with the initial value for  $\tilde{A}_i$  equals to zero. The final unnormalized probability distribution is then given by  $\tilde{\varrho}_r(U, T)$ .

## 2.4.3 Free Energy Perturbation (FEP)

The free energy perturbation method (ZWANZIG, 1954) is one of the most general-purpose strategies to calculate free energy differences. In this method, the thermodynamics of two different systems (A and B) are related to the intention of evaluating differences in intermolecular potentials. This energy change from state A to

state B is calculated by

$$\Delta G_{AB} = -\kappa_b T \ln \langle e^{-\beta(U_B - U_A)} \rangle_A. \quad (2.4)$$

According to the equation above, the free energy difference is calculated by doing an average over the configurations of state A and B obtained during the simulation of state A. This method requires a great overlap between the configurations of the states in order to obtain a rapid convergence of the free energy difference. Overlap between configurations A and B means that the configurations of state B represent only a small perturbation in the configurations of state A. To guarantee overlap, it is possible to carry out simulations in N intermediate states between A and B, so Eq. (2.4) becomes:

$$\Delta G_{AB} = -\kappa_b T \sum_{i=0}^N \ln \langle e^{-\beta(U_{i+1} - U_i)} \rangle_i. \quad (2.5)$$

This way of calculation  $\Delta G$  is also called Exponential Averaging (EXP) (ZWANZIG, 1955; PALIWAL; SHIRTS, 2011). The direction of the transformation is crucial in this method. If the direction is of decreasing entropy, the step is of insertion ( $\Delta G_{AB}$ ), and the method is called insertion exponential averaging (IEXP). When the direction is of increasing entropy, the step is of deletion ( $\Delta G_{BA}$ ), and the method is labeled as deletion exponential averaging (DEXP) (PALIWAL; SHIRTS, 2011). These directions can yield different values of free energy differences due to undersampling in the tail regions of the  $\Delta G_{AB}$  distribution (KLIMOVICH *et al.*, 2015; POHORILLE *et al.*, 2010). These problems make the EXP methods not suited to calculate free energy differences when the system does not have sufficient overlap. For these cases, the Bennett Acceptance Ratio or the Multi-State Bennett Acceptance Ratio are more adequate.

#### 2.4.3.1 Bennett Acceptance Ratio (BAR)

The BAR method (BENNETT, 1976) was developed with the intent of eliminating the directional bias in the free energy estimation with FEP. It uses the uncorrelated samples of the potential energy in both directions ( $A \rightarrow B$  and  $B \rightarrow A$ ) to obtain the free energy differences utilizing the information in a statically optimal way. The free energy difference between two intermediate states is calculated using the potential energy difference ( $\Delta U$ ) between states i and j. The calculation is done by solving self-

consistently the following equations:

$$\Delta G_{ij} = \frac{1}{\beta} \ln \left( \frac{\sum_{k=1}^{N_j} \frac{1}{1 + \exp[-\beta(\Delta U_k^j + C)]}}{\sum_{l=1}^{N_i} \frac{1}{1 + \exp[-\beta(\Delta U_l^i - C)]}} \right) + C - \frac{1}{\beta} \ln \left( \frac{N_j}{N_i} \right), \quad (2.6)$$

$$C = \Delta G_{ij} + \frac{1}{\beta} \ln \left( \frac{N_j}{N_i} \right). \quad (2.7)$$

The total free energy difference between the end states is then given by the sum over differences of consecutive intermediate states. This method also provides a function to obtain the variance for the free energy differences, which is a minimum. The variance equation for any value of  $C$  is given by:

$$s_{ij}^2 = \frac{1}{\beta^2 N_i} \left[ \frac{\langle f^2(x) \rangle_i}{\langle f(x) \rangle_i^2} - 1 \right] + \frac{1}{\beta^2 N_j} \left[ \frac{\langle f^2(x) \rangle_j}{\langle f(x) \rangle_j^2} - 1 \right], \quad (2.8)$$

where  $f(x) = 1/(1+x)$  is the Fermi function and  $x = \exp[\beta(\Delta U - C)]$ . The variance of the free energy difference between end states can be calculated by assuming independent errors and summing over the variances of consecutive intermediate states. However, this assumption is not correct and there is no general formula to obtain a statistically unbiased estimate of an entire transformation with the BAR method (PALIWAL; SHIRTS, 2011).

There are two other methods related to the BAR method that do not solve Eqs. (2.6) and (2.7) self consistently. By doing that, free energy differences will not have minimum variance and the averages of Eqs. (2.6) - (2.8) are accumulated (PALIWAL; SHIRTS, 2011). The two methods are the Unoptimized Bennett Acceptance Ratio (UBAR) and the Range-Based Bennett Acceptance Ratio (RBAR). The first one avoids the self consistent solution of the BAR equations by defining  $C = \beta^{-1} \ln(N_j/N_i)$ . The UBAR method will be close to optimal when each intermediate free energy is relatively near zero. In turn, the RBAR method selects a range of initial guesses of the constant  $C$  to calculate a range of  $\Delta G_{ij}$ . The value of free energy difference corresponding to the minimum variance is then used as input in Eq. (2.7) to calculate the value of  $C$ . Hence, this method requires a good estimation of the initial range of the values of  $C$ . The RBAR can be advantageous when compared to BAR since only the accumulated averages need to be retained for postprocessing (PALIWAL; SHIRTS, 2011).

### 2.4.3.2 Multistate Bennett Acceptance Ratio (MBAR)

The MBAR method (SHIRTS; CHODERA, 2008) is a further development of the BAR method, and is the one chosen to estimate the solvation free energies in this dissertation. This method consists of an estimator that computes free energies and their uncertainties of each  $K$  state by minimizing the  $K \times K$  matrix of variances simultaneously. The estimator solves the following equation for each  $G_i$  self consistently:

$$G_i = -\frac{1}{\beta} \ln \sum_{k=1}^K \sum_{n=1}^{N_k} \frac{\exp[-\beta U_i(x_{kn})]}{\sum_{l=1}^K N_l \exp[\beta(G_l - U_l(x_{kn}))]}. \quad (2.9)$$

The equation above requires the evaluation of the potential energy  $[U_i(x_{kn})]$  of the  $n_{th}$  uncorrelated configuration obtained at state  $K$  and all uncorrelated configuration snapshots ( $N_k$ ) from state  $K$ . Free energy changes between states are given then by  $\Delta G_{ij} = G_j - G_i$ . Much like the majority of the other methods exposed in this section, the MBAR provides a statistical estimator for the free energies and their variances. Hence, the reason we choose MBAR to estimate the free energy differences is that it provides an estimator with lowest variances than the other methods and it is perhaps the most consistently well-performing free energy estimator (PALIWAL; SHIRTS, 2011). Since this is the chosen method, further explanation of MBAR is available in Section 3.5 of the Computational Methods Chapter.

# 3 Computational Methods

The primary goal of this study was to obtain solvation free energy estimates of asphaltene-like molecules using molecular dynamics. In order to do that, we used a variety of computational methods including the molecular dynamics itself, the parameterization of the SAFT- $\gamma$  Mie force field, the solvation free energy simulations, the expanded ensemble method, and the MBAR method, briefly explained in the preceding chapter. The aforementioned methods are then presented and explained here.

## 3.1 Molecular Dynamics

### 3.1.1 Background and Formulation

Molecular Dynamics (MD) uses molecular configurations (Cartesian coordinates and momentum) to extract structural, thermodynamic, and dynamic information of a system. This information is extracted from the time evolution of the system, which is obtained through the numerical integration of the Newton's equations of motion (TUCKERMAN, 2010):

$$\frac{d\vec{p}_i}{dt} = -\frac{\partial U(\vec{r}_N)}{\partial \vec{r}_i}, \quad (3.1)$$

where  $p_i$  is the momentum and  $r_N$  are the coordinates of all the atoms ( $x_1, y_1, z_1, \dots, x_N, y_N, z_N$ ). Alternatively, we can write the equation relative to the velocity ( $v_i$ ):

$$m_i \frac{d\vec{v}_i}{dt} = -\frac{\partial U(\vec{r}_N)}{\partial \vec{r}_i}. \quad (3.2)$$

In order to develop equations for any coordinate system, for instance  $q^N = (r_1, \theta_1, \phi_1)$ , the Hamiltonian formulation, a more general formulation of classical mechanics, is used to develop the equations of motion:

$$\mathcal{H}(q^N, p^N) = K(p^N) + U(q^N). \quad (3.3)$$

In the equation above,  $K$  is the kinetic energy and  $U$  is the potential energy. The equations of motion are then rewritten using the Hamiltonian:

$$\frac{d\vec{p}_i}{dt} = -\frac{\partial \mathcal{H}}{\partial \vec{q}_i}, \quad (3.4)$$

$$\frac{d\vec{q}_i}{dt} = \frac{\partial \mathcal{H}}{\partial \vec{p}_i}. \quad (3.5)$$

In the dynamics described by the equations above, the Hamiltonian is preserved. The coordinate and momentum axes for each atom in a  $6N$  dimensional space is defined as the phase space. The trajectory through the phase space is then the time evolution of a system in a molecular dynamics simulation. This evolution of the simulation may be used to calculate the thermodynamic properties if the system is ergodic, that is, a trajectory in this system will explore with the same probability all regions of the phase space of microstates (points in phase space) with the same energy (SHELL, 2015).

### 3.1.2 Statistical Ensembles

In order to calculate thermodynamic properties, we need to define the control variables of a system. For an isolated system at equilibrium, the control variables are the number of particles ( $N$ ), volume ( $V$ ), and total energy ( $E$ ). The set of configurations under the control variables is then called statistical ensemble. In the example cited above, it is specifically named the microcanonical ensemble. Following the ergodic hypothesis, the system at these conditions will spend the same amount of time in each of the microstates with the fixed Hamiltonian. The number of accessible microstates is defined by the partition function or the density of states, and is given by the following equation for the microcanonical ensemble:

$$\Omega(N, V, E) = \frac{\epsilon_0}{h^{3N} N!} \int dp^N dr^N \delta[\mathcal{H}(p^N, r^N) - E]. \quad (3.6)$$

Here,  $\epsilon_0$  is the energy unit,  $h$  is the Planck constant, and  $\delta$  is the Dirac delta function. As mentioned above, the system will spend the same amount of time at each of the microstates, i. e. each of these microstates have equal probabilities ( $\varrho$ ) of being visited. Such probability is:

$$\varrho(p^N, r^N) = \frac{[\mathcal{H}(p^N, r^N) - E]}{\Omega(N, V, E)}. \quad (3.7)$$

The macroscopic properties from molecular dynamics are then obtained from the relation of the microcanonical partition function to the entropy ( $S$ ). Known as the Boltzmann equation. It is:

$$S = \kappa_b \ln[\Omega(N, V, E)], \quad (3.8)$$

where  $\kappa_b$  is the Boltzmann constant. With these equations, we can derive other relations

to macroscopic properties with the fundamental thermodynamic equations:

$$dS = \frac{1}{T}dE + \frac{P}{T}dV - \frac{\mu}{T}dN, \quad (3.9)$$

$$dE = TdS - PdV + \mu dN. \quad (3.10)$$

As said above, the microcanonical ensemble has  $N$ ,  $V$ , and  $E$  as its control variables. Other ensembles can also be defined according to the macroscopic properties held constant. In the canonical ensemble,  $N$ ,  $V$ , and the temperature ( $T$ ) are held constant and  $N$ , pressure ( $P$ ), and  $T$  are held constant in the isothermal-isobaric ensemble. Other ensembles are the isoenthalpic-isobaric (constant number of particles, pressure, and enthalpy) and the grand canonical (constant chemical potential, volume, and temperature) ones. A variety of physical properties is measured at the conditions of the isothermal-isobaric ensemble such as enthalpies, entropies, redox potential, equilibrium constants, and free energies, what makes this ensemble one of the most important (TUCKERMAN, 2010). This is also the ensemble in which solvation free energy simulations are carried out in this work, hence we are going to briefly describe it. This ensemble is obtained from a Legendre transformation on the canonical ensemble. The Helmholtz free energy  $A(N, V, T)$  becomes the Gibbs free energy  $G(N, P, T)$  by transforming the volume into the external pressure:

$$G(N, P, T) = A(N, V, T) + PV, \quad (3.11)$$

where  $V = V(P)$ . The Gibbs free energy is related to its partition function  $\Delta(N, P, T)$  by:

$$G(N, P, T) = -\kappa_b T \ln \Delta(N, P, T), \quad (3.12)$$

where  $\Delta(N, P, T)$  is given by:

$$\Delta(N, P, T) = \frac{1}{V_0 N! h^{3N}} \int_0^\infty dV \int dp^N dr^N \exp \left[ -\beta \left( \mathcal{H}(r^N, p^N) + PV(r^N) \right) \right]. \quad (3.13)$$

The Helmholtz free energy is then given by:

$$A(N, V, T) = -\kappa_b T \ln Q(N, V, T). \quad (3.14)$$

In the equation above,  $Q(N, V, T)$  is the partition function of the canonical en-

semble:

$$Q(N, V, T) = \frac{1}{h^{3N} N!} \int dp^N dr^N \exp [-\beta \mathcal{H}(r^N, p^N)]. \quad (3.15)$$

From these relations and a differential change in  $G$ , we can obtain the chemical potential ( $\mu$ ), the average volume ( $\langle V \rangle$ ), and entropy relations for isothermal-isobaric ensemble:

$$\mu = \left( \frac{\partial G}{\partial N} \right)_{P, T} = -\kappa_b T \left( \frac{\partial \ln \Delta(N, P, T)}{\partial N} \right)_{P, T}, \quad (3.16)$$

$$\langle V \rangle = \left( \frac{\partial G}{\partial P} \right)_{N, T} = -\kappa_b T \left( \frac{\partial \ln \Delta(N, P, T)}{\partial P} \right)_{N, T}, \quad (3.17)$$

$$S = - \left( \frac{\partial G}{\partial T} \right)_{N, P} = \kappa_b \left[ \ln \Delta(N, P, T) + T \left( \frac{\partial \ln \Delta(N, P, T)}{\partial T} \right)_{N, P} \right]. \quad (3.18)$$

### 3.1.3 Thermostats and Barostats

The isothermal-isobaric and canonical ensembles have external conditions being applied to it (temperature and pressure). For temperature control, the method employed mimics the effect of a thermal reservoir through the use of a thermostat. The thermostats need to be capable of capturing the correct energy fluctuations in the system since the kinetic energy will fluctuate when using a heat bath to control the temperature in a canonical ensemble of a finite system (FRENKEL; SMIT, 2001).

Among the available thermostat are the Berendsen (BERENDSEN *et al.*, 1984), the Andersen (ANDERSEN, 1980), and the Nosé (NOSÉ, 1984) thermostats, but, here, we are going to discuss the most widely used thermostat: the Nosé-Hoover (HOOVER, 1985). This thermostat is based on the formulation of Nosé (NOSÉ, 1984), who used a Lagrangian that contains additional and artificial coordinates and velocities (FRENKEL; SMIT, 2001). In this method, the Hamiltonian in a canonical ensemble is constructed as:

$$\mathcal{H}_{Nosé} = K(p^N) + U(q^N) + \frac{\xi^2 Q}{2} + 3N \kappa_b T \ln s, \quad (3.19)$$

where  $\xi$  is a friction coefficient related to the conjugate momentum of the thermal reservoir to which the system is coupled,  $s$  is the position of the thermal reservoir, and  $Q$  is a parameter that determines the time scale. The velocity update is then done with the friction term added to the equations of motion (SHELL, 2015):

$$\frac{dr_i}{dt} = \frac{p_i}{m_i}, \quad (3.20)$$



$$\frac{dp_i}{dt} = -\frac{\partial U(r^N)}{\partial r_i} - \xi p_i, \quad (3.21)$$

$$\frac{d\xi}{dt} = \frac{\sum p_i^2/m_i - 3N\kappa_b T}{Q}, \quad (3.22)$$

$$\frac{1}{s} \frac{ds}{dt} = \frac{d \ln s}{dt} = \xi. \quad (3.23)$$

This approach proposed by the Nosé-Hoover thermostat only generates a correct canonical distribution for molecular systems in which there are no external forces ( $F_i$ ) and the center of mass is fixed or if there is only one conserved quantity (FRENKEL; SMIT, 2001). To diminish these restrictions of the Nosé-Hoover thermostat, the Nosé-Hoover chains of thermostats method was developed by Martyna *et al.* (1992). This method is the one chosen to control the temperature in our simulations. It proposes the use of another thermal reservoir or a whole chain of thermal reservoirs in order to enhance temperature equilibration (SHELL, 2015). Here, we are going to present its equations of motion since this was the method used in this dissertation. The equations of motion of a system of N particles coupled with M Nosé-Hoover chains is given by

$$\frac{dr_i}{dt} = \frac{p_i}{m_i}, \quad (3.24)$$

$$\frac{dp_i}{dt} = F_i - \frac{p_{\xi_1}}{Q_1} p_i, \quad (3.25)$$

$$\frac{d\xi_k}{dt} = \frac{p_{\xi_k}}{Q_k} \quad k = 1, \dots, M, \quad (3.26)$$

$$\frac{dp_{\xi_1}}{dt} = \sum p_i^2/m_i - 3N\kappa_b T - \frac{p_{\xi_2}}{Q_2} p_{\xi_1}, \quad (3.27)$$

$$\frac{dp_{\xi_k}}{dt} = \frac{p_{\xi_{k-1}}^2}{Q_{k-1}} - \kappa_b T - \frac{p_{\xi_{k+1}}}{Q_{k+1}} p_{\xi_k}, \quad (3.28)$$

$$\frac{dp_{\xi_M}}{dt} = \frac{p_{\xi_{M-1}}^2}{Q_{M-1}} - \kappa_b T. \quad (3.29)$$

The conserved energy for the Nosé Hoover chains is then equal to

$$\mathcal{H}_{Nosé\ Chains} = K(p^N) + U(q^N) + \sum_{k=1}^M \frac{p_{\xi_k}^2}{2Q_k} + 3N\kappa_b T\xi_1 + \sum_{k=2}^M \kappa_b T\xi_k. \quad (3.30)$$

The pressure is controlled with a barostat. It maintains the pressure constant during the simulation by adjusting the simulation volume. Among the available barostats methodologies are the Berendsen (BERENDSEN *et al.*, 1984), in which the pressure is coupled to a pressure bath, and the volume is periodically rescaled, and the Andersen barostat (ANDERSEN, 1980), which serves as a basis for other barostating methods such as the ones developed by Hoover (HOOVER, 1985), Martina-Tobias-Klein (MARTYNA *et al.*, 1994) and Parrinello-Rahman (PARRINELLO; RAHMAN, 1981). The Andersen's idea was to couple the system to a fictional pressure bath and incorporate the volume into the phase space as an additional degree of freedom (TUCKERMAN, 2010). Here, we are going to present the approach developed by Martyna *et al.* (1994) since this was the one used to control the pressure in our simulations. The equations of motion for a chain of barostats of length M are given by

$$\frac{dr_i}{dt} = \frac{p_i}{m_i} + \frac{p_\epsilon}{\mathcal{W}} r_i, \quad (3.31)$$

$$\frac{dp_i}{dt} = F_i - \left(1 + \frac{d}{dN}\right) \frac{p_\epsilon}{\mathcal{W}} p_i - \frac{p_{\epsilon_1}}{Q_1} p_i, \quad (3.32)$$

$$\frac{dV}{dt} = \frac{dV p_\epsilon}{\mathcal{W}} \quad (3.33)$$

$$\frac{dp_\epsilon}{dt} = dV(P_{int} - P_{ext}) + \frac{1}{N} \sum_{i=1}^N \frac{p_i^2}{m_i} - \frac{p_{\xi_1}}{Q_1} p_\epsilon, \quad (3.34)$$

$$\frac{d\xi_k}{dt} = \frac{p_{\xi_k}}{Q_k} \quad k = 1, \dots, M, \quad (3.35)$$

$$\frac{dp_{\xi_1}}{dt} = \sum \frac{p_i^2}{m_i} + \frac{p_\epsilon^2}{\mathcal{W}} - (dN + 1)\kappa_b T - \frac{p_{\xi_2}}{Q_2} p_{\xi_1}, \quad (3.36)$$

$$\frac{dp_{\xi_k}}{dt} = \frac{p_{\xi_{k-1}}^2}{Q_{k-1}} - \kappa_b T - \frac{p_{\xi_{k+1}}}{Q_{k+1}} p_{\xi_k} \quad k = 2, \dots, M - 1, \quad (3.37)$$

$$\frac{dp_{\xi_M}}{dt} = \frac{p_{\xi_{M-1}}^2}{\mathcal{Q}_{M-1}} - \kappa_b T. \quad (3.38)$$

In the equations above,  $\epsilon$  is equal to

$$\epsilon = \ln \left[ \frac{V}{V(0)} \right], \quad (3.39)$$

where  $V$  is the volume of the system and  $V(0)$  is the volume at  $t = 0$ . The parameter ( $\mathcal{W}$ ) associated to  $\epsilon$ , the momentum ( $p_\epsilon$ ) conjugate to  $\epsilon$ , the external pressure ( $P_{ext}$ ), and the internal pressure ( $P_{int}$ ) are also featured in the equations of motion.  $P_{ext}$  is imposed as we do with the temperature in the thermostat and  $P_{int}$  is calculated during the simulation with the following equation (TUCKERMAN, 2010):

$$P_{int} = \frac{1}{dV} \left[ \sum_{i=1}^N \left( \frac{p_i^2}{m_i} + r_i \cdot F_i \right) - dV \frac{\partial U}{\partial V} \right]. \quad (3.40)$$

Finally, the conserved energy for the chain of barostats proposed by Martyna *et al.* (1994) is equal to

$$\mathcal{H}_{N,P_{ext},T} = K(p^N) + U(q^N) + \frac{p_\epsilon^2}{\mathcal{W}} + \sum_{k=1}^M \frac{p_{\xi_k}^2}{\mathcal{Q}_k} + (dN + 1)\kappa_b T \xi_1 + \kappa_b T \sum_{k=1}^M \xi_k + P_{ext} V. \quad (3.41)$$

All the equations of motion above are then numerically integrated using the methodologies described in the next section.

### 3.1.4 Integration of the equations of motion

With the formalism defined for the equations of motions and the statistical ensemble, we can now derive discrete-time numerical approximations for them. The basic idea is to solve the trajectory of atoms as a function of time [ $r^N(t)$ ] by updating the positions at discrete time intervals or time steps. To do that, the classical time evolution approach is used to preserve the Hamiltonian of the system in the numerical integration methods. In this approach, we consider the time evolution of an arbitrary function  $a(x_t)$  along a trajectory  $x_t$  (TUCKERMAN, 2010). Doing the time derivative of  $a(x_t)$ :

$$\frac{da}{dt} = \sum_{\alpha=1}^{3N} \left[ \frac{\partial \mathcal{H}}{\partial p_\alpha} \frac{\partial a}{\partial q_\alpha} - \frac{\partial \mathcal{H}}{\partial q_\alpha} \frac{\partial a}{\partial p_\alpha} \right]. \quad (3.42)$$

In the equation above, we can represent the time evolution of  $a(x_t)$  with the

Poisson bracket:

$$\frac{da}{dt} = \{a, \mathcal{H}\}. \quad (3.43)$$

The Poisson bracket is equal to applying the Liouville operator ( $i\mathcal{L}$ ) on the phase space. Hence,

$$\frac{da}{dt} = i\mathcal{L}a. \quad (3.44)$$

Substituting Eq. 3.44 in Eq. 3.42, we have the following equation:

$$i\mathcal{L}a = \sum_{\alpha=1}^{3N} \left[ \frac{\partial \mathcal{H}}{\partial p_{\alpha}} \frac{\partial a}{\partial q_{\alpha}} - \frac{\partial \mathcal{H}}{\partial q_{\alpha}} \frac{\partial a}{\partial p_{\alpha}} \right]. \quad (3.45)$$

The solution of Eq. 3.44 for  $a(x_t)$  is given by

$$a(x_t) = \exp(i\mathcal{L}t)a(x_0). \quad (3.46)$$

Here,  $\exp(i\mathcal{L}t)$  is known as the classical propagator. The effect of this operator in a function can not be evaluated. However, we can develop approximate solutions for the Hamiltonian's equations with this operator (TUCKERMAN, 2010). Rewriting Eq. 3.45 as

$$i\mathcal{L} = i\mathcal{L}_1 + i\mathcal{L}_2, \quad (3.47)$$

where

$$\begin{aligned} i\mathcal{L}_1 &= \sum_{\alpha=1}^N \frac{\partial \mathcal{H}}{\partial p_{\alpha}} \frac{\partial}{\partial q_{\alpha}}, \\ i\mathcal{L}_2 &= - \sum_{\alpha=1}^N \frac{\partial \mathcal{H}}{\partial q_{\alpha}} \frac{\partial}{\partial p_{\alpha}}. \end{aligned} \quad (3.48)$$

The operators  $i\mathcal{L}_1$  and  $i\mathcal{L}_2$  in the equations above are non-commuting operators, that is, the order in which the operator is applied is important (TUCKERMAN, 2010). This fact implies that we can not separate the classical propagator  $\exp(i\mathcal{L}t)$  into the product  $\exp(i\mathcal{L}_1t) \exp(i\mathcal{L}_2t)$ . Though we can not do that, we can still express the propagator in terms of these two factors by using the symmetric Trotter theorem or Strang splitting formula (TROTTER, 1959; STRANG, 1968). Applying this theorem to

the classical propagator, we then obtain

$$\begin{aligned} \exp(i\mathcal{L}t) &= \exp(i\mathcal{L}_1t + i\mathcal{L}_2t) = \\ \lim_{P \rightarrow \infty} [\exp(i\mathcal{L}_2t/2P) \exp(i\mathcal{L}_1t/P) \exp(i\mathcal{L}_2t/2P)]^P, \end{aligned} \quad (3.49)$$

where  $P$  is an integer. Defining a time step  $\Delta t = t/P$  and using it in Eq. 3.49, we have

$$\exp(i\mathcal{L}t) = \lim_{P \rightarrow \infty, \Delta t \rightarrow 0} [\exp(i\mathcal{L}_2\Delta t/2) \exp(i\mathcal{L}_1\Delta t) \exp(i\mathcal{L}_2\Delta t/2)]^P. \quad (3.50)$$

In order to obtain an approximation for  $\exp(i\mathcal{L}t)$ , we do not take the limits and consider that  $P$  is a finite number (TUCKERMAN, 2010). The resulting approximation for the classical propagator is then

$$\exp(i\mathcal{L}t) \approx [\exp(i\mathcal{L}_2\Delta t/2) \exp(i\mathcal{L}_1\Delta t) \exp(i\mathcal{L}_2\Delta t/2)]^P + \vartheta(P\Delta t^3), \quad (3.51)$$

using  $P = t/\Delta t$

$$\exp(i\mathcal{L}\Delta t) \approx \exp(i\mathcal{L}_2\Delta t/2) \exp(i\mathcal{L}_1\Delta t) \exp(i\mathcal{L}_2\Delta t/2) + \vartheta(\Delta t^3). \quad (3.52)$$

Now we can use Eq. 3.52 as a numerical propagation for a single time step ( $\Delta t$ ). Using this propagation on a single particle moving with Hamiltonian, where  $i\mathcal{L}_1 = K(p)$  and  $i\mathcal{L}_2 = U(r)$ , we obtain

$$\exp(i\mathcal{L}\Delta t) \approx \exp\left(-\frac{\Delta t}{2} \frac{\partial U}{\partial r} \frac{\partial}{\partial p}\right) \exp\left(\Delta t \frac{p}{m} \frac{\partial}{\partial r}\right) \exp\left(-\frac{\Delta t}{2} \frac{\partial U}{\partial r} \frac{\partial}{\partial p}\right), \quad (3.53)$$

where the derivatives of the intermolecular potential  $-\frac{\partial U(r)}{\partial r}$  are equal to the force ( $F$ ) acting on the particle. We now are able to replace the exact solution of Eq. 3.46 with the approximation of Eq. 3.53. The approximation evolution from a initial condition  $(r(t), p(t))$  is then

$$\begin{aligned} \begin{bmatrix} r(t + \Delta t) \\ p(t + \Delta t) \end{bmatrix} &\approx \exp\left(\frac{\Delta t}{2} F(r(t)) \frac{\partial}{\partial p}\right) \times \exp\left(\Delta t \frac{p(t)}{m} \frac{\partial}{\partial r}\right) \\ &\times \exp\left(\frac{\Delta t}{2} F(r(t)) \frac{\partial}{\partial p}\right) \begin{bmatrix} r(t) \\ p(t) \end{bmatrix}. \end{aligned} \quad (3.54)$$

The propagation is determined by acting each of the three operators starting

from the right on  $r$  and  $p$ . The result of applying the operator in a function  $g(r)$  is the Taylor expansion  $g(r + c)$ . Hence, after applying the first operator, we obtain the equation bellow (TUCKERMAN, 2010):

$$\exp \left\{ \frac{\Delta t}{2} F[r(t)] \frac{\partial}{\partial p} \right\} \begin{bmatrix} r(t) \\ p(t) \end{bmatrix} = \begin{bmatrix} r(t) \\ p(t) + \frac{\Delta t}{2} F(r(t)) \end{bmatrix}. \quad (3.55)$$

Acting the second operator in the preceding equation, the result is the following:

$$\exp \left[ \Delta t \frac{p(t)}{m} \frac{\partial}{\partial r} \right] \begin{bmatrix} r(t) \\ p(t) + \frac{\Delta t}{2} F[r(t)] \end{bmatrix} = \begin{bmatrix} r(t) + \frac{\Delta t}{m} p(t) \\ p(t) + \frac{\Delta t}{2} F[r(t) + \frac{\Delta t}{m} p(t)] \end{bmatrix}. \quad (3.56)$$

and, finally, we find the following equations after applying the third operator :

$$\exp \left\{ \frac{\Delta t}{2} F[r(t)] \frac{\partial}{\partial p} \right\} \begin{bmatrix} r(t) + \frac{\Delta t}{m} p(t) \\ p(t) + \frac{\Delta t}{2} F[r(t) + \frac{\Delta t}{m} p(t)] \end{bmatrix} = \begin{bmatrix} r(t) + \frac{\Delta t}{m} \{ p(t) + \frac{\Delta t}{2} F[r(t)] \} \\ p(t) + \frac{\Delta t}{2} F[r(t)] + \frac{\Delta t}{2} F \{ r(t) + \frac{\Delta t}{m} [p + \frac{\Delta t}{2} F(r(t))] \} \end{bmatrix}. \quad (3.57)$$

Using the equations above and substituting  $p/m$  for  $v$ , the final position  $r(t + \Delta t)$  can be written as

$$r(t + \Delta t) = r(t) + v(t)\Delta t + \frac{F(t)}{2m} \Delta t^2. \quad (3.58)$$

Eq. 3.58 is the position update part of the velocity-Verlet algorithm (VERLET, 1967). In this method, the positions are updated by a time step of  $\Delta t$  by using the positions at the previous time steps and forces. These recalculations of forces at each time step are the most computationally expensive part of the simulation since we have to take the derivative of the potential energy at each time step. The equation to update the velocity can also be derived from the equations above. It is given by

$$v(t + \Delta t) = v(t) + \frac{F(t + \Delta t) + F(t)}{2m} \Delta t. \quad (3.59)$$

Instead of using a time step of  $\Delta t$ , the velocities can be updated at  $\Delta t/2$ . This is the strategy proposed by the leap frog algorithm:

$$v(t + \Delta t/2) = v(t - \Delta t/2) + \frac{F(t)}{m} \Delta t, \quad (3.60)$$

$$r(t + \Delta t) = r(t) + v(t + \Delta t/2)\Delta t. \quad (3.61)$$

Although using different time steps or formulations, both of the methods described here generate the same trajectory for a given initial configuration.

### 3.1.5 Initial Configuration and Periodic Boundary Condition

The equations from the preceding section require an overlap-free initial configuration with positions and velocities for all atoms in the system. The initial velocities follow a temperature-dependent Maxwell-Boltzmann distribution, which is

$$\varrho(v_{x,i}) = \left( \frac{m_i}{2\pi\kappa_b T} \right)^{\frac{1}{2}} \exp \left( -\frac{m_i v_{x,i}^2}{2\kappa_b T} \right). \quad (3.62)$$

Random velocities are then found with the equation above for each of the 3N components of the velocity, and the initial positions can be obtained by several approaches. The initial configuration can be taken from an x-ray or a nuclear magnetic resonance (NMR) spectroscopy, the atoms can be placed randomly in the simulation volume, or the atoms can be placed in idealized or approximate geometries. The generally used method to acquire the configurations places the molecules on a cubic lattice (SHELL, 2015). Among the available software to optimize this placement, there is the Packmol software (MARTÍNEZ *et al.*, 2009). It treats the initial configuration problem as a packing optimization problem. The molecules are packed in such way that atoms from different molecules keep a safe pairwise distance and, due to the optimization function and gradient evaluations, this strategy enables the packing of millions of atoms in reasonable time (MARTÍNEZ *et al.*, 2009).

Independently of the technique or software used, certain restrictions should be applied in the initial configurations to carry out molecular simulations. As an example, the cubic lattice has a finite size, but a finite box would result in simulations dominated by surface effects. To avoid that and be able to simulate bulk phase, we can create a box periodically repeated in all directions by applying the so-called Periodic Boundary Conditions (PBC) (FRENKEL; SMIT, 2001). The periodic box has a primitive cell, which contains the N particles, replicated in a periodic lattice of infinite cells as represented in Figure 3.1.1.

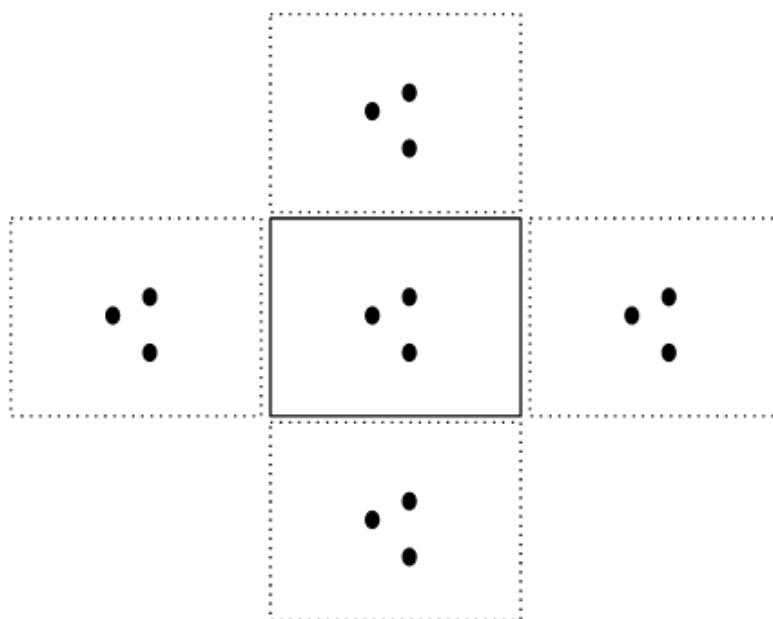


Figure 3.1.1 – Representation of the periodic boundary condition.

The application of the PBC results in particles interacting not only with each other but also with their images. This fact significantly increases the number of interacting pairs and, consequently, the computational time. To overcome that, we need to choose a limited range potential using the minimum image convention criterion. This criterion only allows a particle to interact with the nearest particle or image. This is technically done during the simulations by neglecting the interactions between two particles at or beyond a maximum length, which is given the name of cutoff radius ( $R_c$ ). This cutoff should be equal to or less than half of the box length. This process of examining each pair separation can also be expensive depending on the number of distinct pairs. That is the reason molecular dynamics algorithms use pair listings. This method defines a 'skin' around the cutoff radius with a radius  $R_{List}$ , as represented in Figure 3.1.2. The pair list is initially built of all the neighbor particles within a distance  $R_{List}$  of each particle. Over the course of the simulation, only pairs in the pair list interact. We then will have a list of all particles  $j$  with which a particle  $i$  interacts as an example. The pair list must then be updated from time to time during the simulation since particles typically diffuse (TUCKERMAN, 2010).



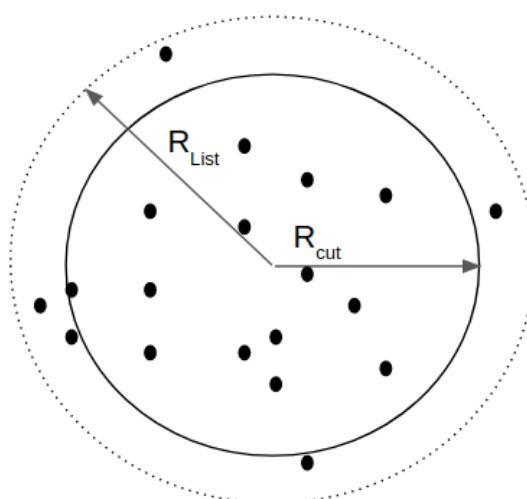


Figure 3.1.2 – Representation of the cutoff radius and the pair list radius. Adapted from [Tuckerman \(2010\)](#).

### 3.1.6 Force Fields

As said previously, we need to calculate the derivative of the potential energy function in relation to the coordinates in order to update the positions during a simulation. Therefore, we need a model for this potential energy functions. These models are called force fields. They can describe structural characteristics such as van der Waals interactions, bond lengths, bond angles, and torsion in a molecular simulation. The description is done by approximating the potential energy function  $[U(r^N)]$ , which has contributions due to intermolecular and intramolecular interactions. The intramolecular interactions include bond stretching, angle bending, and bond torsion. An illustration of these interactions can be seen in Figure 3.1.3.

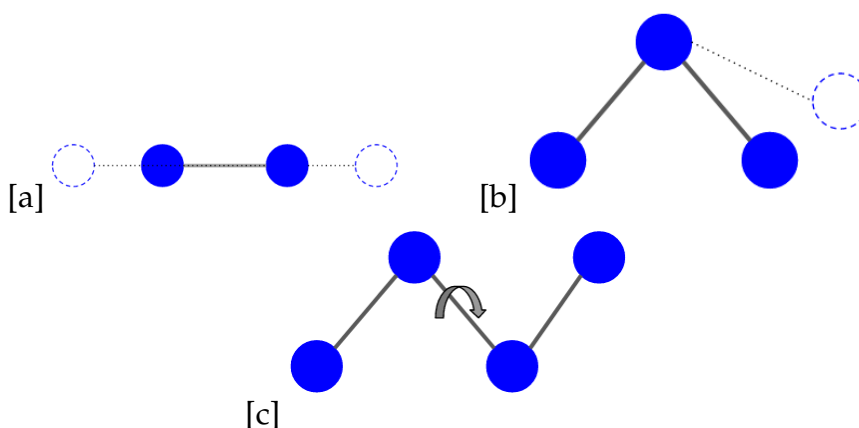


Figure 3.1.3 – Representation of bond stretching [a], angle bending [b], and bond torsion [c].

In this dissertation, we are going to present the equations which are most used

to represent these interactions. The contribution to the bond stretching (bs ) is usually given by the harmonic approximation around the energy minimum:

$$u_{bs}(d) = k_{bs}(d - d_0)^2. \quad (3.63)$$

Here,  $d$  is the bond length,  $d_0$  is the equilibrium bond length, and  $k_{bs}$  is a bond stretching constant. The angle bending (ab) contribution corresponds to deviations from the preferred geometry and is often given by:

$$u_{ab}(\theta) = k_{ab}(\theta - \theta_0)^2, \quad (3.64)$$

where  $k_{ab}$  and  $\theta_0$  are constants defined by the force field and  $\theta$  is the bond angle between three atoms. The bond torsion (bt) interactions correspond to the energies of rotations around bonds, and it happens among four atoms. A commonly used model is

$$u_{bt}(\omega) = \sum_{n=0}^N c_n \cos^n(\omega), \quad (3.65)$$

where  $N$  is the number of terms,  $c_n$  is the coefficient defined by the force field, and  $\omega$  is the torsional angle also defined by the force field.

The other type of interactions, the intermolecular interactions or non-bonded interactions, can include interactions due to charges, point-dipole moment, and van der Waals interactions. The first one represents the interaction between two atoms  $i$  and  $j$  with partial charges ( $q_i$  and  $q_j$ ) and they are usually represented by Coulomb's Law:

$$u_{qq}(r_{ij}) = \frac{q_i q_j}{4\pi\epsilon_0 r_{ij}}. \quad (3.66)$$

In the equation above,  $\epsilon_0$  is the free space permittivity constant and  $r_{ij}$  is the distance between atoms  $i$  and  $j$ . The interaction between the partial charges and the dipole moments of particles ( $u_{qp}$ ) and the dipole-dipole ( $u_{pp}$ ) interactions can be expressed as:

$$u_{qp}(r_{ij}) = \frac{q}{r_{ij}^3} (p \bullet \vec{r}_{ij}). \quad (3.67)$$

$$u_{pp}(r_{ij}) = \frac{1}{r_{ij}^3} (\vec{p}_i \bullet \vec{p}_j) - \frac{3}{r_{ij}^5} (\vec{p}_i \bullet \vec{r}_{ij})(\vec{p}_j \bullet \vec{r}_{ij}). \quad (3.68)$$

where  $p_i$  and  $p_j$  are dipole moments of the two particles.

In many force fields, the van der Waals interaction between particles  $i$  and  $j$  is

modeled by the Lennard-Jones Potential:

$$u_{LJ}(r_{ij}) = 4\epsilon \left[ \left( \frac{\sigma}{r_{ij}} \right)^{12} - \left( \frac{\sigma}{r_{ij}} \right)^6 \right], \quad (3.69)$$

where  $\epsilon$  is the depth of the potential well,  $\sigma$  is the distance corresponding to a zero intermolecular potential. The graphical representation of the Lennard-Jones potential is presented in Figure 3.1.4.

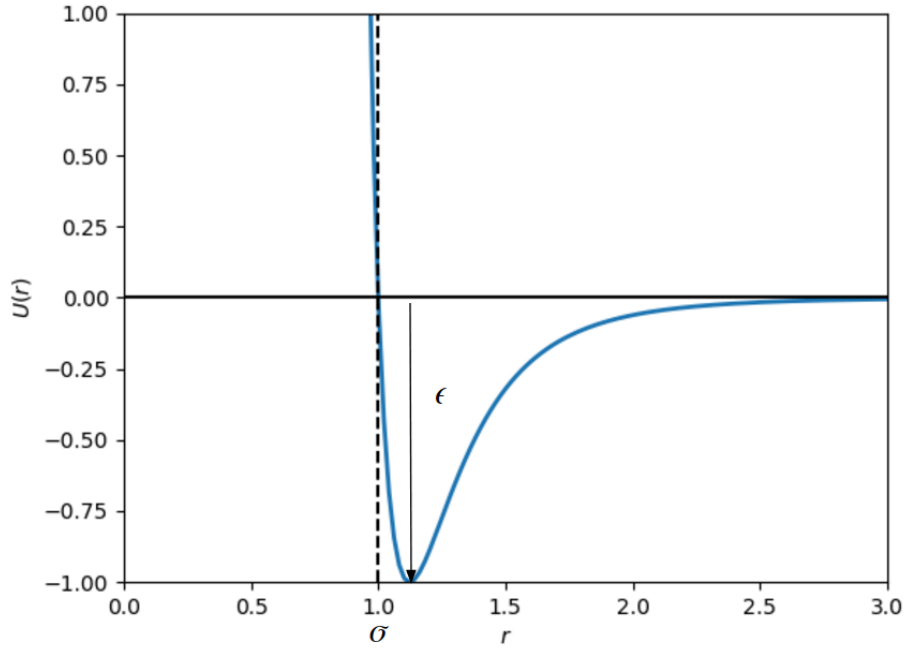


Figure 3.1.4 – Lennard-Jones potential representation for  $\sigma = 1$  and  $\epsilon = 1$ .

The potential in Figure 3.1.4 tends to zero and becomes negligible after a specific large value of  $r$ . In practice, this potential is truncated beyond a certain cutoff. The point in which the cutoff is defined is generally the one in which the radial distribution function  $[g(r)]$  is approximately constant. Also, only interactions with the nearest periodic image of the cell are considered for short-range interactions as explained in Section 3.1.5. With this conditions, the calculations of forces and velocities are computationally feasible. The final potential energy function defined by the force field is then expressed by summing all the interactions above:

$$U(r^N) = u_{bs}(d) + u_{ab}(\theta) + u_{bt}(\omega) + u_{qq}(r_{ij}) + u_{qp}(r_{ij}) + u_{pp}(r_{ij}) + u_{LJ}(r_{ij}). \quad (3.70)$$

## 3.2 SAFT- $\gamma$ Mie Force Field

### 3.2.1 SAFT-VR Mie Equation of State (EoS)

The SAFT-VR Mie equation of state (LAFITTE *et al.*, 2013) is the basis for the SAFT- $\gamma$  Mie coarse-grained force field (AVENDAÑO *et al.*, 2011), which was the force field used to model the molecules used in the simulations carried out in this dissertation. This EoS was initially developed by Lafitte *et al.* (2013) to describe chain molecules formed from fused segments interacting via the Mie attractive and repulsive potential. The Mie potential is a type of generalized Lennard-Jones potential that can be used to explicitly describe repulsive interactions of different hardness/softness and attractive interactions of different ranges, and is given by

$$U_{Mie}(r) = \epsilon \frac{\lambda_r}{\lambda_r - \lambda_a} \left( \frac{\lambda_r}{\lambda_a} \right)^{\left( \frac{\lambda_a}{\lambda_r - \lambda_a} \right)} \left[ \left( \frac{\sigma}{r} \right)^{\lambda_r} - \left( \frac{\sigma}{r} \right)^{\lambda_a} \right], \quad (3.71)$$

where  $\lambda_r$  is the repulsive exponent and  $\lambda_a$  is the attractive exponent. The SAFT-VR Mie equation uses the Barker and Henderson (1976) high perturbation expansion of the Helmholtz free energy up to the third order and an improved expression for the radial distribution function (RDF) of Mie monomers at contact to obtain an equation able to give an accurate theoretical description of the vapor-liquid equilibrium and second derivative properties (LAFITTE *et al.*, 2013). For a non-associating fluid, the Helmholtz free energy is

$$\frac{A^{SAFT}}{N\kappa_b T} = a^{SAFT} = a^{IDEAL} + a^{MONO} + a^{CHAIN}, \quad (3.72)$$

or, depending on the molecule type, the free energy can be equal to

$$\frac{A^{SAFT}}{N\kappa_b T} = a^{SAFT} = a^{IDEAL} + a^{MONO} + a^{RING}. \quad (3.73)$$

Here,  $a^{IDEAL}$  is the ideal contribution for a mixture. It is given by

$$a^{IDEAL} = \sum_{i=1}^{N_c} x_i \ln (\rho_i \Lambda_i^3) - 1, \quad (3.74)$$

where  $x_i = N_i/N_c$  is the molar fraction of component  $i$ ,  $N_i$  is the number of molecules of each component,  $N_c$  is the total number of molecules of the system,  $\rho_i = N_i/V$  is the number density, and  $\Lambda_i^3$  is the de Broglie thermal wavelength. Also in Eq. 3.72,  $a^{MONO}$  is the monomer contribution, which describes interactions between Mie segments and

can be expressed, for a mixture, as

$$a^{MONO} = \left( \sum_{i=1}^{N_c} x_i m_{s,i} \right) a^M. \quad (3.75)$$

In the equation above,  $m_{s,i}$  is the number of spherical segments making up the molecule  $i$  and  $a^M$  is the monomer dimensionless Helmholtz free energy.  $a^M$  is expressed as a third-order perturbation expansion in the inverse temperature (BARKER; HENDERSON, 1976):

$$a^M = a^{HS} + \beta a_1 + \beta^2 a_2 + \beta^3 a_3. \quad (3.76)$$

Here,  $a^{HS}$  is the hard-sphere dimensionless Helmholtz free energy for a mixture and is given by:

$$a^{HS} = \frac{6}{\pi \rho_s} \left[ \left( \frac{\zeta_2^3}{\zeta_3^2} - \zeta_0 \right) \ln(1 - \zeta_3) + \frac{3\zeta_1 \zeta_2}{1 - \zeta_3} + \frac{\zeta_2^3}{\zeta_3(1 - \zeta_3)^2} \right]. \quad (3.77)$$

The variable  $\rho_s = \rho \sum_i^{N_c} x_i m_{s,i}$  is the total number density of spherical segments and  $\zeta_l$  are the moments of the number density:

$$\zeta_l = \frac{\pi \rho_s}{6} \left( \sum_{i=1}^{N_c} x_{s,i} d_{ii}^l \right), \quad l = 0, 1, 2, 3, \quad (3.78)$$

where  $x_{s,i}$  is the mole fraction of segments. It is related to the mole fractions of all component ( $x_i$ ) by:

$$x_{s,i} = \frac{m_{s,i} x_i}{\sum_{k=1}^{N_c} m_{s,k} x_k}. \quad (3.79)$$

The effective hard-sphere diameter  $d_{ii}$  for the segments is

$$d_{ii} = \int_0^{\sigma_{ii}} \{1 - \exp[-\beta U_{ii}^{Mie}(r)]\} dr. \quad (3.80)$$

The integral in Eq. (3.80) is normally obtained by means of a 5-point Gauss-Legendre quadrature (PAPAIOANNOU *et al.*, 2014). For brevity, the detailing of the other terms of Eq. (3.76) are available in the Appendix A. The term  $a^{CHAIN}$  in Eq. 3.72 corresponds to the chain contribution. This chain formation of  $m_s$  tangentially bonded Mie segments is based on the first-order perturbation theory (TPT1) (PAPAIOANNOU *et al.*, 2014) and can be expressed as

$$a^{CHAIN} = - \sum_{i=1}^{N_c} x_i (m_{s,i} - 1) \ln [g_{ii}^{Mie}(\sigma_{ii})]. \quad (3.81)$$

The term  $g_{ij}^{Mie}(\sigma_{ij})$  correspond to the radial distribution function (RDF) of the hypothetical Mie system evaluated at the effective diameter. It is obtained with the following perturbation expansion

$$g_{ij}^{Mie}(\sigma_{ij}) = g_{d,ij}^{HS}(\sigma_{ij}) \exp \left[ \beta \epsilon \frac{g_{1,ij}(\sigma_{ij})}{g_{d,ij}^{HS}(\sigma_{ij})} + (\beta \epsilon)^2 \frac{g_{2,ij}(\sigma_{ij})}{g_{d,ij}^{HS}(\sigma_{ij})} \right]. \quad (3.82)$$

In the equation above,  $g_{d,ij}^{HS}$  is equal to

$$g_{d,ij}^{HS}(\sigma_{ij}) = \exp(k_0 + k_1 x_{0,ij} + k_2 x_{0,ij}^2 + k_3 x_{0,ij}^3), \quad (3.83)$$

where  $x_{0,ij} = \sigma_{ij}/d_{ij}$  and  $k_1, k_2$ , and  $k_3$  are density dependent coefficients. These coefficients and the other terms of Eq. 3.82 are available in Appendix A.

The ring contribution ( $a^{RING}$ ) in Eq. 3.73 have two forms for rings formed from tangentially bonded segments. The first was developed by Lafitte *et al.* (2012). It considers that the difference between a chain and a ring molecule is that the latter has one more bond that is connecting the first segment to the last. With this assumption, Eq. (3.81) can be adapted to rings by

$$a^{RING} = - \sum_{i=1}^{N_c} x_i m_{s,i} \ln[g_{ii}^{Mie}(\sigma_{ii})]. \quad (3.84)$$

According to Lafitte *et al.* (2012), Eq. (3.84) needs an additional parameterization with molecular simulation data so that the EoS can be used in molecular simulations, but this additional parameterization is not necessary when we are modeling chain molecules. Recently, Müller and Mejía (2017) tried to correct this inconsistency. They developed a Helmholtz free energy equation for rings based on the work of Müller and Gubbins (1993), who obtained rigorous expressions for hard-sphere fluids with molecular geometries of rings with  $m_s = 3$ . The final expression developed for the dimensionless Helmholtz free energy due to ring formation is

$$a^{RING} = - \sum_{i=1}^{N_c} x_i (m_{s,i} - 1 + \chi_i \eta_i) \ln [g_{ii}^{Mie}(\sigma_{ii})], \quad (3.85)$$

where  $\eta_i = m_{s,i} \rho_i \sigma_{ii}^3/6$  is the packing fraction of the atom  $i$  and  $\chi_i$  is a parameter which depends on  $m_{s,i}$  and the geometry of the ring of each component  $i$ . For a value of  $\chi = 0$ , Eq. (3.85) is equal to Eq. (3.81). In addition, the equation corresponds to a system of hard sphere triangles when  $\chi = 1.3827$ . Müller and Mejía (2017) also calculated values of  $\chi$  for  $m_s = 3, m_s = 4, m_s = 5$ , and  $m_s = 7$  using a geometry formed by a system of equilateral triangles. In these calculations, they used pseudo-experimental data from

molecular dynamics (MD) for a defined pure fluid with  $\epsilon/\kappa_b = 250K$ ,  $\sigma = 3.0\text{\AA}$ ,  $\lambda_r = 11$ , and  $\lambda_r = 6$ . Values of  $\chi$  for some of the geometry estimated in their article can be seen in Figure 3.2.1.

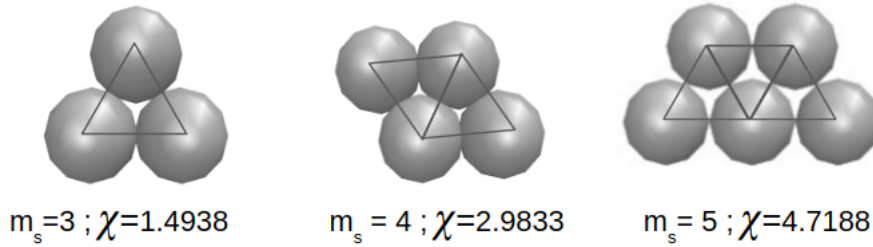


Figure 3.2.1 – Values for parameter  $\chi$  according to the ring geometry. Adapted from Müller and Mejía (2017).

Lafitte *et al.* (2013) also suggested mixing rules for this EoS parameters based on Lorentz-Berthelot combining rules (ROWLINSON; SWINTON, 1982):

$$\sigma_{ij} = \frac{\sigma_{ii} + \sigma_{jj}}{2}, \quad (3.86)$$

$$d_{ij} = \frac{d_{ii} + d_{jj}}{2}, \quad (3.87)$$

$$\lambda_{k,ij} - 3 = \sqrt{(\lambda_{k,ii} - 3)(\lambda_{k,jj} - 3)}, \quad k = r, a, \quad (3.88)$$

$$\epsilon_{ij} = (1 - k_{ij}) \frac{\sqrt{\sigma_{ii}^3 \sigma_{jj}^3}}{\sigma_{ij}^3} \sqrt{\epsilon_{ii} \epsilon_{jj}}. \quad (3.89)$$

The  $k_{ij}$  is a binary interaction parameter to correct the deviations of the mixing rule. This parameter can be described as a scaling factor. It accounts for the interactions among chemically distinct compounds, which are not explicitly considered by the SAFT-VR Mie EoS. These mixing rules are the ones available in the literature and employed by other papers that used this force field. Therefore, we ended up using Eqs. 3.86 to 3.89 in our study. However, the binary interaction parameter was only necessary for aqueous mixtures in our study. With the dimensionless Helmholtz free energy provided by the equations above, we can now calculate the derivative properties with the fundamental thermodynamic equation (Eq. 3.10) using the EoS. The output from these calculations are then used to estimate the parameters of the SAFT- $\gamma$  Mie force field.

### 3.2.2 Parameter Estimation for the SAFT- $\gamma$ Mie Force Field

The SAFT- $\gamma$  Mie Force Field uses a top-down coarse-graining methodology in its parameterization. This methodology aims to obtain the intermolecular parameters

from macroscopic experimental data such as fluid-phase equilibrium or interfacial tension data. The idea is that the force field parameters estimated with the SAFT-VR Mie EoS can be used in molecular simulations since both the equation of state and the force field use the same explicit intermolecular potential model (Mie potential). This correspondence between models has been used to parametrize a variety of fluids (ERVIK *et al.*, 2016b). This force field has the advantage of incorporating the degrees of freedom provided by the use of the Mie Potential (HERDES *et al.*, 2015). This flexibility offers the exploration of a vast parameter space without using an iterative simulation scheme (AVENDAÑO *et al.*, 2011). Despite these advantages, the force field can be restricted by the shortcomings of the equation of state. As an example, the lack of an association term in the equation can cause an inadequate representation of the properties of hydrogen bonding compounds.

Each substance has initially five parameters to be estimated ( $m_s$ ,  $\sigma$ ,  $\epsilon$ ,  $\lambda_r$ , and  $\lambda_a$ ) according to Eq. (3.71). The number of segments is usually fixed in an integer value since each segment represents one pseudo atom. The attractive parameter is generally fixed due to its high correlation with the repulsive parameter. Usually, the chosen value for this parameter is 6, corresponding to the London model, which is a good representation of the dispersion scale of most simple fluids that do not have strong polar interactions (RAMRATTAN *et al.*, 2015; HERDES *et al.*, 2015). There are two strategies to obtain the parameters: one is by fitting the SAFT-VR Mie EoS to experimental data such as vapor pressure, liquid density and the other one is by using corresponding states parametrization. The first was the one followed in this dissertation. Generally, this approach minimizes the following unweighted least-squares objective function:

$$F_{obj} = \sum_{i=1}^{N_p} \left[ \frac{P_v^{SAFT}(T_i, \sigma, \epsilon, \lambda_r) - P_v^{exp}(T_i)}{P_v^{exp}(T_i)} \right]^2 + \sum_{i=1}^{N_p} \left[ \frac{\rho_l^{SAFT}(T_i, \sigma, \epsilon, \lambda_r) - \rho_l^{exp}(T_i)}{\rho_l^{exp}(T_i)} \right]^2, \quad (3.90)$$

where  $N_p$  is the number of experimental points,  $P_v$  is the vapor pressure, and  $\rho_l$  is the saturated liquid density. Other properties that can be used in the estimation are interfacial tension and speed of sound, for instance. The multiple parameters of the model make it necessary the use of a wide range of experimental data since multiple solutions may be found for the fit. Therefore, one needs to be careful in deciding the level of coarse-graining (i.e. the choice of parameter  $m_s$ ) and the subsequent parameter space so as to avoid some physical inconsistencies such as a premature freezing (LOBANOVA *et al.*, 2015).

Lafitte *et al.* (2012) suggested that two correction factors ( $c_\sigma$  and  $c_\epsilon$ ) should be estimated with simulation data when using Eq. (3.84) for the ring contribution. They



are related to the EoS parameters by scaled parameters:

$$\sigma^{scaled} = c_\sigma \sigma^{SAFT}. \quad (3.91)$$

$$\epsilon^{scaled} = c_\epsilon \epsilon^{SAFT}. \quad (3.92)$$

According to [Lafitte \*et al.\* \(2012\)](#), these corrections are necessary because the approximations employed in the EoS theory generate discrepancies between molecular simulations and the EoS for ring molecules modeled with Eq. (3.84). However, this new parameterization is not necessary when using Eq. (3.85) as the ring contribution or when we are modeling chain molecules with Eq. 3.81. This fact makes the strategy of [Lafitte \*et al.\* \(2012\)](#) inconsistent since parameterization with molecular simulation should not be necessary according to the overall idea of this force field. Furthermore, the use of molecular simulation data highly increases the time spent on the parameterization process. The objective function for the estimation of the correction parameter is given by

$$F_{obj} = \sum_{i=1}^{N_p} \left[ \frac{P_v^{sim}(T_i, \sigma^{SAFT}, \epsilon^{SAFT}) - P_v^{SAFT}(T_i, \sigma^{scaled}, \epsilon^{scaled})}{P_v^{sim}(T_i, \sigma^{SAFT}, \epsilon^{SAFT})} \right]^2 + \sum_{i=1}^{N_p} \left[ \frac{\rho_{liq}^{sim}(T_i, \sigma^{SAFT}, \epsilon^{SAFT}) - \rho_{liq}^{SAFT}(T_i, \sigma^{scaled}, \epsilon^{scaled})}{\rho_{liq}^{sim}(T_i, \sigma^{SAFT}, \epsilon^{SAFT})} \right]^2. \quad (3.93)$$

The repulsive parameter is maintained in the value found on the minimization of Eq. (3.90). The refined values for  $\sigma$  and  $\epsilon$  are

$$\sigma^{sim} = \sigma^{SAFT} / c_\sigma, \quad (3.94)$$

$$\epsilon^{sim} = \epsilon^{SAFT} / c_\epsilon, \quad (3.95)$$

The other method to obtain the force field parameters is the corresponding states parametrization ([MEJÍA \*et al.\*, 2014](#)). This method considers that the unweighted volume average of the attractive contribution to the Mie intermolecular potential,  $a_1$ , is the following mean-field approximation

$$a_1 = 2\pi\rho\sigma^3\epsilon\alpha. \quad (3.96)$$

The van der Waals constant,  $\alpha$ , considering  $\lambda_a = 6$  is related by the Mie exponents by

$$\alpha = \frac{1}{\epsilon\sigma^3} \int_{\sigma}^{\infty} \phi(r)r^2 dr = \frac{\lambda_r}{3(\lambda_r - 3)} \left( \frac{\lambda_r}{6} \right)^{6/(\lambda_r - 6)}. \quad (3.97)$$

The parameterization in this method starts by using the experimental acentric factor,  $\omega$ , for each molecule with a fixed value of  $m_s$  in order to obtain the value of the repulsive exponent with the following Padé series:

$$\lambda_r = \frac{\sum_{i=0} a_i \omega^i}{1 + \sum_{i=1} b_i \omega^i}, \quad (3.98)$$

where  $a_i$  and  $b_i$  are parameters that are dependent on the number of segments. A table with these parameters is presented in the original paper (MEJÍA *et al.*, 2014). The van der Waals constant can be found by substituting  $\lambda_r$  into Eq. (3.97). The reduced critical temperature  $T_c^*$  is related to  $\alpha$  by a Padé series:

$$T_c^* = \frac{\sum_{i=0} c_i \alpha^i}{1 + \sum_{i=1} d_i \alpha^i}. \quad (3.99)$$

The values of  $c_i$  and  $d_i$  are also available in the original paper. The reduced temperature of the equation above is used in conjunction with the experimental critical temperature,  $T_c$ , to find the energy parameter with the relation below:

$$T_c^* = \frac{\kappa_b T_c}{\epsilon}. \quad (3.100)$$

The diameter parameter, however, is not obtained with the critical properties, but with the reduced liquid density at the reduced temperature of 0.7 ( $\rho_{T_r=0.7}$ ). This density is also obtained with a Padé series using parameters by Mejía *et al.* (2014):

$$\rho_{T_r=0.7}^* = \frac{\sum_{i=0} e_i \alpha^i}{1 + \sum_{i=1} f_i \alpha^i}. \quad (3.101)$$

The coefficients  $a_i, b_i, \dots, f_i$  depend on the number of segments, and they are available in the literature for ring and chain geometries (MÜLLER; MEJÍA, 2017; MEJÍA *et al.*, 2014). The relation between the Eq. 3.101,  $\sigma$  and the experimental density is given by

$$\rho_{T_r=0.7}^* = \rho_{T_r=0.7} \sigma^3 N_{av}, \quad (3.102)$$

where  $N_{av}$  is the Avogadro number. This corresponding states method has the advan-

tage of only requiring critical data, which is available for a great range of fluids, and liquid density data. The parameters obtained with these parameterization strategies are available at a large online database for this force field (ERVIK *et al.*, 2016b). Here, we only show the coarse-grained geometries and the sets of parameters for the compounds used in this dissertation. They are available in Tables 3.2.1 and 3.2.2. The parameters for water were retrieved from Lobanova *et al.* (2016); for carbon dioxide, propane, and hexane from Herdes *et al.* (2015); for 1-octanol from Ervik *et al.* (2016b); and for the aromatic compounds from Müller and Mejía (2017).

Table 3.2.1 – Parameters of the SAFT- $\gamma$  Mie force field for each substance used in this work.

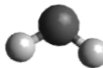



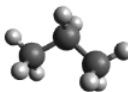

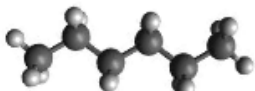

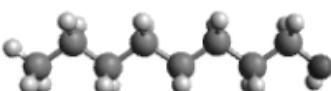
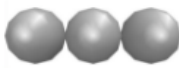
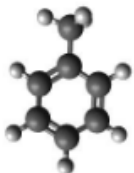

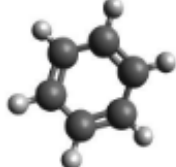
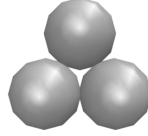
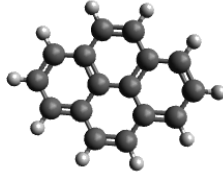

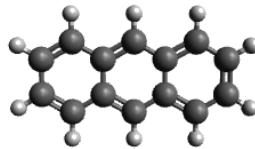
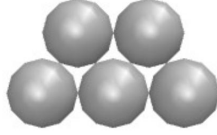
	$m_s$	$\epsilon/\kappa_b$ (K)	$\sigma$ (Å)	$\lambda_r$
Water	1	305.21	2.902	8.0
Carbon dioxide	2	194.94	2.848	14.65
Propane	1	426.08	4.871	34.29
Hexane	2	376.35	4.508	19.57
1-octanol	3	495.71	4.341	28.79
Toluene	3	268.24	3.685	11.80
Benzene	3	230.30	3.441	10.45
Pyrene	4	459.04	4.134	15.79
Anthracene	5	259.68	3.631	9.55

When modeling mixtures with this force field, it can be necessary to estimate the binary interaction parameter  $k_{ij}$  of Eq. (3.89). This parameter is normally estimated by minimizing the difference between experimental binary vapor-liquid equilibrium or interfacial tension data and the SAFT-VR Mie EoS output data (MÜLLER; MEJÍA, 2017; LOBANOVA *et al.*, 2016). The objective function is similar to:

$$F_{obj} = \sum_{k=1}^{N_p} \left( \frac{P_v^{SAFT}(T_k, x, k_{ij}) - P_v^{exp}(T_k, x)}{P_v^{exp}(T_k, x)} \right)^2 + \sum_{k=1}^{N_p} \left( \frac{\rho_l^{SAFT}(T_k, x, k_{ij}) - \rho_l^{exp}(T_i)}{\rho_l^{exp}(T_i)} \right)^2. \quad (3.103)$$

However, Ervik *et al.* (2016a) used molecular simulation results to fit the parameter to the interfacial tension data. The strategy they followed was to carry out simulations in three values of  $k_{ij}$  first and, after, refine the parameter until a value in good agreement with the experimental data is found. We decided to follow this strategy in our estimations of  $k_{ij}$  since the estimation with the EoS did not provide satisfactory results for the hydration free energy calculations.

Table 3.2.2 – Coarse-grained geometries of the substances used in this work for the SAFT- $\gamma$  Mie force field.

Molecule	Structure	Coarse grained structure
Water		
Carbon dioxide		
Propane		
Hexane		
1-octanol		
Toluene		
Benzene		
Pyrene		
Anthracene		

### 3.3 Solvation Free Energy Calculations Based on Molecular Dynamics

Using the SAFT- $\gamma$  Mie Force Field described in the section above, we carried out solvation free energy molecular dynamics simulations. The free energies we are trying to calculate can be expressed as averages over ensembles of atomic configurations generated using Monte Carlo or Molecular Dynamics techniques. In the canonical

ensemble, the free energy is given by

$$F(N, V, T) = -\kappa_b T \ln Q(N, V, T). \quad (3.104)$$

Recall that  $Q(N, V, T)$  is the partition function of the canonical ensemble, expressed as

$$Q(N, V, T) = \frac{1}{h^{3N} N!} \int dp^n dr^n \exp \left[ -\beta \left( \sum_{i=1}^N \frac{p_i^2}{2m_i} + U(r_1, \dots, r_n) \right) \right]. \quad (3.105)$$

The Gibbs free energy, the object of study in this dissertation, is given by

$$G(N, P, T) = -\kappa_b T \ln \Delta(N, P, T), \quad (3.106)$$

where  $\Delta(N, P, T)$  is the partition function of the isothermal-isobaric ensemble:

$$\Delta(N, P, T) = \frac{1}{V_0 N! h^{3N}} \int_0^\infty dV \int dp^n dr^n \exp \left[ -\beta \left( \sum_{i=1}^N \frac{p_i^2}{2m_i} + U(r_1, \dots, r_n) + PV(r_1, \dots, r_n) \right) \right]. \quad (3.107)$$

Evaluating the partition function is an often unfeasible task, but we are interested in calculating only the Gibbs free energy difference between two states of a system, which is

$$\Delta G_{AB} = G_B - G_A = -\kappa_b T \ln \left( \frac{\Delta_B}{\Delta_A} \right). \quad (3.108)$$

Since the masses of particles in states A and B of a system are the same and the Hamiltonian is separable in  $K(p)$  and  $U(r)$ , the moment integrals in the ratio  $\Delta_B/\Delta_A$  can be simplified into the ratio of the configuration integrals:

$$\frac{Z_B}{Z_A} = \frac{\int_0^\infty dV \int dr^n \exp \{ -\beta [U(r_1, \dots, r_n) + PV(r_1, \dots, r_n)]_B \}}{\int_0^\infty dV \int dr^n \exp \{ -\beta [U(r_1, \dots, r_n) + PV(r_1, \dots, r_n)]_A \}}. \quad (3.109)$$

This identity results in the following equation for the Gibbs free energy difference, which does not require the calculation of the partition function at each state:

$$\Delta G_{AB} = G_B - G_A = -\kappa_b T \ln \left( \frac{Z_B}{Z_A} \right). \quad (3.110)$$

In the case of a study concerning the solvation of a single molecule, the Gibbs free energy difference between end states *A* and *B* are, more specifically, the difference between the solute alone in the gas phase and the solute interacting with the solvent. In order to have accurate results for free energy differences, the states need to have sufficient overlap ([KLIMOVICH \*et al.\*, 2015](#)). The overlap can be achieved by calculating the free energy difference among a series of intermediates states. The result of these differences is independent of the path chosen since free energy is a state function. That is why alchemical states (without physical sense) can be used to link physical states of interest. The solvation free energy calculations are then done through a thermodynamic path in which the solute molecule is gradually inserted into the solvent as illustrated in [Figure 3.3.1](#). According to this path, the free energy of solvation is expressed as

$$\Delta G_{solv} = \Delta G_{1 \rightarrow 4} = \Delta G_{1 \rightarrow 2} + \Delta G_{2 \rightarrow 3} + \Delta G_{3 \rightarrow 4} - \kappa_b T \ln \frac{V^*}{V_1}. \quad (3.111)$$

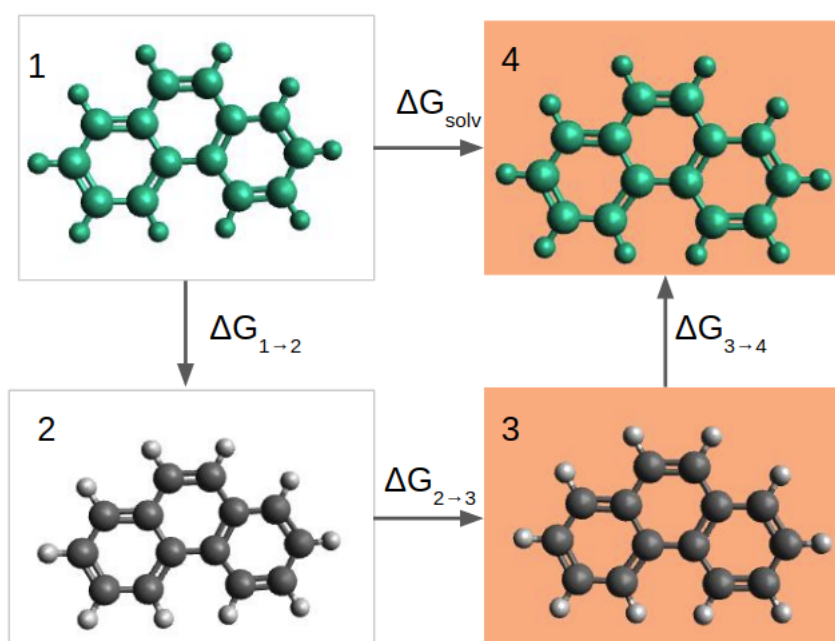


Figure 3.3.1 – Thermodynamic path for computing solvation free energy of a single solute molecule with molecular dynamics. Adapted from [Klimovich \*et al.\* \(2015\)](#).

The last term in Eq. 3.111 accounts for the difference between the mean volume of the simulation box with the solute inserted ( $V_1$ ) and the mean volume of the simulation box with only solvent molecules in it ( $V^*$ ). However, [Shirts \*et al.\* \(2003\)](#) have shown that this term is negligible with respect to the statistical uncertainty of calculating  $\Delta G$ . In addition, when we have other solute molecules in solution, another term in Eq. 3.111 may be necessary ([SHIRTS \*et al.\*, 2003](#)). The term  $\Delta G_{1 \rightarrow 2}$ , also represented in [Figure 3.3.1](#), is the solvation free energy associated with turning off the non bonded

interactions of the molecule in the gas phase. In the next transformation,  $\Delta G_{2 \rightarrow 3}$  is the free energy of moving the non-interacting molecule from the gas phase to the solvent and is equal to zero since the transformation of a non-interacting molecule does not depend on the environment. Lastly,  $\Delta G_{3 \rightarrow 4}$  is the free energy required for the non-interaction molecule in the solvent phase to regain its non-bonded interactions with the solvent. The solvation free energy calculation can be classified according to the types of non-bonded interactions that are turned off/on in the  $1 \rightarrow 2$  and  $3 \rightarrow 4$  parts of the path. If both non-bonded interactions with the environment and internal interactions are turned off, this is an annihilation free energy calculation. On the other hand, if only non-bonded interactions with the environment are turned off, this is a decoupling free energy calculation (KLIMOVICH *et al.*, 2015). In the latter case,  $\Delta G_{1 \rightarrow 2} = 0$  and the  $\Delta G_{solv} = \Delta G_{3 \rightarrow 4}$ .

The methods used to carry out the transformations of Figure 3.3.1 during the simulation scale the solute charges to zero and then turn off the interactions corresponding to the Lennard-Jones or Mie potential. In order to carry out the latter process, a modified potential with a coupling parameter ( $\lambda$ ) is used. Each  $\lambda$  represents an alchemical state. When  $\lambda = 0$ , there is no interaction with the solvent and, when  $\lambda = 1$ , interactions are fully activated. The coupling of the  $\lambda$  parameter could be linear, as it is demonstrated in Figure 3.3.2, but it could generate numerical problems related to the exponential part of the potential (SHIRTS *et al.*, 2003).

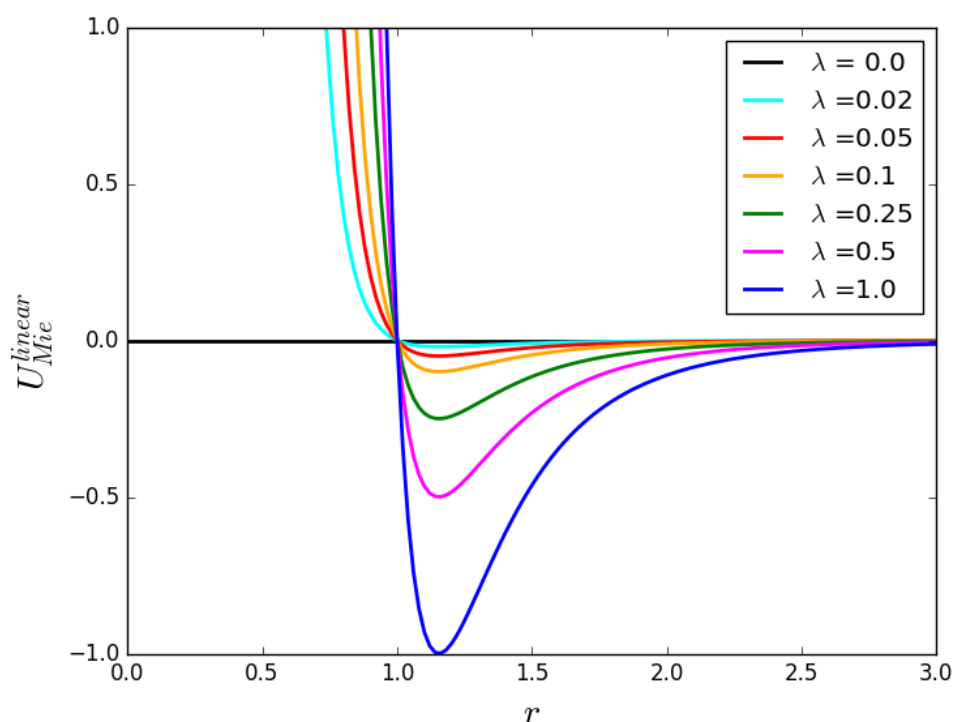


Figure 3.3.2 – Linear coupling of the potential energy,  $U_{Mie}^{linear} = \lambda U_{Mie}$ , for different values of  $\lambda$ . Here,  $\sigma = 1$ ,  $\epsilon = 1$ ,  $\lambda_a = 6$ , and  $\lambda_r = 8$ .

Observing Figure 3.3.2, we can see that a linear coupling of the parameter would cause an abrupt change in the potential energy when the  $\lambda$  reaches zero. That is the reason the non-linear softcore scheme (BEUTLER *et al.*, 1994) is used to couple the  $\lambda$ . This scheme makes the potential behave more smoothly in relation to the change of  $\lambda$ . The softcore potential is equal to

$$U_{LJ}^{sc}(r) = 4\lambda\epsilon \left\{ \frac{1}{[\alpha(1-\lambda) + (r/\sigma)^6]^2} - \frac{1}{\alpha(1-\lambda) + (r/\sigma)^6} \right\}, \quad (3.112)$$

where  $\alpha$  is a constant whose value is normally assumed to be 0.5. Based on Eq. 3.112, we propose a generalized softcore Mie potential for any value of  $\lambda_r$  and  $\lambda_a$ . It is given by:

$$U_{Mie}^{sc}(r) = \lambda\epsilon \frac{\lambda_r}{\lambda_r - \lambda_a} \left( \frac{\lambda_r}{\lambda_a} \right)^{\left( \frac{\lambda_a}{\lambda_r - \lambda_a} \right)} \left\{ \frac{1}{[\alpha(1-\lambda) + (r/\sigma)^{\lambda_a}]^{\lambda_r/\lambda_a}} - \frac{1}{\alpha(1-\lambda) + (r/\sigma)^{\lambda_a}} \right\}. \quad (3.113)$$

With the intention of observing the behavior of this generalized softcore potential, we plotted the  $U_{Mie}^{sc}(r)$  with repulsive exponents of different hardness/softness for a range of values of  $\lambda$  in Figures 3.3.3 to 3.3.6. We see that this softcore scheme makes the potential behave more smoothly in relation to the change of  $\lambda$ , mainly for the softer repulsive exponents.

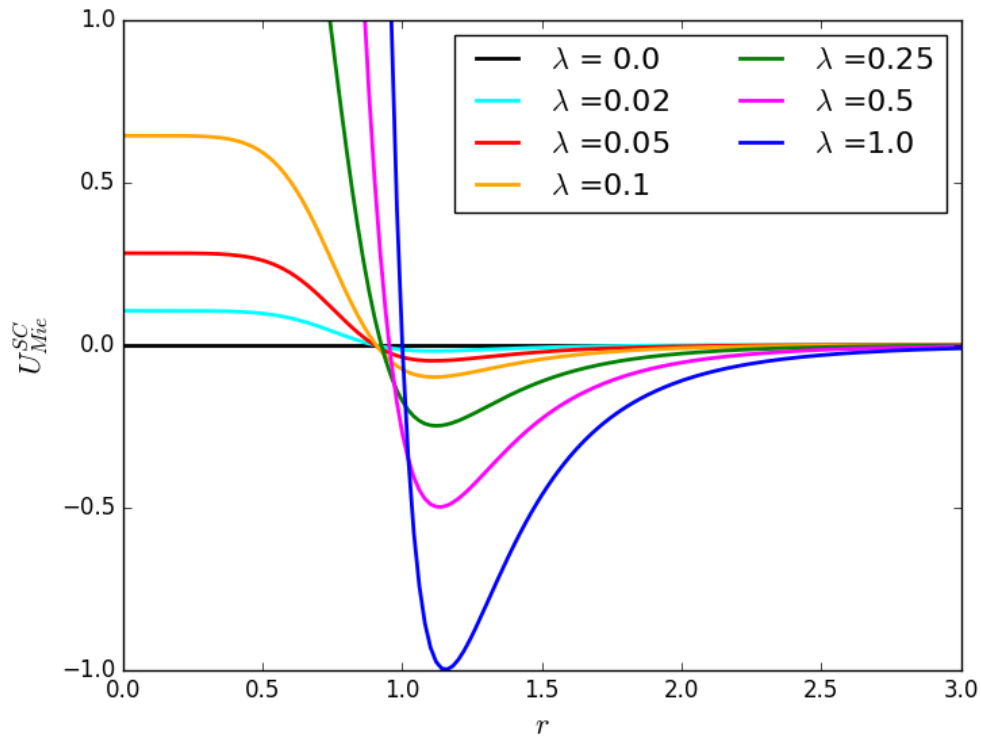


Figure 3.3.3 – Generalized softcore Mie potential, Eq. 3.113, for different values of  $\lambda$ . Here,  $\sigma = 1$ ,  $\epsilon = 1$ ,  $\lambda_a = 6$ , and  $\lambda_r = 8$



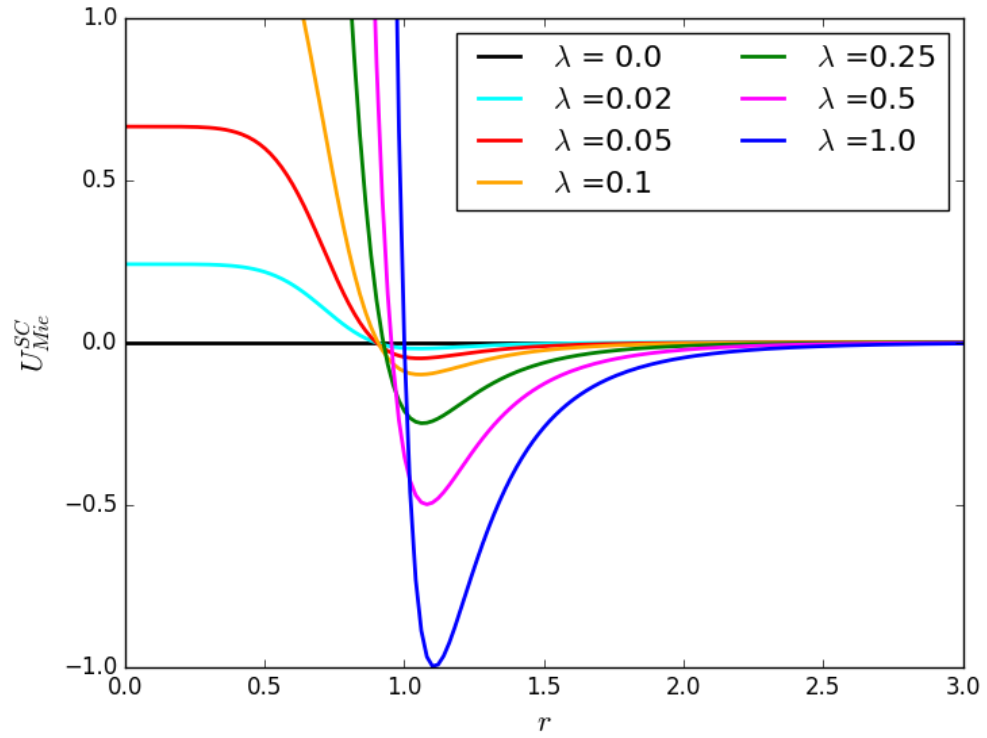


Figure 3.3.4 – Generalized softcore Mie potential, Eq. 3.113, for different values of  $\lambda$ . Here,  $\sigma = 1$ ,  $\epsilon = 1$ ,  $\lambda_a = 6$ , and  $\lambda_r = 15$

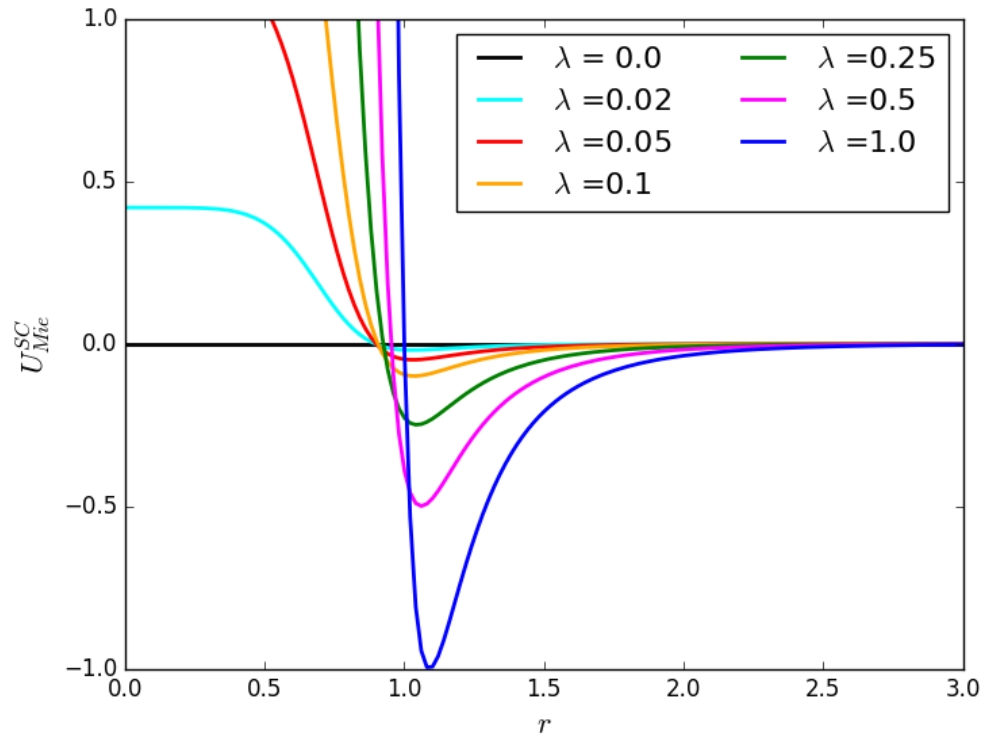


Figure 3.3.5 – Generalized softcore Mie potential, Eq. 3.113, for different values of  $\lambda$ . Here,  $\sigma = 1$ ,  $\epsilon = 1$ ,  $\lambda_a = 6$ , and  $\lambda_r = 20$

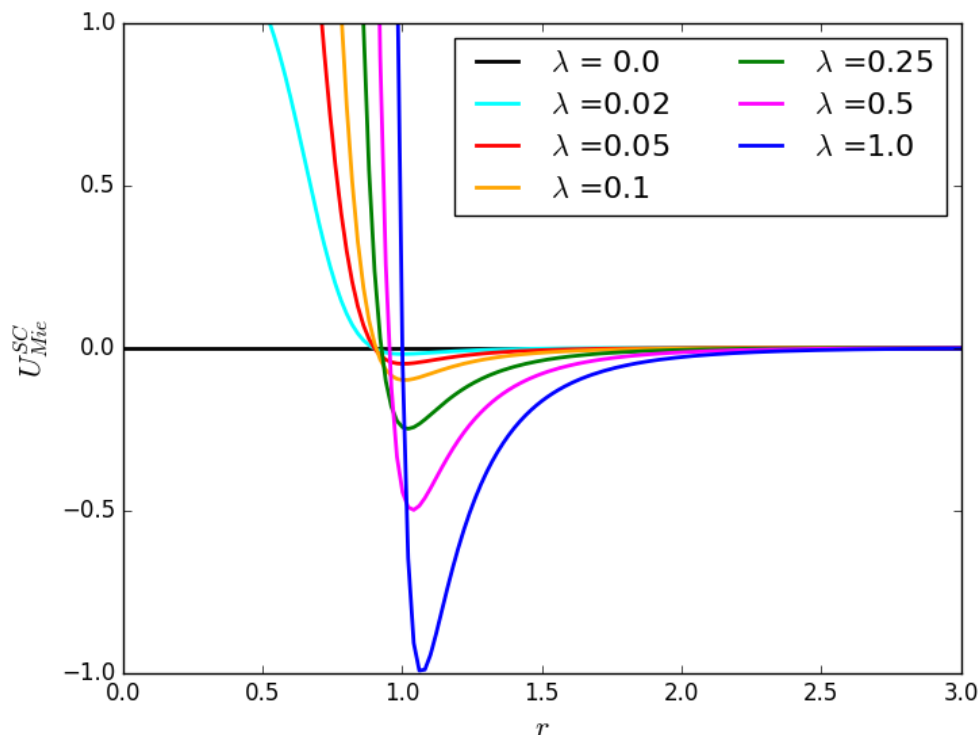


Figure 3.3.6 – Generalized softcore Mie potential, Eq. 3.113, for different values of  $\lambda$ . Here,  $\sigma = 1$ ,  $\epsilon = 1$ ,  $\lambda_a = 6$ , and  $\lambda_r = 30$

Now that we defined our coupled potential, we can then obtain the potential energies related to each alchemical state by doing independent simulations in different values of  $\lambda$  or by doing expanded ensemble simulations (LYUBARTSEV *et al.*, 1992). The latter was the method used in this dissertation, and it is described in Section 3.4. With the potential energies, the next step is to use post-processing methods, such as the MBAR used in this study, to effectively calculate  $\Delta G_{3 \rightarrow 4}$ . The solvation free energies can then be used to calculate other properties such as the partition coefficient. This property is a measure of the partitioning of one solute in two solvents (a and b) with different physicochemical characteristics at a temperature T. It is defined by the following equation when the activity coefficients are assumed to be one (ESSEX *et al.*, 1992):

$$P = \frac{[solute]_a}{[solute]_b}, \quad (3.114)$$

where  $[solute]_a$  and  $[solute]_b$  are the concentration of the solute in the solvent a and b, respectively. Since  $P$  is an equilibrium constant, it can be related to free energy change associated with transferring the solute from the phase a to the phase b. Hence, we can define the relation between the partition coefficient and the difference in free energy

with the equation below (ESSEX *et al.*, 1992):

$$2.303RT \log P^{a/b} = \Delta G_{solv}^a - \Delta G_{solv}^b, \quad (3.115)$$

where 2.303 is a conversion factor.

### 3.4 Expanded Ensemble Method

We decided to use the Expanded Ensemble method (LYUBARTSEV *et al.*, 1992) in our solvation free energy simulations since it allows a non-Boltzmann sampling scheme of different states in a single simulation. Lyubartsev *et al.* (1992) initially proposed in their paper a sampling scheme of different temperatures, but this idea can be generalized to a sampling scheme of different states or  $\lambda$ 's (ESCOBEDO; MARTINEZ-VERACOECHEA, 2007). In this scheme, the sampling is done by biasing the phase space exploration process with weights not related to the statistical ensemble. The partition function of the statistical expanded ensemble,  $Z^{EE}$ , is obtained from the probability distributions corresponding to each  $\lambda$ . Hence,  $Z^{EE}$  is defined as a sum of subensembles  $Z_i$  in different values of  $\lambda$ , that is,

$$Z^{EE} = \sum_{i=1}^N Z_i(\lambda_i) \exp(\eta_i), \quad (3.116)$$

where  $N$  is the number of alchemical states,  $\eta_i$  is the arbitrary weight of the subensemble at each state, and  $Z_i$  is the configurational partition function of state  $i$ . For the isothermal-isobaric ensemble,  $Z_i$  is given by

$$Z_i = \frac{1}{V_0} \int_0^\infty dV \int dr^n \exp \{ -\beta_i [U(\lambda, r_1, \dots, r_n) + P_i V(r_1, \dots, r_n)] \}. \quad (3.117)$$

In solvation free energy calculations with molecular dynamics,  $\lambda$  corresponds to the coupling parameter of the softcore potential (Eq. 3.112). Since we are carrying out molecular dynamic simulations, the sampling of the expanded ensemble is done by performing an arbitrary number of MD steps followed by a  $\lambda$  transition. Chodera and Shirts (2011) proved that this type of sampling of the expanded ensemble is similar to the Gibbs sampling method (GEMAN; GEMAN, 1984; LIU, 2002). Following the Gibbs method, the sampling of the configuration space  $x$  for one state  $\lambda_k$  during the MD steps is done by using the conditional distribution:

$$\pi(x|\lambda_k) = \frac{\exp[-\beta u(x, \lambda_k)]}{\int dx \exp[-\beta u(x, \lambda_k)]}. \quad (3.118)$$

The state transition in the MD simulation uses the following conditional distribution:

$$\pi(\lambda_k|x) = \frac{\exp[-\beta u(x, \lambda_k) + \eta_k]}{\sum_{k=1}^K \exp[-\beta u(x, \lambda_k) + \eta_k]}, \quad (3.119)$$

where  $u(x, \lambda_k)$  is the reduced potential function for the NPT ensemble. There is a variety of acceptance schemes to do the expanded sampling using Eq. (3.119), but [Chodera and Shirts \(2011\)](#) suggested that the independence sampling ([LIU, 2002](#)) is the best strategy to increase the number of uncorrelated configurations. The implementation they suggested consists of updating the state index from  $i$  to  $j$  by first generating a uniform random number  $R$  on the interval  $[0, 1)$  and then selecting the smallest new value of  $j$  that satisfies the relation

$$R < \sum_{i=1}^j \pi(\lambda_i|x). \quad (3.120)$$

The sampling strategy above depends on a proper selection of weights in order to guarantee an adequate sampling of the states. If there is not a sufficient number of visits to each state, the expanded ensemble becomes deficient in obtaining input data to estimate free energy differences with the methods exposed in Section 2.4. Here, we followed the flat-histogram approach ([BERG; NEUHAUS, 1992; LEE, 1993; DAYAL et al., 2004](#)) to calculate the weights. This strategy aims to obtain adequate sampling by ensuring that all the states have an equal number of visits, i.e., the ratio of the probability of sampling state  $i$  ( $\pi_i$ ) to the probability of sampling state  $j$  ( $\pi_j$ ) is equal to one. Given that  $\pi_i$  is equal to:

$$\pi_i = \frac{Z_i(\lambda_i) \exp(\eta_i)}{Z^{EE}}, \quad (3.121)$$

Substituting Eq. 3.110 in Eq. 3.121, the following relation can be obtained for  $\pi_i/\pi_j = 1$ :

$$(\eta_i - \eta_j) = \beta(G_i - G_j). \quad (3.122)$$

Eq. (3.122) proposes that the choice of the weights is dependent on the free energies that we are attempting to obtain. This equation is then solved iteratively with trial simulations. For the first simulation, the values of  $\eta$  are set to zero, and the histogram of the states visited is obtained. With this histogram, it is possible to estimate the free energy differences and, since the weights are related to the free energies by Eq. (3.122), the next values of  $\eta$  can be calculated. This iteration goes on until a uniform

distribution is attained. The weights found are then used in a longer simulation to obtain the final solvation free energies.

The choice of the  $\lambda$  set corresponding to overlapping alchemical states are crucial to acquire accurate solvation free energies. In this work, the method chosen to obtain the optimal staging of the  $\lambda$  domain is the one developed by [Escobedo and Martinez-Veracoechea \(2007\)](#) with a basis in the study of [Katzgraber \*et al.\* \(2006\)](#). This method targets "bottlenecks" in the simulation. It does that by optimizing  $\lambda$  through the minimization of the number of round trips per CPU time between the lowest (0) and highest (1) values of  $\lambda$ . This is specifically done by maximizing the steady-state flow  $\phi$  of the simulation, which "walks" among the values of  $\lambda$ . This flow is estimated from a Fick's diffusion type of law:

$$\phi = D(\Lambda)\Pi(\Lambda)\frac{dx(\Lambda)}{d\Lambda}. \quad (3.123)$$

In the equation above,  $\Lambda$  is the actual continuous value of the coupling parameter. This continuous function of  $\lambda$ 's is obtained by interpolating the  $\lambda$  set linearly.  $D(\Lambda)$  is the diffusivity at state  $\Lambda$  and  $x(\Lambda)$  is the fraction of times that the trial simulation at state  $\Lambda_i$  has most recently visited the state  $\lambda = 1$  as opposed to state  $\lambda = 0$ . The derivative  $dx(\Lambda)/d\Lambda$  is approximated with the central finite differences method. Finally,  $\Pi(\Lambda)$  is the probability of visiting  $\Lambda$ :

$$\Pi(\Lambda) = \frac{C'\bar{\Pi}(\lambda)}{\Lambda_{i+1} - \Lambda_i}. \quad (3.124)$$

The  $C'$  term in the equation above represents a constant and  $\bar{\Pi}(\lambda)$  is the arithmetic average of the frequency of visits to the  $\Lambda$  state:

$$\bar{\Pi}_i(\lambda) = \frac{\pi_{i+1} - \pi_i}{2}. \quad (3.125)$$

The  $\phi$  is maximum when the optimal probability  $\Pi'(\Lambda_i)$  of visiting state  $\Lambda_i$  is proportional to  $1/\sqrt{D(\Lambda)}$  ([TREBST \*et al.\*, 2004](#)). With that information, it is possible to estimate the diffusivity using one trial simulation with the following equation:

$$D(\Lambda) = \frac{\Lambda_{i+1} - \Lambda_i}{\bar{\Pi}(\lambda)dx(\Lambda)/d\Lambda}, \quad \Lambda_i < \Lambda < \Lambda_{i+1}. \quad (3.126)$$

Hence, we can calculate  $\bar{\Pi}$  and, consequently, the cumulative probability ( $\Phi$ ), which is used to obtain the new  $\lambda$  states by

$$\Phi = \int_{\lambda=0}^{\lambda=1} \Pi'(\Lambda_i)d\Lambda = \frac{i}{K}, \quad (3.127)$$

where  $K$  is the total number of  $\lambda$  states. In order to carry out our solvation free energy simulations, we obtained these cumulative probabilities for every  $\lambda$  set we estimated. A graphical demonstration of the relation between the optimized coupling parameters and the cumulative probability of Eq. 3.127 is presented in Figure 3.4.1.

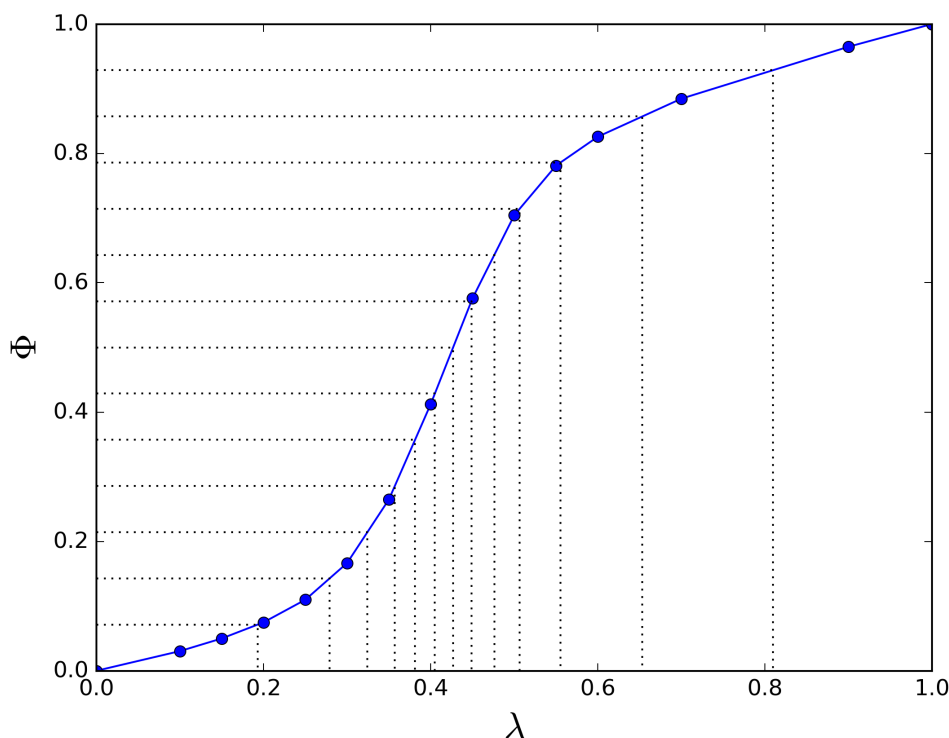


Figure 3.4.1 – Relation between the optimized coupling parameters and the cumulative probability used to obtain them.

## 3.5 Multistate Bennett Acceptance Ratio (MBAR)

We presented in the sections above the methods used to obtain the total potential energies of each alchemical state with molecular dynamics, and, in this section, we are going to discuss the methodology utilized to estimate the solvation free energies with these data. The MBAR method (SHIRTS; CHODERA, 2008) is based on the free energy perturbation. It is a maximum likelihood method which computes free energies and their uncertainties of all  $K$  states by minimizing the  $K \times K$  matrix of variances for a simulation with  $N_j$  uncorrelated samples in equilibrium. For each of the  $\{x_{i,n}\}_{n=1}^{N_i}$  configurations of  $i$ , the following probability distributions are sampled:

$$p_i(x) = \frac{q_i(x)}{c_i}, \quad (3.128)$$

$$c_i = \int dx q_i(x), \quad (3.129)$$

where  $q_i(x) = \exp[-u_i(x)]$  and  $u_i$  is the reduced potential energy of each state, defined for a an alchemical transformation by  $u_i(x) = \beta_i[U_i(x)]$ . In addition,  $c_i$  is a normalization constant. The free energies are estimated from the ratio of this constant in each state, since

$$\Delta f_{ij} = f_i - f_j = -\ln \frac{c_j}{c_i} = -\ln \frac{\int dx q_j(x)}{\int dx q_i(x)}. \quad (3.130)$$

Shirts and Chodera (2008) then proposed the following arbitrary function:

$$c_i \langle \alpha_{ij} q_j \rangle_i = c_j \langle \alpha_{ij} q_i \rangle_j. \quad (3.131)$$

Using the equation above for every state K, the following relation is obtained:

$$\sum_{j=1}^K \frac{\hat{c}_i}{N_i} \sum_{n=1}^{N_i} \alpha_{ij} q_j(x_{i,n}) = \sum_{j=1}^K \frac{\hat{c}_j}{N_j} \sum_{n=1}^{N_j} \alpha_{ij} q_i(x_{j,n}). \quad (3.132)$$

Shirts and Chodera (2008) suggested the following equation for the arbitrary term  $\alpha_{ij}$  in order to minimize the variance:

$$\alpha_{ij}(x) = \frac{N_j \hat{c}_i^{-1}}{\sum_{k=1}^K N_k c_i^{-1} q_k(x)}. \quad (3.133)$$

Assuming that the sampling is carried out following Boltzmann statistics, Eqs. (3.132) and (3.133) can be rearranged to obtain the free energy estimator, which is solved self consistently:

$$f_i = -\ln \sum_{k=1}^K \sum_{n=1}^{N_k} \frac{\exp[-u_i(x_{kn})]}{\sum_{l=1}^K N_l \exp\{[f_l - u_l(x_{kn})]\}}. \quad (3.134)$$

The equation above requires the evaluation of the potential energy of every uncorrelated configuration  $n$  for all K states  $[u_i(x_{kn})]$  and for all uncorrelated configuration snapshots ( $N_k$ ) from state  $k$ . With the free energies, we compute the free energy differences between states with Eq. 3.130. The uncertainty resulting from free energy estimation can be computed by the covariance matrix ( $s$ ):

$$\delta_{ij}^2 \Delta f_{ij} = s_{ii} + s_{jj} - 2s_{ij}. \quad (3.135)$$

The MBAR method explained here can be considered as a limiting case of the Weighted Histogram Analysis Method (WHAM) (KUMAR *et al.*, 1992) for computing free energies. WHAM equations become equal to Eq. (3.134) if the histogram width

tends to zero. Despite this, the MBAR is still more suited than the WHAM because it does not have the bias associated with the discretization and allows the calculation of an error estimate (SHIRTS; CHODERA, 2008).

## 3.6 Gibbs Ensemble Monte Carlo (GEMC)

In the initial steps of this research, we estimated the SAFT- $\gamma$  Mie force field parameters of phenanthrene with the methodology proposed by Lafitte *et al.* (2012). This approach required liquid-vapor equilibrium data obtained with molecular simulation, as it is described in Section 3.2.2. Hence, we carried out Monte Carlo simulations at the Gibbs Ensemble (PANAGIOTOPOULOS, 1987) since this ensemble is commonly used to study phase coexistence with molecular simulation. In addition to that, this method does not use an explicit interface, which can hinder the determination of bulk phase behavior of small systems with long-range interactions (RAI; MAGINN, 2012).

Before talking in more detail about this ensemble, we are going to discourse on Monte Carlo simulations briefly. The Monte Carlo (MC) approach is another method for generating atomic trajectories in order to obtain macroscopic properties. Rather than using the numerical integration of Newton's equations of motion, the trajectories are obtained stochastically in the Monte Carlo approach. The positions are evolved by random moves or perturbations (MC steps) acquired with the Metropolis method (METROPOLIS *et al.*, 1953). Hence, the trajectories are not predictable from the set of initial positions. The Metropolis method is a Markov process, that is, a stochastic process in which the configurations change randomly with time and only depends on the states and their directly preceding states, but not on the previous configurations (RAABE, 2017). The random move is constructed in such a way that the probability of visiting a particular point  $r^N$  is proportional to the Boltzmann factor  $\exp[-\beta U(r^N)]$  (FRENKEL; SMIT, 2001). The construction of a particle displacement according to Metropolis *et al.* (1953) can be briefly summed up as:

1. Pick a random particle, and calculate its energy  $U(r^N)$ .
2. Perturb the particle by randomly displacing it,  $r' = r + \Delta r$ . Where  $\Delta r$  is a perturbation randomly chosen from a defined interval of maximum displacement  $([-\delta_{max}, \delta_{max}])$ . Calculate the energy with the new positions  $U(r'^N)$ .
3. Accept the move from  $r^N$  to  $r'^N$  with the probability:

$$acc_{A \rightarrow B} = \min\{1, \exp[-\beta U(r'^N) + \beta U(r^N)]\}. \quad (3.136)$$

The values of maximum displacement are defined iteratively in order to obtain acceptance rates of 25-50% in step 3 (FRENKEL, 2013). Monte Carlo simulations are



interesting when we need to calculate properties in different thermodynamic ensembles, such as the Gibbs Ensemble used in this dissertation. The phase coexistence at this ensemble is obtained with simultaneous Monte Carlo (MC) simulations of two boxes with periodic boundary conditions, representing a two-phase system. The boxes exchange molecules, energy, and volume between them. Equilibrium is obtained through MC steps that consist of translation and rotation moves, volume exchange moves, and random exchanges of molecules between the boxes. For the phase equilibrium of multi-component systems, the GEMC simulations should be carried out at the NPT (constant number of particles, pressure, and temperature) ensemble to obey the requirement of an additional degree of freedom for mixtures. In turn, the simulation of single component systems is carried out at a constant number of particles, temperature, and volume (NVT) since the two-phase region would be a line for this system at constant pressure and temperature (FRENKEL; SMIT, 2001). The partition function of the GEMC-NVT ensemble is obtained by considering that the particles in both boxes are subjected to the same intermolecular interactions. Also, volumes and number of particles of the box ( $N_1, N_2, V_1$  and  $V_2$ ) can vary while the total volume ( $V$ ) and the total number of particles ( $N$ ) remain constant ( $N = N_1 + N_2, V = V_1 + V_2$ ). Therefore, the partition function is

$$Q(NVT) \equiv \sum_{N_1}^N \frac{1}{V \Lambda^{3N} N_1! (N - N_1)!} \int_0^V V_1^{N_1} V_2^{N_2} dV_1 \int \exp[-\beta U(x_1^{N_1})] dx_1^{N_1} \int \exp[-\beta U(x_2^{N_2})] dx_2^{N_2}. \quad (3.137)$$

In order to define the acceptance rules for the MC moves and compute any property of interest, it is necessary to know the probability of finding the configuration with  $N_1$  particles in box 1 with volume  $V_1$  and positions  $x_1^{N_1}$  and  $x_2^{N_2}$ . This probability is given by:

$$\pi(x_1^{N_1}, x_2^{N_2}, N_1, N_2, V_1, V_2) \propto \frac{V_1^{N_1} V_2^{N_2}}{N_1! N_2!} \exp[-\beta U(x_1^{N_1}) + \beta U(x_2^{N_2})]. \quad (3.138)$$

The acceptance criterion for the translation and rotation moves from configuration A to configuration B is similar to the conventional NVT MC method and is equal to:

$$acc_{A \rightarrow B} = \min\{1, \exp[\beta U(x_A^{N_1}) - \beta U(x_B^{N_1})]\}. \quad (3.139)$$

The volume exchange moves take place by exchanging an amount  $\Delta V$  between the boxes to achieve pressure equilibrium.  $\Delta V$  can be chosen from a uniform distribution

based on the maximum variation of volume ( $\delta V_{max}$ ) defined with probability  $1/\delta V_{max}$  (FRENKEL; SMIT, 2001). The acceptance rule for these moves is:

$$acc_{A \rightarrow B} = \min \left\{ 1, \left( \frac{V_1^B}{V_1^A} \right)^{N_1+1} \left( \frac{V_2^B}{V_2^A} \right)^{N_2+1} \exp[\beta U(x_A^N) - \beta U(x_B^N)] \right\}. \quad (3.140)$$

Particle exchange moves are carried out to obtain the equality of chemical potential between the boxes. One particle from one box is removed and then added to a random location in the other box. The criteria to accept or reject this type of move is:

$$acc_{A \rightarrow B} = \min \left\{ 1, \frac{N_1 V_2}{(N_2 + 1) V_1} \exp[\beta U(x_A^N) - \beta U(x_B^N)] \right\}. \quad (3.141)$$

This method has been widely used to calculate phase equilibrium, but its performance is poor for the region near the critical point due to large density fluctuations. The GEMC method also has poor performance for dense systems since the particle exchange moves have a low acceptance rate (WESTMORELAND *et al.*, 2002).

## 4 Methodology

In this study, we had first to obtain the parameters of phenanthrene using an equation for rings for the SAFT- $\gamma$  Mie force field since these parameters were not available on this force field database (ERVIK *et al.*, 2016b). Hence, we divided this chapter into two sections. The first one describes how we parametrized the phenanthrene molecule with the SAFT- $\gamma$  Mie force field and the second one explains how we carried out the solvation free energy simulations.

### 4.1 Phenanthrene Parameterization

We implemented the two parameterization strategies for molecules with aromatic rings described in Section 3.2.2 for phenanthrene. For both of them, only vapor pressure data (MORTIMER; MURPHY, 1923; OSBORN; DOUSLIN, 1975) were used due to the unavailability of saturated liquid density. We did not estimate the attractive exponent,  $\lambda_a$ . Instead, the value of six was given to it, as recommended by Ramrattan *et al.* (2015) and Herdes *et al.* (2015), due to its high correlation with the repulsive exponent. The parameterization with the ring equation of Müller and Mejía (2017) was carried out with the number of segments equal to five and with a geometry such as that in Figure 4.1.1, since this level of coarse-graining was also used for a similar molecule (anthracene) in the original paper.

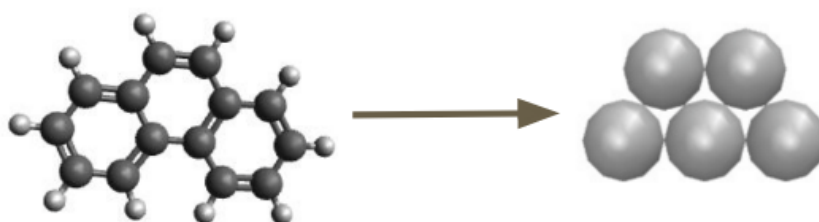


Figure 4.1.1 – Representation of phenanthrene with the geometry proposed by Müller and Mejía (2017).

The minimization was done using the Particle Swarm Optimization (PSO) method (SCHWAAB *et al.*, 2008) with 50 particles and 50 interactions. The objective function used was equal to

$$F_{obj} = \sum_{i=1}^{N_p} \left[ \frac{P_v^{SAFT}(T_i, \sigma, \epsilon, \lambda_r) - P_v^{exp}(T_i)}{P_v^{exp}(T_i)} \right]^2. \quad (4.1)$$

Here,  $P_v^{exp}$  is the experimental vapor pressure and  $P_v^{SAFT}$  is the vapor pressure

obtained with the SAFT-VR Mie EoS. According to the fundamental thermodynamic equation, the vapor pressure is given by

$$P_v^{SAFT} = -\frac{\partial A^{SAFT}}{\partial V} \quad (4.2)$$

This derivative and all the derivatives required to calculate  $A^{SAFT}$ , such as the ones in Eqs. A.24 and A.26, were obtained numerically with the finite difference method. The routine used to calculate the bubble point with the EoS was the one proposed by Smith *et al.* (2007). The parameters ( $\sigma$ ,  $\epsilon$ , and  $\lambda_r$ ) from the minimization of the objective function in Eq. (4.1) are the final force field parameters used in molecular simulations.

The parameterization with the ring equation the Lafitte *et al.* (2012) was carried out with  $m_s = 3$ , so that every bead would represent one aromatic ring, as depicted in Figure 4.1.2.

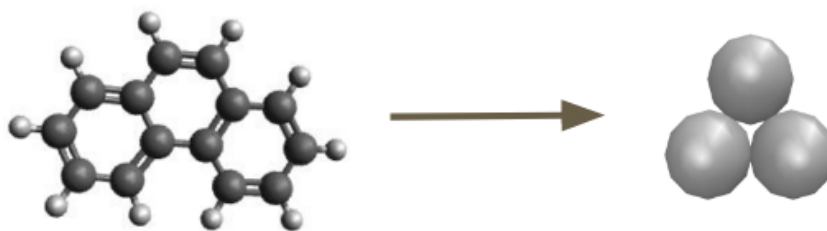


Figure 4.1.2 – Representation of phenanthrene with the geometry proposed by Lafitte *et al.* (2012).

The first part of the estimation followed the same procedure described above for the Müller and Mejía (2017) equation. However, as explained in Section 3.2.2, the Lafitte *et al.* (2012) equation requires the estimation of correction factors  $c_\sigma$  and  $c_\epsilon$  (Eqs. (3.91) and (3.92)). We then estimated these parameters by using the PSO method with Eq. (3.93). In this equation, vapor pressures and saturated liquid densities from molecular simulations are required. We then decided to use the Gibbs Ensemble Monte Carlo method on the NVT ensemble, explained in Section 3.6, to obtain these equilibrium properties at eight different temperatures.

The boxes for the GEMC-NVT simulations were generated by inserting 400 molecules of phenanthrene into one liquid box and 100 molecules of phenanthrene into one vapor box using the Playmol package (ABREU, 2017), which is integrated with the Packmol package (MARTÍNEZ *et al.*, 2009). Initial densities of each box were made equal to the saturated densities found with the SAFT-VR Mie Eos, aiming at avoiding the migration of all molecules to a single phase during the simulation. The GEMC-NVT simulations were executed using the Cassandra software (SHAH; MAGINN, 2011), which was developed to perform Monte Carlo simulations. The equilibration and production times lasted around  $10^4$  and  $5 \times 10^4$  MC cycles, respectively. Each MC cycle

corresponded to  $10^3$  rotation trials,  $10^3$  translation trials,  $10^2$  molecule insertion trials,  $10^2$  molecule deletion trials, and 10 volume exchange trials. The cut-off distance was equal to 20 Å and we did not use long-range interactions. The saturated vapor density ( $\rho_{vap}$ ), the saturated liquid density ( $\rho_{liq}$ ), and the vapor pressure ( $P_v$ ) were sampled at each 100 MC cycles. Later on, these data were divided into five blocks for calculation of their averages and standard deviations. With the correction factors found after the estimation with the simulation data, we calculated, with Eqs. (3.94) and (3.95), the  $\sigma$  and  $\epsilon$  parameters. Lafitte *et al.* (2012) proposes that these are the final parameters to be used in molecular simulations. Hence, an iterative simulation is not required, and the set of optimal parameters can be obtained with one group of molecular simulations.

## 4.2 Solvation Free Energy Simulations

Using the parameters for phenanthrene estimated with the Müller and Mejía (2017) approach and the SAFT- $\gamma$  Mie force field parameters available for other compounds, we carried out molecular dynamics simulations to estimate the solvation free energies. The chosen software package to perform the simulations was LAMMPS (PLIMPTON, 1995). In this package, the equations of motion were integrated with the velocity-Verlet algorithm (VERLET, 1967) with a time step of 2 fs. As required by the coarse-grained model, molecules with more than one bead were treated as rigid bodies. The thermostat and the barostat were the Nosé-Hoover chains as described in Hoover (1985), Martyna *et al.* (1992), and Martyna *et al.* (1994) with damping factors of 100 and 1000 time steps, respectively. For the rigid bodies in our simulations, we used the rigid-body algorithm of Kamberaj *et al.* (2005). Electrostatics interactions are not explicitly accounted for by the SAFT- $\gamma$  Mie force field. Hence, we did not compute Coulombic interactions. The potential cutoff was equal to 20 Å (MÜLLER; MEJÍA, 2017) with a neighbor list skin of 2 Å. The initial configurations of the solvated systems were also generated using the Playmol package, which is integrated with the Packmol package. For the binary mixtures, one molecule of solute and a varying number of solvent molecules- 700 molecules of toluene, 700 molecules of octanol, 1024 molecules of hexane, 3000 molecules of water - were randomly added to a cubic box. Besides the systems with pure substances acting as solvents, we performed simulations to study the solvation free energy of phenanthrene in a mixture of toluene and carbon dioxide with different weight fractions ( $w_{CO_2}$ ). The system consisted of one molecule of phenanthrene for all the cases and 123 molecules of  $CO_2$  and 618 molecules of toluene ( $w_{CO_2} = 0.087$ ); 166 molecules of  $CO_2$  and 589 molecules of toluene ( $w_{CO_2} = 0.119$ ); 232 molecules of  $CO_2$  and 545 molecules of toluene ( $w_{CO_2} = 0.169$ ); 380 molecules of  $CO_2$  and 446 molecules of toluene ( $w_{CO_2} = 0.289$ ). These substances used in our study were selected with the intention of testing the force field with standard sets used as

a benchmark in solvation free energy calculations, with aromatic substances used as models to asphaltenes and with water, which probably is the most used solvent in computational studies of solvation free energies.

All simulations were performed with the constant temperature and pressure values of 298 K and 1 bar, except the ones containing carbon dioxide. These had the temperature of 298 K and the pressure of the experimental liquid-phase equilibrium corresponding to each composition of the system  $CO_2$ +toluene (CHANG, 1992). For all simulations, the initial box was equilibrated at the NPT ensemble for 2 ns, and the resulting configurations were used as the initial configuration of the expanded ensemble simulations. These were carried out with the LAMMPS user package for expanded ensemble simulations with the Mie potential developed by our research group, available at <https://github.com/atoms-ufrj/USER-ALCHEMICAL>.

During these expanded ensemble simulations, the sampling of a new alchemical state was tried at every 10 MD steps. To define the optimal values of  $\lambda$  and  $\eta$  corresponding to each state, trial simulations, having around 9 ns of production time, were carried out. In the first simulation, we chose the group of  $\lambda$  values arbitrarily, and we either set all  $\eta$ 's to zero or assigned values previously found for similar solute-solvent pairs. The subsequent group of  $\eta$ 's were estimated with the flat histogram approach (Eq. (3.122)). We then performed another trial simulation with the new weights. The results of this simulation were used to optimize the group of  $\lambda$ 's by minimizing the number of round trips, as described in Section 3.4. The  $\eta$ 's corresponding to the newest group of  $\lambda$ 's were interpolated linearly from the free energy differences. With the final values of  $\eta$  and  $\lambda$  defined for each mixture, larger simulations with a production time of 20 ns were carried out.

Since the employed force field considers that the beads do not have charges, there are no Coulombic interactions and the  $\Delta G$  in Eq. (3.111) becomes equal to  $\Delta G_{3 \rightarrow 4}$ . The post-processing method used to effectively calculate free energy differences with the potential energies obtained from the expanded ensemble simulations was the Multistate Bennett Acceptance Ratio (MBAR) method, described in Section 3.5. The software alchemical-analysis (KLIMOVICH *et al.*, 2015) was utilized to obtain the  $\Delta G_{solv}$  with MBAR and to assess the quality of the results. After the first estimations, we realized that the binary interaction parameter of Eq. (3.89) was necessary for systems containing water. Hence, we estimated  $k_{ij}$  for these pairs and, for all the other pairs, we set  $k_{ij}$  to zero. The estimation was done by performing trial expanded ensemble simulations in three values of  $k_{ij}$ , as suggested by Ervik *et al.* (2016a). With the  $\Delta G_{solv}$  obtained with these simulations, we did a linear fit to obtain the refined value of the parameter. We used this strategy because the estimation with SAFT-VR Mie EoS gave poor results for the hydration free energies.

# 5 Results and Discussion

## 5.1 Phenanthrene parameterization

The first part of this work consisted in obtaining phenanthrene parameters for the SAFT- $\gamma$  Mie Force Field as described in Section 4.1. This part was necessary since these parameters were not available for the ring geometry on the force field database (ERVIK *et al.*, 2016b). We carried out this parameterization using the free energy of Helmholtz equations for rings of (LAFITTE *et al.*, 2012), Eq. 3.84, and of (MÜLLER; MEJÍA, 2017), Eq. 3.85. The parameters obtained with these two strategies and the mean percentage error (MPE) to the experimental data (MORTIMER; MURPHY, 1923; OSBORN; DOUSLIN, 1975) were those observed in Table 5.1.1.

Table 5.1.1 – Estimated SAFT- $\gamma$  Mie force field parameters for phenanthrene.

$m_s$	$\epsilon/\kappa_b$ (K)	$\sigma$ (Å)	$\lambda_r$	MPE(%)
3 (LAFITTE <i>et al.</i> , 2012)	485.55	4.197	14.34	1.48   9.74
5 (MÜLLER; MEJÍA, 2017)	262.74	4.077	9.55	0.88

The MPE value of 1.64 for the Lafitte *et al.* (2012) strategy in the Table 5.1.1 is the error between the vapor pressure calculated with the equation of state and the experimental data. The second MPE value for the Lafitte *et al.* (2012) strategy (9.74) is the error between the vapor pressure calculated with the equation of state and the vapor pressure obtained in the GEMC simulations. We also show, in Figure 5.1.1, the vapor pressure curve obtained using the EoS with the two Helmholtz free energy equations for rings. It is clear, observing the figure mentioned above and the MPE values for both equations for rings, that both approaches for the EoS performed very similarly and were able to reproduce the experimental data. However, the parameters estimated with the equation of Lafitte *et al.* (2012) could not be directly transferred to molecular simulations because this strategy needs an estimation with molecular simulation. This additional procedure is not necessary when estimating parameters for the chain equation (AVENDAÑO *et al.*, 2011) or the ring equation of Müller and Mejía (2017). In addition to that, this use of molecular simulation data to acquire the parameters negates the overall idea proposed by (AVENDAÑO *et al.*, 2011). They developed this force field with the intention of obtaining the parameters in a more straightforward way than other force fields since the SAFT- $\gamma$  Mie model would not have the computational time associated with doing molecular simulations in its parameterization. Due to these specific characteristics of the model of Lafitte *et al.* (2012), we only studied the solvation free energy of phenanthrene with the set of parameters estimated with the strategy of Müller and Mejía (2017). In



fact, we only followed the approach of Lafitte *et al.* (2012) because it was the only one available when we first started this research. The sets of parameters and geometries for the other compounds were retrieved from the literature, and they are displayed in Tables 3.2.1 and 3.2.2.

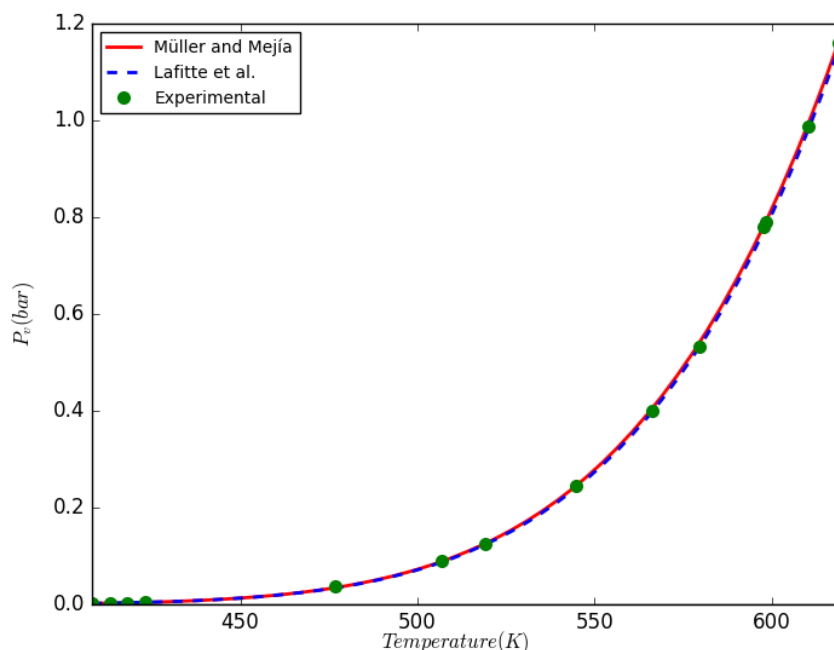


Figure 5.1.1 – Vapor pressure ( $P_v$ ) of phenanthrene calculated with the two strategies of the SAFT-VR Mie EoS for modeling molecules formed by rings.

## 5.2 Solvation free energies

Our primary intention with this study is to assess the capability of the SAFT- $\gamma$  Mie force field to represent solvation free energies. Hence, we chose benchmark solutes used in the literature (benzene, propane) and polyaromatic solutes (pyrene, phenanthrene, and anthracene), and, for the solvents, we picked non-polar (hexane), aromatic (toluene), and hydrogen bonding (1-octanol, water) substances. It would be interesting to do a study with a bigger database of pairs solvent-solute. However, the time required for performing each of the solvation free energy simulations, some difficulties related to the available computational structure, and the fact that a better model of aromatic compounds with this force field was only published in the middle of our study prevented us from doing a more extensive study. The solvation free energy simulations for the pairs chosen were carried out with binary interaction parameters equal to zero since these parameters were not necessary according to our preliminary studies. Since the force field does not account for charges, we only calculated the Mie contribution (Eq. (3.113)) to the solvation free energy. A total of 15 to 18  $\lambda$ 's, depending on the solute-solvent



pairs, and their respective  $\eta'$ s were estimated as described in Sections 3.4 and 4.2. The simulations carried out using these optimized weights deviated from the requirement of equal number visits by an average of 5% for all of the solvent+solute pairs. The final  $\lambda$  set for all the pairs was found using the cumulative probability distribution (Eq. (3.127)). The probability distribution for the hexane(solvent)+benzene(solute) pair can be seen in Figure 5.2.1. From now on we are going to use the terminology solvent+solute. The optimized values of  $\lambda$  and  $\eta$  for this pair and all the other pairs are available in Tables 5.2.1 to 5.2.4. By observing the coupling parameters found for all the pairs, we can see that they are concentrated on the region with a steeper slope as it is expected in this method.

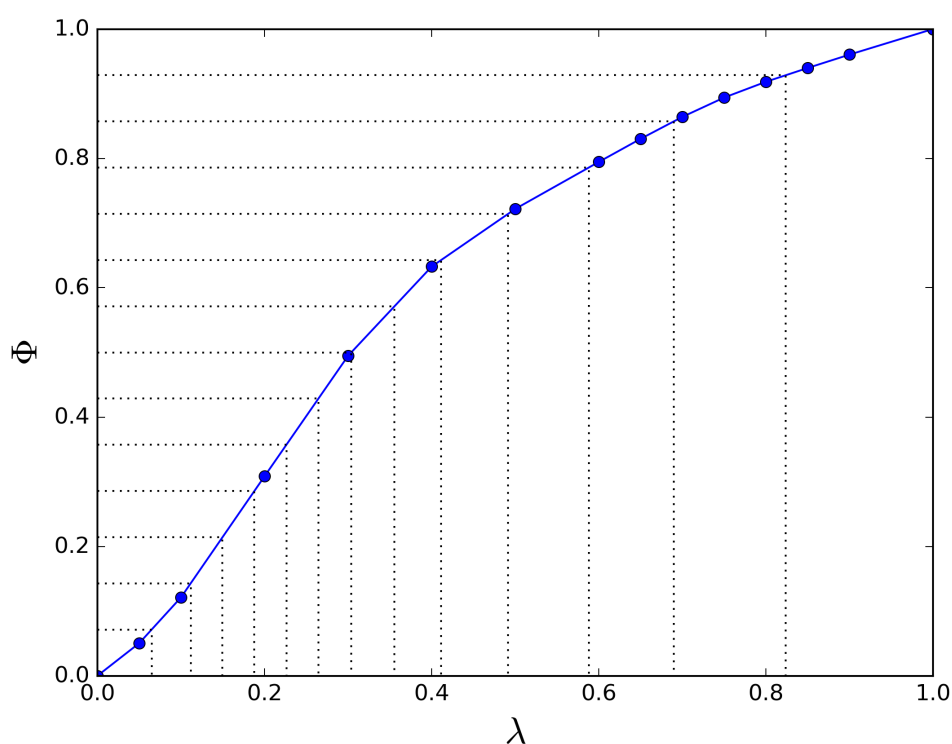


Figure 5.2.1 – Cumulative probability used to obtain the optimized values of  $\lambda$ 's for the pair hexane+benzene.

Table 5.2.1 – Optimized values of  $\lambda$  and  $\eta$  for the hexane+solute pairs.

benzene		pyrene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000
0.065	0.708	0.076	4.234	0.090	1.981
0.112	1.385	0.107	5.620	0.132	3.461
0.15	1.892	0.132	6.499	0.161	4.494
0.188	2.399	0.152	6.690	0.185	5.185
0.226	2.519	0.170	6.643	0.205	5.552
0.264	2.457	0.189	6.461	0.224	5.725
0.304	2.367	0.213	6.091	0.244	5.722
0.356	1.921	0.242	5.566	0.268	5.523
0.411	1.411	0.280	4.729	0.305	4.975
0.492	0.524	0.355	2.853	0.372	3.576
0.588	-0.663	0.483	-0.778	0.500	0.297
0.69	-2.016	0.678	-6.947	0.560	-1.390
0.824	-3.922	0.788	-10.631	0.722	-6.309
1.000	-6.583	1.000	-18.141	1.000	-15.448

Table 5.2.2 – Optimized values of  $\lambda$  and  $\eta$  for the 1-octanol+solute pairs.

propane		anthracene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000
0.027	3.126	0.078	3.932	0.049	2.578
0.050	5.109	0.111	6.178	0.091	5.663
0.073	6.093	0.130	7.426	0.125	8.575
0.095	6.570	0.143	8.201	0.144	10.069
0.117	6.826	0.154	8.717	0.157	10.978
0.142	6.956	0.164	9.085	0.169	11.599
0.174	6.969	0.174	9.357	0.180	12.040
0.215	6.847	0.184	9.556	0.192	12.340
0.269	6.554	0.197	9.676	0.206	12.499
0.337	6.050	0.214	9.681	0.225	12.478
0.427	5.228	0.238	9.490	0.253	12.161
0.545	3.955	0.274	8.958	0.298	11.280
0.720	1.843	0.326	7.906	0.371	9.406
1.000	-1.903	0.399	6.088	0.484	5.891
		0.515	2.777	0.664	-0.516
		0.695	-2.960	0.802	-5.908
		1.000	-13.768	1.000	-14.073

Table 5.2.3 – Optimized values of  $\lambda$  and  $\eta$  for the toluene+solute pairs.

pyrene		anthracene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000
0.090	2.563	0.119	0.218	0.136	0.726
0.130	4.338	0.174	1.210	0.191	2.307
0.154	5.439	0.209	2.052	0.223	3.430
0.172	6.181	0.236	2.664	0.246	4.233
0.188	6.670	0.261	3.122	0.264	4.780
0.204	6.986	0.283	3.378	0.281	5.149
0.222	7.121	0.306	3.449	0.299	5.354
0.244	7.025	0.332	3.311	0.318	5.389
0.278	6.520	0.360	2.936	0.340	5.222
0.340	5.010	0.399	2.209	0.372	4.717
0.462	1.247	0.466	0.567	0.425	3.440
0.616	-4.283	0.564	-2.211	0.524	0.444
0.788	-11.032	0.715	-6.983	0.701	-5.814
1.000	-19.814	1.000	-16.923	1.000	-17.803

Table 5.2.4 – Optimized values of  $\lambda$  and  $\eta$  for the phenanthrene+ $CO_2$ + toluene mixture with different values of  $w_{CO_2}$ .

$w_{CO_2} = 0.087$		$w_{CO_2} = 0.119$		$w_{CO_2} = 0.169$		$w_{CO_2} = 0.289$	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
0.128	0.604	0.128	0.732	0.064	0.883	0.066	0.806
0.184	2.067	0.186	2.223	0.108	0.764	0.111	0.760
0.217	3.164	0.219	3.319	0.175	1.969	0.172	1.983
0.240	3.940	0.244	4.098	0.214	3.156	0.204	2.967
0.260	4.472	0.267	4.704	0.240	3.974	0.227	3.627
0.277	4.823	0.289	5.031	0.258	4.457	0.245	4.082
0.295	5.035	0.313	5.084	0.273	4.750	0.262	4.395
0.318	5.059	0.339	4.950	0.287	4.921	0.279	4.583
0.347	4.762	0.373	4.371	0.305	4.962	0.299	4.621
0.397	3.753	0.425	3.055	0.326	4.885	0.325	4.423
0.491	1.031	0.488	1.196	0.361	4.401	0.365	3.739
0.670	-5.148	0.525	-0.027	0.419	2.990	0.428	2.198
0.791	-9.713	0.730	-7.185	0.527	-0.299	0.530	-0.842
1.000	-18.098	1.000	-17.769	0.697	-6.180	0.701	-6.763
				1.000	-17.998	1.000	-18.163

It is also essential to analyze the reliability of solvation free energy estimations through the overlapping of the intermediate states. Insufficient overlap among states when using FEP based methods such as MBAR may result in the underestimation of variance and, consequently, in substantially incorrect solvation free energies (KLIMOVICH *et al.*, 2015). The overlap matrix for the solvation free energy of benzene in hexane is presented in Figure 5.2.2 and the matrices for the other pairs are available in Appendix B. Each element  $ij$  of these matrices is the average probability of observing a configuration sampled from state  $i$  in state  $j$ . As an example, the average probability of finding a configuration sampled from state 3 in state 4 is 0.11 in Figure 5.2.2. According to Klimovich *et al.* (2015), a tridiagonal overlap matrix is an indication of reliable free energy estimates, as long as the resulting error is sufficiently low. They define a tridiagonal matrix as one matrix with elements appreciably different from zero (the values should be as low as 0.03) in the main diagonal and the first diagonals above and below the main one. This requirement was met for all the pairs in our study. Some of the overlap matrices, including the one in Figure 5.2.2, had more than three diagonals, and, consequently, an apparent unnecessary number of intermediate states. However, this number of intermediate states was indispensable in our study because the error estimate of the solvation free energies significantly increased when we removed some of the intermediate states. Hence, we maintained these intermediate states in order to obtain low error values. After this analysis, we present in Table 5.2.5 the results for solvation free energy calculations, the errors from these estimations with MBAR, and the absolute deviations from experimental data (KATRITZKY *et al.*, 2003).

Table 5.2.5 – Calculated and experimental values for the solvation free energies (kcal/mol) of solutes in non-aqueous solvents.

Solute	Solvent	$\Delta G_{solv}^{exp}$	$\Delta G_{solv}^{Mie}$	Absolute Deviation
benzene	hexane	-3.96	-3.76 $\pm$ 0.01	0.20
pyrene	hexane	-11.53	-10.82 $\pm$ 0.02	0.71
phenanthrene	hexane	-10.01	-9.16 $\pm$ 0.01	0.85
propane	1-octanol	-1.32	-1.36 $\pm$ 0.03	0.04
anthracene	1-octanol	-11.72	-8.12 $\pm$ 0.03	3.61
phenanthrene	1-octanol	-10.22	-8.34 $\pm$ 0.03	1.47
pyrene	toluene	-12.86	-11.74 $\pm$ 0.01	1.11
anthracene	toluene	-11.31	-9.90 $\pm$ 0.01	1.41

$\lambda$	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14
0	.42	.29	.16	.08	.03	.01	.01								
1	.33	.27	.18	.10	.06	.03	.02	.01							
2	.23	.23	.18	.12	.09	.06	.04	.02	.01	.01					
3	.14	.16	.16	.13	.11	.10	.08	.06	.03	.02	.01				
4	.06	.09	.11	.11	.12	.12	.13	.11	.07	.04	.02	.01	.01		
5	.02	.03	.05	.07	.09	.12	.16	.16	.12	.08	.05	.02	.01	.01	
6		.01	.02	.03	.06	.09	.15	.18	.16	.12	.08	.04	.02	.01	
7			.01	.02	.03	.07	.13	.17	.18	.16	.11	.07	.04	.02	.01
8				.01	.02	.04	.09	.14	.17	.18	.15	.10	.06	.03	.01
9					.01	.02	.06	.11	.15	.18	.17	.13	.09	.05	.02
10						.01	.04	.07	.11	.16	.18	.17	.14	.08	.03
11						.01	.02	.04	.08	.12	.17	.19	.19	.14	.06
12							.01	.02	.05	.08	.13	.18	.22	.19	.11
13							.01	.01	.03	.05	.09	.15	.22	.25	.19
14								.01	.01	.03	.06	.11	.19	.29	.30

Figure 5.2.2 – Overlap matrix for hexane+benzene.

The numerical values for solvation free energies in hexane had overall smaller absolute deviations from experimental data than the deviations in the other solvents. Additionally, this force field presented better results for the pair hexane+benzene than the TraPPE force field ( $-4.35 \pm 0.05$  kcal/mol) (GARRIDO *et al.*, 2011) and the ELBA coarse-grained force field ( $-2.92 \pm 0.01$  kcal/mol) (GENHEDEN, 2016). TraPPE is a force field parametrized with fluid-phase equilibria data that uses the Lennard-Jones potential to describe the non-bonded interactions. In the cited paper, they used the united-atom description of the TraPPE force field for the alkyl group, the all-atom description for the polar groups and the explicit-hydrogen approach for the aromatic groups. In the explicit-hydrogen approach, the interaction sites for all hydrogen atoms, some lone pair electrons, and bond centers are accounted for (RAI; SIEPMANN, 2007). In turn, the ELBA force field is a coarse-grained model that comprises six independent parameters. This force field models three carbons as one Lennard-Jones site and one water molecule as a single Lennard-Jones site with a point dipole. We also present the solvation free energies corresponding to each alchemical state ( $\lambda$ ) for all the pairs studied here in Figures 5.2.3 to 5.2.5. In these figures, we did not represent the errors of the estimation with MBAR because they were too small. Specifically observing the solvation free energy in hexane (Figure 5.2.3), we can see the effect of the molecule's

size on the entropic region of the free energy curve. This region corresponds to the first values of  $\lambda$  where the cavity required for the insertion of the solute is formed.

We expected that a force field based on an EoS that does not explicitly account for hydrogen bond would not perform well for 1-octanol in mixtures since the parameterization of this molecule did not explicitly account for the interactions of association. All the beads representing 1-octanol have the same intermolecular parameter, and there is no distinction between the polar and apolar groups. Despite this, the solvation free energies of propane and phenanthrene in 1-octanol lied in the desired deviation range of 1-2 kcal/mol (MOBLEY; GILSON, 2017). For propane, the observed deviation in solvation free energies was much smaller when compared to the other solutes, which can be attributed to the non-polarity of propane and its smoother free energy curve, presented in Figure 5.2.4. Such solvation free energy of propane in 1-octanol also had a smaller deviation than the prediction of the ELBA force field ( $-0.92 \pm 0.01$ ) (GENHEDEN, 2016). The absolute deviation of the solvation free energy computed for anthracene in 1-octanol is much higher than the one calculated for phenanthrene in 1-octanol. The anthracene and phenanthrene molecules have the same geometry (Figure 4.1.1) in the SAFT- $\gamma$  Mie model, although anthracene is a linear molecule and phenanthrene is not, and also similar physical properties. Hence, this high deviation of the solvation free energy of anthracene in 1-octanol may indicate a problem in the geometry chosen for anthracene in the SAFT- $\gamma$  Mie force field and the importance of the geometry in modeling the molecules with this force field.

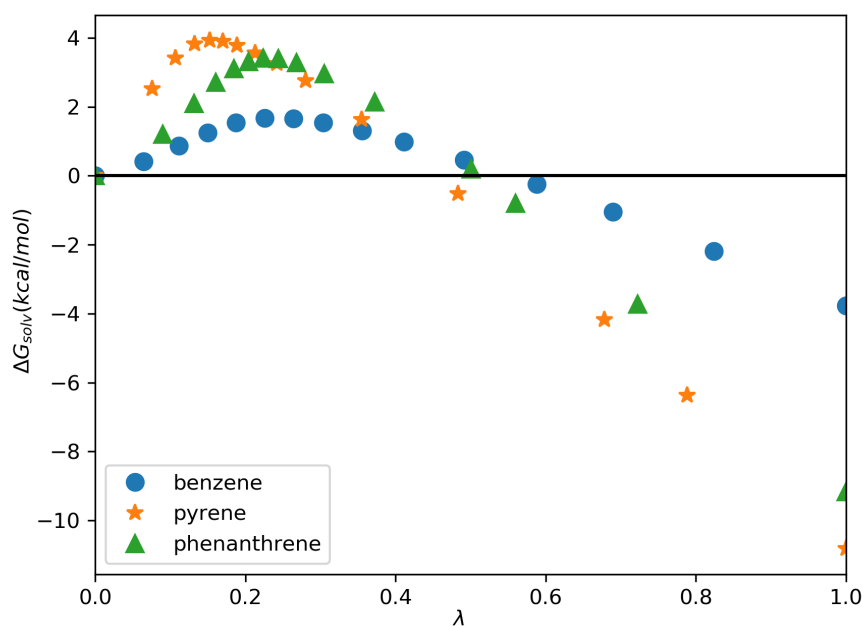


Figure 5.2.3 – Representation of the solvation free energies of different solutes in hexane estimated at each alchemical state.

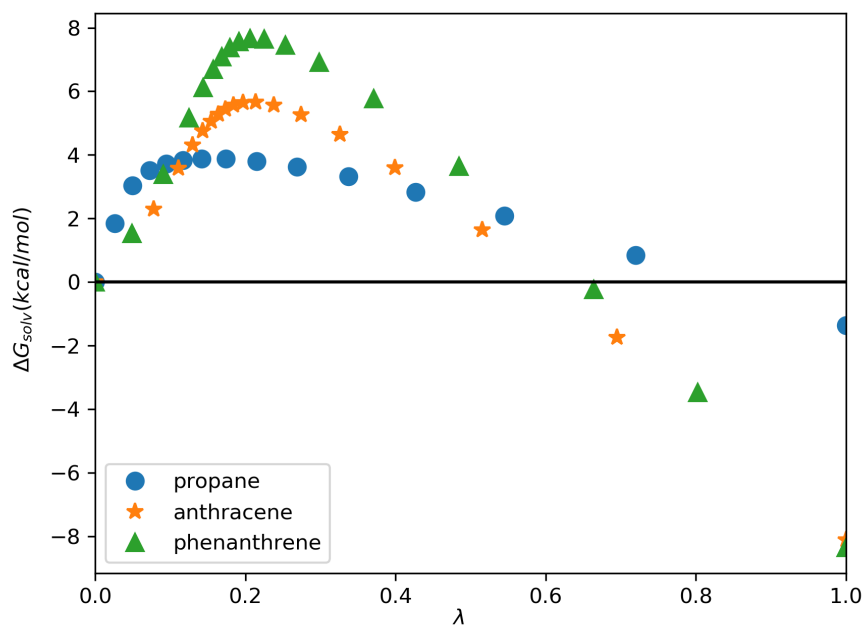


Figure 5.2.4 – Representation of solvation free energies of different solutes in 1-octanol estimated at each alchemical state.

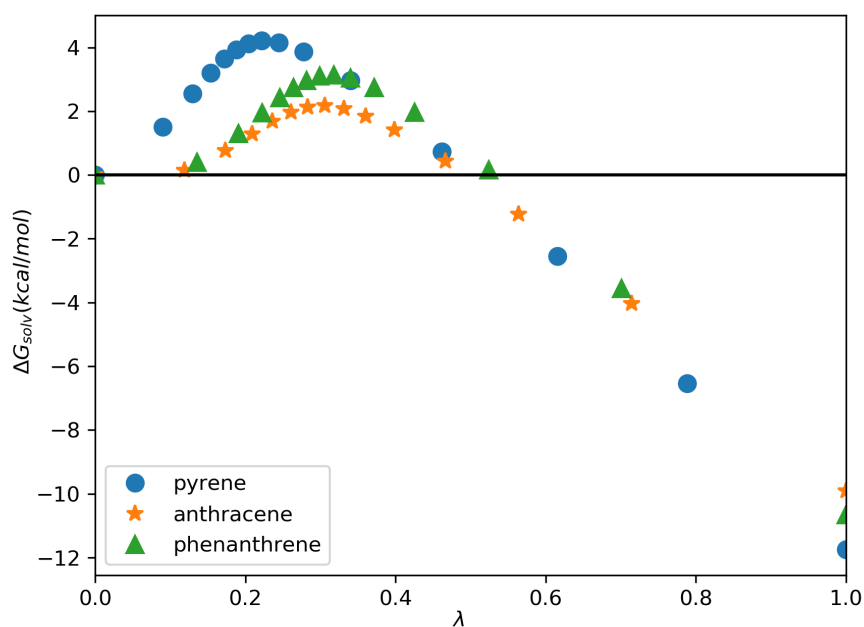


Figure 5.2.5 – Representation of solvation free energies of different solutes in toluene estimated at each alchemical state.

The results also indicate a reasonable capability of the force field for predicting the solvation free energies of polyaromatic solutes in aromatic solvents. The influence

of the molecular geometry on the solvation free energy curves was the same as the one observed for other solvents, as can be seen in Figure 5.2.5. We also calculated the  $\Delta G_{solv}$  for phenanthrene in pure toluene and in toluene+ $CO_2$  mixtures. To the best of our knowledge, there are no available experimental data for these solvation free energies, but the previous results for phenanthrene in other solvents showed that the force field is adequate to describe the solvation phenomenon of phenanthrene in a pure aromatic solvent. Hence, we decided to carry out a qualitative study of the influence of  $CO_2$  in the solvation free energies of phenanthrene in toluene in order to evaluate the description of this system with the SAFT- $\gamma$  Mie force field. The results for these sets are exposed in Table 5.2.6.

Table 5.2.6 – Calculated values for the solvation free energies (kcal/mol) of phenanthrene in toluene+ $CO_2$ .

$w_{CO_2}$	$\Delta G_{solv}^{Mie}$
0.0	$-10.65 \pm 0.02$
0.087	$-10.73 \pm 0.02$
0.119	$-10.78 \pm 0.02$
0.169	$-10.71 \pm 0.02$
0.289	$-10.69 \pm 0.02$

The increase of the mass fraction of  $CO_2$  in toluene caused a small effect on the solvation free energies in the range of weight fractions (0.087-0.289) studied in this dissertation. First, the  $\Delta G_{solv}$  decreased with the increase of  $w_{CO_2}$  up to 0.119. After this, the effect was reversed, and carbon dioxide became an anti-solvent. Soroush *et al.* (2014) reported that asphaltene precipitation occurs when carbon dioxide mass fractions became higher than 0.10 in the system asphaltene+toluene+carbon dioxide, which is in agreement with the anti-solvent effect of carbon dioxide observed in the values calculated here. In the Figure 5.2.6, we present the free energy profiles of the solvation free energies in the toluene +  $CO_2$  mixtures. Although we noticed the anti-solvent effect, the differences observed are pretty small. These minor differences may indicate that the effect of  $CO_2$  is negligible in the solvation of phenanthrene in toluene when using the SAFT- $\gamma$  Mie force field to model the molecules. Nevertheless, more studies need to be done to make a safe assertion about it. It is also worth remarking that this is a qualitative study due to the lack of experimental data. Overall, the methodology proposed by the SAFT- $\gamma$  Mie force field was satisfactory in predicting the solvation free energies of the pairs solvent-solute studied here. For the pair 1-octanol+anthracene, the performance obtained was not as good as it was for the other pairs. This result highlights the importance of choosing a correct geometry for this coarse-grained force field.



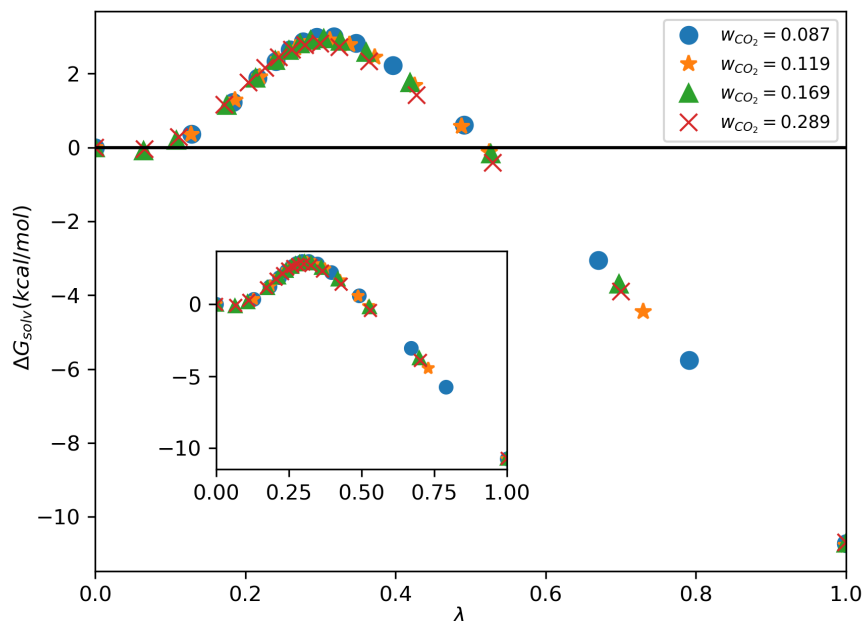


Figure 5.2.6 – Representation of solvation free energies of phenanthrene in toluene+ $CO_2$  estimated at each alchemical state.

### 5.3 Hydration free energies

Water is a solvent extensively used in experimental and computational studies. Because of this importance and the fact that water has unique properties, such as density maximum at 277 K and increased diffusivity upon compression, developing an accurate computational model for water is an ongoing quest (HADLEY; MCCABE, 2012). Hence, we also calculated the solvation free energies in water (hydration free energies) with the SAFT- $\gamma$  Mie force field. With these calculations, we intend to verify if this coarse-grained model would represent the water molecule correctly and would be a good alternative to decrease the computational cost of solvation studies with asphaltene models. The simulations with water as a solvent were carried out using widely studied solutes (propane) and aromatic solutes (benzene, toluene, and phenanthrene) with a set of fifteen intermediate states. We obtained these sets of  $\lambda$  and  $\eta$  with the same methodology used to acquire the sets for the solvation free energies with non-aqueous solvents, and they are exposed in Table 5.3.1. At first in our simulations, the binary interaction parameters of all aqueous mixtures were set to zero, but preliminary results for hydration free energies, displayed in Table 5.3.2, exhibited a high deviation from experimental data (ABRAHAM *et al.*, 1990; RIZZO *et al.*, 2006).

Table 5.3.1 – Optimized values of  $\lambda$  and  $\eta$  for the water+solute pairs.

propane		benzene		toluene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
0.107	2.673	0.193	-0.295	0.177	0.182	0.142	-2.462
0.157	4.703	0.279	1.468	0.262	2.432	0.256	0.597
0.186	6.047	0.324	2.931	0.307	4.244	0.319	4.504
0.210	7.148	0.357	4.168	0.336	5.552	0.358	7.762
0.230	8.017	0.381	5.091	0.360	6.696	0.384	10.104
0.250	8.883	0.405	5.891	0.380	7.558	0.407	12.185
0.272	9.291	0.427	6.443	0.400	8.233	0.427	13.607
0.294	9.700	0.449	6.770	0.422	8.678	0.446	14.490
0.328	9.900	0.476	6.900	0.443	8.859	0.469	14.834
0.381	9.930	0.506	6.805	0.473	8.810	0.494	14.667
0.484	9.463	0.555	6.392	0.514	8.452	0.533	13.832
0.623	8.195	0.653	5.109	0.606	7.148	0.620	11.069
0.781	6.378	0.810	2.421	0.755	4.273	0.806	3.279
1.000	3.333	1.000	-1.480	1.000	-1.547	1.000	-6.122

Table 5.3.2 – Calculated values using  $k_{ij} = 0$  and experimental values for the hydration free energies (kcal/mol) of solutes in water.

Solute	$\Delta G_{solv}^{exp}$	$\Delta G_{solv}^{Mie}$	Absolute Deviation
propane	$2.00 \pm 0.20$	$1.10 \pm 0.01$	0.90
benzene	$-0.86 \pm 0.20$	$-4.45 \pm 0.03$	3.59
toluene	$-0.83 \pm 0.20$	$-10.98 \pm 0.30$	10.15
phenanthrene	$-3.88 \pm 0.60$	$-10.90 \pm 0.04$	7.02

With these results, the need for binary interaction parameters became clear. First, we estimated  $k_{ij}$  with the SAFT-VR Mie EoS and experimental vapor pressure data, but this strategy also provided results that had high absolute deviations from the experimental data. Therefore, we used the approach of estimating  $k_{ij}$  with the output from solvation free energy calculations with molecular dynamics, as described in the last paragraph of Section 4.2. We initially found individual values for the interaction parameter of each pair, but, since the parameters for aromatic solutes were very similar (0.148, 0.162, 0.152), we averaged these values. By doing that, we obtained a general parameter for the water+aromatic pairs, which is exposed in Table 5.3.3. Also in this table, we display the binary interaction parameter for the pair water+propane.

Table 5.3.3 – Binary interaction parameters employed.

Pair	$k_{ij}$
water+propane	0.067
water+aromatic	0.154

The relatively large  $k_{ij}$  value of the interaction between aromatic solutes and water can be related to the lack of an explicit association term in the equation of state used to obtain the parameters for water. Actually, the SAFT-VR Mie has an association term (LAFITTE *et al.*, 2013), but it was not incorporated in the force field. The SAFT- $\gamma$  Mie model for water (LOBANOVA *et al.*, 2016) has two different temperature-dependent sets of parameters. The parameters utilized in this work were those estimated with experimental interfacial tension data. Hence, we tested the only binary interaction parameter for water+toluene estimated with MD interfacial data available in the literature (HERDES *et al.*, 2017). Nevertheless, the result also had a high absolute deviation, and this parameter could not be transferred to the calculation of the solvation free energy of toluene in water.

These issues faced by SAFT- $\gamma$  Mie model can also be related to the problems of modeling water with a coarse-grained force field. One of the main difficulties is the choice of which water molecules are going to be represented by which specific beads since water molecules move independently and interact by non-bonded interactions (HADLEY; MCCABE, 2010; HADLEY; MCCABE, 2012). The SAFT- $\gamma$  Mie water considers that one water molecule corresponds to one bead. This strategy only saves a small amount of simulation time, but it can predict properties at physiological temperatures unlike other more aggressive models such as the MARTINI, which considers that one bead represents various water molecules. In light of all these problems related to modeling the water molecule, the SAFT- $\gamma$  Mie force field appears to be a good alternative when working close to room temperatures, but the necessity of additional parameters estimated with molecular simulation indicates severe flaws in the methodology. This estimation of the binary parameter increased significantly the simulation time required to calculate the hydration free energies, since we had to carry out three additional simulations for every pair water-solute and then three other simulations for the three water+aromatic solutes in order to test the averaged binary interaction parameter. If these simulations are necessary for every time a new mixture with water is going to be studied with the SAFT- $\gamma$  Mie force field, the use of this model can become impractical. With this idea in mind, a useful investigation to be made is to check how accurate would be the prediction of the hydration free energy of other aromatic solutes by the SAFT- $\gamma$  Mie force field with the binary interaction parameter estimated here. Using these binary interaction parameters calculated with data from molecular dynamics, we then obtained the final hydration free energies presented in Table 5.3.4.

Table 5.3.4 – Calculated and experimental hydration free energies (kcal/mol) of solutes in water.

Solute	$\Delta G_{solv}^{GAFF}$	$\Delta G_{solv}^{ELBA}$	$\Delta G_{solv}^{exp}$	$\Delta G_{solv}^{Mie}$	Absolute Deviation
propane	2.50 $\pm$ 0.02	2.76 $\pm$ 0.02	2.00 $\pm$ 0.20	2.01 $\pm$ 0.01	0.01
benzene	-0.81 $\pm$ 0.02	-0.69 $\pm$ 0.01	-0.86 $\pm$ 0.20	-1.12 $\pm$ 0.01	0.26
toluene	-0.79 $\pm$ 0.03	-0.76 $\pm$ 0.01	-0.83 $\pm$ 0.20	-0.84 $\pm$ 0.01	0.01
phenanthrene	-5.26 $\pm$ 0.03	N/A	-3.88 $\pm$ 0.60	-3.47 $\pm$ 0.02	0.41

Hydration free energies calculated using the SAFT- $\gamma$  Mie force field with  $k_{ij} \neq 0$  had low absolute deviations from the experimental data, as expected since the parameters were adjusted to fit these experimental data. In the table above, we also show the results obtained by [Genheden \(2016\)](#) with the ELBA force field and by [Mobley and Guthrie \(2014\)](#) with the GAFF force field for the solutes and with the TIP3P model for water. The GAFF (General Amber Force Field) force field is an all-atom model that consists of bonded and non-bonded parameters and is suitable for the study of a significant number of molecules. In turn, the TIP3P model considers that water is a rigid monomer represented by three interacting sites with non-bonded interactions and Coulombic interactions ([JORGENSEN et al., 1983](#)). Both the GAFF and the TIP3P models use the Lennard-Jones potential to calculate the non-bonded interactions. The solvation free energies for the ELBA force field were estimated with thermodynamic integration and the solvation free energies with the GAFF force field were estimated with MBAR.

Comparing the results of the three aforementioned force fields, the root mean square error (RMSE) of all the pairs tested with the SAFT- $\gamma$  Mie model was 0.24, the RMSE for hydration free energies obtained with the GAFF force field was 0.73, and that for the ELBA coarse-grained force field was 0.44. The difference in absolute deviations between the GAFF and SAFT- $\gamma$  Mie force fields is significantly high for phenanthrene, hence the coarse-grained force field with a binary parameter is preferred if the application requires a high level of accuracy. The results also indicated that the SAFT- $\gamma$  Mie model with the binary interaction parameter performed better than the ELBA force field in modeling the solvation phenomenon of the pairs studied in this work, but performed worse with the binary parameter set to zero. This difference in performance occurred despite the fact that both the SAFT- $\gamma$  Mie and ELBA models have the same level of coarse-graining for the solvent (one bead represents one water molecule). Therefore, the choice between the two coarse-grained models is dependent on the availability and transferability of binary interaction parameters of the Mie Model. We also present, for the SAFT- $\gamma$  Mie force field, the hydration free energy profiles in Figure 5.3.1. Bigger molecules had steeper free energy profiles, as it was for the solvation free energy study in other solvents. We also observe that the hydration free energy for the first non-zero

$\lambda$  is negative for benzene and toluene when a positive value is expected since free energy is required for cavity formation in the solvent for the insertion of the solute. This anomaly can be caused by the fact that the exponential parameters in the Mie potential compensate for cavity formation.

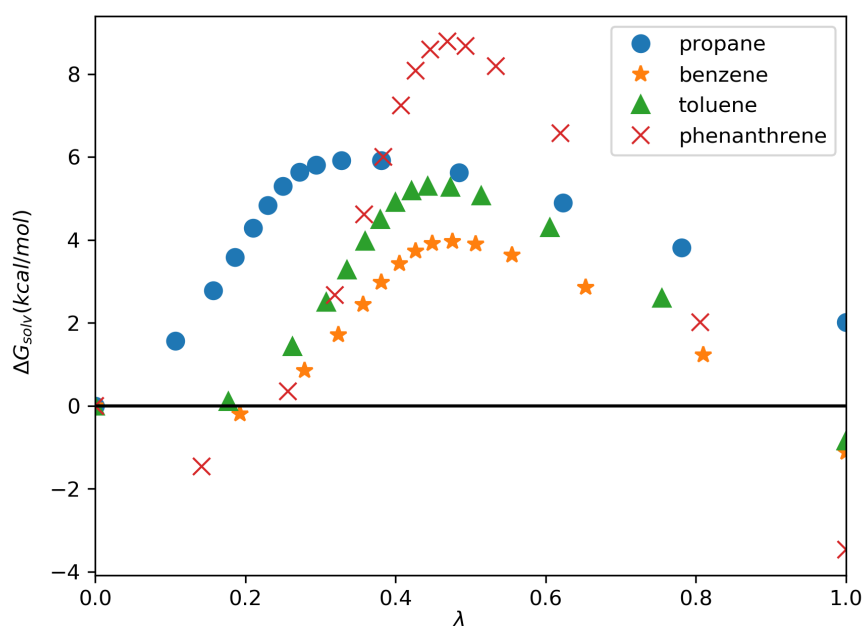


Figure 5.3.1 – Representation of hydration free energies of different solutes estimated at each alchemical state.

The results found here for both the solvation free energies and hydration free energies fulfilled the intentions of this dissertation. We assessed the prediction capability of the SAFT- $\gamma$  Mie force field and provided satisfactory solvation free energy estimates of PAHs using a coarse-grained force field. In addition to that, we found flaws in the methodology used by the SAFT- $\gamma$  Mie force field to model the water molecule. Hence, these shortcomings of this model can now be addressed, and the force field can even be improved by using other mixing rules to avoid the use of a binary parameter or, even, using hydration free energy estimates in the parameterization of water. These results also encourage us to calculate solvation free energies of more complex molecules mimicking asphaltenes in non-aqueous solvents in future studies.

## 5.4 Partition Coefficients

Using the solvation free energies estimated in the sections above, we also calculated partition coefficients by means of Eq. (3.115), for the pairs water/1-octanol and water/hexane with the intention of testing the modeling capabilities of the SAFT- $\gamma$

Mie force field again. The partition functions studied here have many experimental data available in the literature due to their environmental importance (SANGSTER, 1997). Besides this, the calculations of these specific partition coefficients are relevant because 1-octanol is used to quantify hydrophobicity and can serve as a model for biological lipids and different soils (RUELLE, 2000), and hexane is a model for an apolar, hydrophobic phase. Calculated values and experimental data are shown in Table 5.4.1. The experimental data of the partition coefficients were taken from Poole *et al.* (2000), Sangster (1997) for the coefficient of water/1-octanol and from Schulte *et al.* (1998) for the coefficient of water/hexane.

Table 5.4.1 – Partition Coefficient Calculated from MD simulations and from experimental data.

	Molecular Dynamics	Experimental	Absolute Deviation
$\log P^{water/1-octanol}$			
propane	2.47	2.40	0.07
phenanthrene	3.57	4.46	0.89
$\log P^{water/hexane}$			
benzene	1.93	2.06	0.13
phenanthrene	4.17	4.49	0.32

Overall absolute deviations were small for pairs with smaller solvation free energy deviations such as propane and benzene. The water/1-octanol partition coefficient of phenanthrene had higher deviation due to the higher deviation of the free energy of solvation of this compound in 1-octanol. Comparing with other force fields, Garrido *et al.* (2009) reported average absolute deviations for the water/1-octanol partition coefficient of 0.4 with the GROMOS 53a6 force field (OOSTENBRINK *et al.*, 2004), 0.3 for TraPPE, and 0.9 for OPLS-AA/TraPPE force fields. However, they attribute the low deviations of TraPPE to the cancellation of errors between the two solvation free energies. Additionally, Genheden (2016) found average absolute deviations of 0.86 for the water/hexane partition coefficients and of 0.75 for the water/1-octanol partition coefficients with the ELBA coarse-grained force field. In this dissertation, we performed a small study of partition coefficients with the SAFT- $\gamma$  Mie force field. Hence, a larger set would be necessary to do a complete evaluation of the performance of this force field in the prediction of partition coefficients.

## 6 Conclusions

This dissertation consisted in the study of solvation free energy calculations of aromatic solutes in different solvents by using the SAFT- $\gamma$  Mie coarse-grained force field. Solvation free energy studies are mostly done using water as a solvent and with all-atom force fields based on the Lennard-Jones potential. Therefore, with this study, we were able to provide data about the capability of a coarse-grained force field based on the Mie potential in calculating solvation free energies. Additionally, the solvation free energy estimations carried out here can help improve the SAFT- $\gamma$  Mie force field since these calculations are helpful in identifying errors and shortcomings in the modeling process. The SAFT- $\gamma$  Mie uses the SAFT-VR Mie EoS in its parameterization, which results in a relatively straightforward top-down method of obtaining parameters. Following this strategy, the phenanthrene parameters, which were not available in the original database of this force field, were obtained using vapor-liquid equilibrium data and two different ring equations and geometries. However, only the parameters estimated with the ring equation proposed by Müller and Mejía (2017) were utilized in the solvation free energy simulations since this equation did not require molecular simulation data in its parameterization.

To perform our expanded ensemble simulations, we optimized the coupling parameters and their respective simulation weights. The resulting potential energies from the expanded ensemble simulations then served as input to estimate solvation free energy differences with the MBAR method. The results for non-aqueous solvents had absolute deviations from the experimental data of less than 2.0 kcal/mol, except for the pair 1-octanol+anthracene. We also observed the geometry effect on the free energy curves - larger molecules had steeper curves and more substantial absolute deviations. The influence of carbon dioxide on the solvation free energy of phenanthrene in toluene was found to be negligible according to the SAFT- $\gamma$  Mie force field.

Hydration free energy calculations with the SAFT- $\gamma$  Mie model required the use of relatively large values of  $k_{ij}$  to produce satisfactory results. We chose to estimate the parameter with the output from molecular dynamics data since the strategy of using the SAFT-VR Mie EoS provided high absolute deviations from the experimental data. This necessity of one additional parameter probably happens due to the lack of a term to account for the hydrogen bond in the EoS on which this force field is based and due to problems associated with the coarse-graining of water molecules. The results obtained with  $k_{ij}$  estimated with MD output were satisfactory, the absolute deviations from the experimental data found were smaller than the ones for the GAFF and ELBA force fields. We also used the solvation free energies to calculate partition coefficients in water/1-

octanol and water/hexane. The obtained absolute deviations from experimental data were similar to the ones found for all-atom force fields (GROMOS, TraPPE and OPLS-AA/TraPPE) and another coarse-grained force field (ELBA).

Overall, the SAFT- $\gamma$  Mie force field proved to be a suitable model to represent the solvation phenomenon of non-aqueous solvents. It correctly described solvation free energies of solutes mimicking asphaltenes dissolved in hexane, toluene, 1-octanol. However, the calculation of hydration free energies required the use of a binary interaction parameter estimated with MD output, which increased the simulation time significantly. This fact evidenced flaws in the methodology used by the SAFT- $\gamma$  force field and raised questions about the feasibility of this model for hydration free energy calculations. Nevertheless, the SAFT- $\gamma$  Mie force field for water used here does not predict freezing at room temperature as other force fields do, which is essential for our hydration free energy calculations. Therefore, it would be relevant to test if the binary interaction parameter for our aromatic solutes estimated here can be used in hydration free energy calculations of other aromatic solutes and if we could use MBAR to obtain the  $k_{ij}$  through reweighting. Hence, we would only need to carry out molecular dynamics simulations with one value of  $k_{ij}$ , and then use this output to estimate with MBAR the results with other  $k'_{ij}$ s. We also have some ideas that could be developed in the future using the results from this dissertation. The SAFT- $\gamma$  Mie force field could be used to model larger asphaltene models and, consequently, increase the scale of the simulations we performed. Additionally, it would be a valid investigation to study new methodologies to calculate solubility with solvation free energies using the SAFT- $\gamma$  Mie force field.



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# APPENDIX A – Detailing of the SAFT-VR Mie Equation of State

The term  $a_1$  in Eq. 3.76 is the first-order mean-attractive energy of the mixture (LAFITTE *et al.*, 2013), and is given by

$$a_1 = \sum_{i=1}^n \sum_{j=1}^n x_{s,i} x_{s,j} a_{1,ij}, \quad (\text{A.1})$$

where  $a_{1,ij}$  is the term corresponding to the pair interactions. It is equal to

$$a_{1,ij} = \mathcal{C}_{ij} \{ x_{0,ij}^{\lambda_{a,ij}} [a_{1,ij}^s(\rho_s; \lambda_{a,ij}) + B_{ij}(\rho_s; \lambda_{a,ij})] - x_{0,ij}^{\lambda_{r,ij}} [a_{1,ij}^s(\rho_s; \lambda_{r,ij}) + B_{ij}(\rho_s; \lambda_{r,ij})] \}, \quad (\text{A.2})$$

with  $\mathcal{C}_{ij}$  equals to

$$\mathcal{C}_{ij} = \frac{\lambda_{r,ij}}{\lambda_{r,ij} - \lambda_{a,ij}} \left( \frac{\lambda_{r,ij}}{\lambda_{a,ij}} \right)^{\left( \frac{\lambda_{a,ij}}{\lambda_{a,ij} - \lambda_{r,ij}} \right)}. \quad (\text{A.3})$$

Also in Eq. A.2,  $B_{ij}(\rho_s; \lambda_{ij})$  is equal to

$$B_{ij}(\rho_s; \lambda_{ij}) = 2\pi\rho_s d_{ij}^3 \epsilon_{ij} \left[ \frac{1 - \zeta_x/2}{(1 - \zeta_x)^3} I_{\lambda,ij} - \frac{9\zeta_x(1 + \zeta_x)}{2(1 - \zeta_x)^3} J_{\lambda,ij} \right], \quad (\text{A.4})$$

$$\zeta_x = \frac{\pi\rho_s}{6} \sum_{i=1}^n \sum_{j=1}^n x_{s,i} x_{s,j} d_{ij}^3. \quad (\text{A.5})$$

Here,  $I_{\lambda,ij}$  and  $J_{\lambda,ij}$  two terms that depend of the Mie potential. They are given by

$$I_{\lambda,ij} = \frac{(x_{0,ij})^{3-\lambda_{ij}} - 1}{\lambda_{ij} - 3}, \quad (\text{A.6})$$

$$J_{\lambda,ij} = \frac{(x_{0,ij})^{4-\lambda_{ij}}(\lambda_{ij} - 3) - (x_{0,ij})^{3-\lambda_{ij}}(\lambda_{ij} - 4) - 1}{(\lambda_{ij} - 3)(\lambda_{ij} - 4)}. \quad (\text{A.7})$$

The terms  $a_{1,ij}^s$  are the mean-attractive energies of a Sutherland potential of variable range and are obtained using a SAFT-VR treatment (AVENDAÑO *et al.*, 2013).

The resulting equation is:

$$a_{1,ij}^S(\rho_s; \lambda_{ij}) = -2\rho_s \left( \frac{\pi \epsilon_{ij} d_{ij}^3}{\lambda_{ij} - 3} \right) \frac{1 - \zeta_x^{eff}(\lambda_{ij})/2}{[1 - \zeta_x^{eff}(\lambda_{ij})]^3}, \quad (\text{A.8})$$

where  $\zeta_x^{eff}(\lambda_{ij})$  is the effective packing fraction obtained within a van der Waals one-fluid approximation (LAFITTE *et al.*, 2013). It is equal to

$$\zeta_x^{eff}(\lambda_{ij}) = c_1(\lambda_{ij})\zeta_x + c_2(\lambda_{ij})\zeta_x^2 + c_3(\lambda_{ij})\zeta_x^3 + c_4(\lambda_{ij})\zeta_x^4. \quad (\text{A.9})$$

Here, the coefficients  $c_1$ ,  $c_2$ ,  $c_3$  and  $c_4$  are

$$\begin{bmatrix} c_1 \\ c_2 \\ c_3 \\ c_4 \end{bmatrix} = \begin{bmatrix} 0.81096 & 1.7888 & -37.578 & 92.284 \\ 1.0205 & -19.341 & 151.26 & -463.50 \\ -1.9057 & 22.845 & -228.14 & 973.92 \\ 1.0885 & -6.1962 & 106.98 & -677.64 \end{bmatrix} \cdot \begin{bmatrix} 1 \\ 1/\lambda_{ij} \\ 1/\lambda_{ij}^2 \\ 1/\lambda_{ij}^3 \end{bmatrix}. \quad (\text{A.10})$$

The second-order fluctuation term,  $a_2$ , in Eq. 3.76 has a similar formulation to  $a_1$ . It is given by

$$a_2 = \sum_{i=1}^n \sum_{j=1}^n x_{s,i} x_{s,j} a_{2,ij}, \quad (\text{A.11})$$

where  $a_{2,ij}$  is equal to

$$\begin{aligned} a_{2,ij} = & \frac{1}{2} K^{HS} (1 + \chi_{ij}) \epsilon_{ij} \mathcal{C}_{ij}^2 \{ x_{0,ij}^{2\lambda_{a,ij}} [a_{1,ij}^s(\rho_s; 2\lambda_{a,ij}) + B_{ij}(\rho_s; 2\lambda_{a,ij})] \\ & - 2x_{0,ij}^{2\lambda_{a,ij}+2\lambda_{r,ij}} [a_{1,ij}^s(\rho_s; \lambda_{a,ij} + \lambda_{r,ij}) + B_{ij}(\rho_s; \lambda_{a,ij} + \lambda_{r,ij})] \\ & + x_{0,ij}^{2\lambda_{r,ij}} [a_{1,ij}^s(\rho_s; 2\lambda_{r,ij}) + B_{ij}(\rho_s; 2\lambda_{r,ij})] \}. \end{aligned} \quad (\text{A.12})$$

In Eq. A.12,  $K^{HS}$  is the isothermal compressibility of the mixture of hard spheres. It is equal to

$$K^{HS} = \frac{(1 - \bar{\zeta}_x)^4}{1 + 4\bar{\zeta}_x + 4\bar{\zeta}_x^2 + 4\bar{\zeta}_x^3 + \bar{\zeta}_x^4}, \quad (\text{A.13})$$

where  $\bar{\zeta}_x$  is the packing fraction for the mixture. It is equal to

$$\bar{\zeta}_x = \frac{\pi \rho_s}{6} \sum_{j=1}^n x_{s,i} x_{s,j} \sigma_{ij}^3, \quad (\text{A.14})$$

In Eq. A.12,  $\chi_{ij}$  is an empirical function of  $\bar{\zeta}_x$ . It is given by

$$\chi_{ij} = f_1(\alpha_{ij})\bar{\zeta}_x + f_2(\alpha_{ij})\bar{\zeta}_x^5 + f_3(\alpha_{ij})\bar{\zeta}_x^8, \quad (\text{A.15})$$

where  $\alpha_{ij}$  represents the equation for mixtures for the reduced attractive van der Waals constant of the Mie potential. It is given by

$$\alpha_{ij} = C_{ij} \left( \frac{1}{\lambda_{a,ij} - 3} - \frac{1}{\lambda_{r,ij} - 3} \right). \quad (\text{A.16})$$

Finally,  $a_3$  in Eq. 3.76 is equal to

$$a_3 = \sum_{i=1}^n \sum_{j=1}^n x_{s,i} x_{s,j} a_{3,ij}, \quad (\text{A.17})$$

where  $a_{3,ij}$  is equal to

$$a_{3,ij} = -\epsilon_{ij}^3 f_4(\alpha_{ij}) \bar{\zeta}_x \exp[f_5(\alpha_{ij}) \bar{\zeta}_x + f_6(\alpha_{ij}) \bar{\zeta}_x^2]. \quad (\text{A.18})$$

The functions  $f_k$  ( $k = 1, \dots, 6$ ) are obtained with

$$f_k(\alpha_{ij}) = \frac{\sum_{n=0}^{n=3} \phi_{k,n} \alpha_{ij}^n}{1 + \sum_{n=4}^{n=6} \phi_{k,n} \alpha_{ij}^{n-3}}, \quad (\text{A.19})$$

where  $\phi_{k,n}$  are coefficients defined in the original paper by Lafitte *et al.* (2013).

The density dependent coefficients of Eq. 3.83 are given by the following equations:

$$k_0 = -\ln(1 - \zeta_x) + \frac{42\zeta_x - 39\zeta_x^2 + 9\zeta_x^3 - 2\zeta_x^4}{6(1 - \zeta_x)^3}, \quad (\text{A.20})$$

$$k_1 = \frac{\zeta_x^4 + 6\zeta_x^2 - 12\zeta_x}{2(1 - \zeta_x)^3}, \quad (\text{A.21})$$

$$k_2 = \frac{-3\zeta_x^2}{8(1 - \zeta_x)^2}, \quad (\text{A.22})$$

$$k_3 = \frac{-\zeta_x^4 + 3\zeta_x^2 + 3\zeta_x}{6(1 - \zeta_x)^3}. \quad (\text{A.23})$$

In Eq. 3.82, the term  $g_{1,ij}(\sigma_{ij})$  is the first-order contribution to the radial distribu-

tion function. It has the following form:

$$g_{1,ij}(\sigma_{ij}) = \frac{1}{2\pi\epsilon_{ij}d_{ij}^3} \left[ 3 \frac{\partial a_{1,ij}}{\partial \rho_s} - \mathcal{C}_{ij} \lambda_{a,ij} x_{0,ij}^{\lambda_{a,ij}} \frac{a_{1,ij}^s(\rho_s; \lambda_{a,ij}) + B_{ij}(\rho_s; \lambda_{a,ij})}{\rho_s} \right. \\ \left. + \mathcal{C}_{ij} \lambda_{r,ij} x_{0,ij}^{\lambda_{r,ij}} \frac{a_{1,ij}^s(\rho_s; \lambda_{r,ij}) + B_{ij}(\rho_s; \lambda_{r,ij})}{\rho_s} \right]. \quad (\text{A.24})$$

Also in Eq. 3.82, the second-order contribution to the radial distribution function ( $g_{2,ij}(\sigma_{ij})$ ) is equal to

$$g_{2,ij}(\sigma_{ij}) = (1 + \gamma_{c,ij}) g_{2,ij}^{MCA}(\sigma_{ij}), \quad (\text{A.25})$$

where  $g_{2,ij}^{MCA}(\sigma_{ij})$  is equal to

$$g_{2,ij}(\sigma_{ij}) = \frac{1}{2\pi\epsilon_{ij}d_{ij}^3} \left[ 3 \frac{\partial \frac{a_2}{1+\gamma_{ij}}}{\partial \rho_s} - \epsilon_{ij} K^{HS} \mathcal{C}_{ij}^2 \lambda_{r,ij} x_{0,ij}^{2\lambda_{r,ij}} \frac{a_{1,ij}^s(\rho_s; 2\lambda_{r,ij}) + B_{ij}(\rho_s; 2\lambda_{r,ij})}{\rho_s} \right. \\ \left. + \epsilon_{ij} K^{HS} \mathcal{C}_{ij}^2 (\lambda_{r,ij} + \lambda_{a,ij}) (x_{0,ij})^{\lambda_{r,ij} + \lambda_{a,ij}} \frac{a_{1,ij}^s(\rho_s; \lambda_{r,ij} + \lambda_{a,ij}) + B_{ij}(\rho_s; \lambda_{r,ij} + \lambda_{a,ij})}{\rho_s} \right. \\ \left. - \epsilon_{ij} K^{HS} \mathcal{C}_{ij}^2 \lambda_{a,ij} x_{0,ij}^{2\lambda_{a,ij}} \frac{a_{1,ij}^s(\rho_s; 2\lambda_{a,ij}) + B_{ij}(\rho_s; 2\lambda_{a,ij})}{\rho_s} \right], \quad (\text{A.26})$$

and  $\gamma_{c,ij}$  is a correction factor obtained from the equation bellow:

$$\gamma_{c,ij} = \phi_{7,0} \bar{\zeta}_x \theta_{ij} \exp(\phi_{7,3} \bar{\zeta}_x + \phi_{7,4} \bar{\zeta}_x^2) \{1 - \tanh[\phi_{7,1}(\phi_{7,2} - \alpha_{ij})]\}, \quad (\text{A.27})$$

where  $\theta_{ij} = \exp(\beta\epsilon_{ij}) - 1$ .

## APPENDIX B – Overlap Matrices



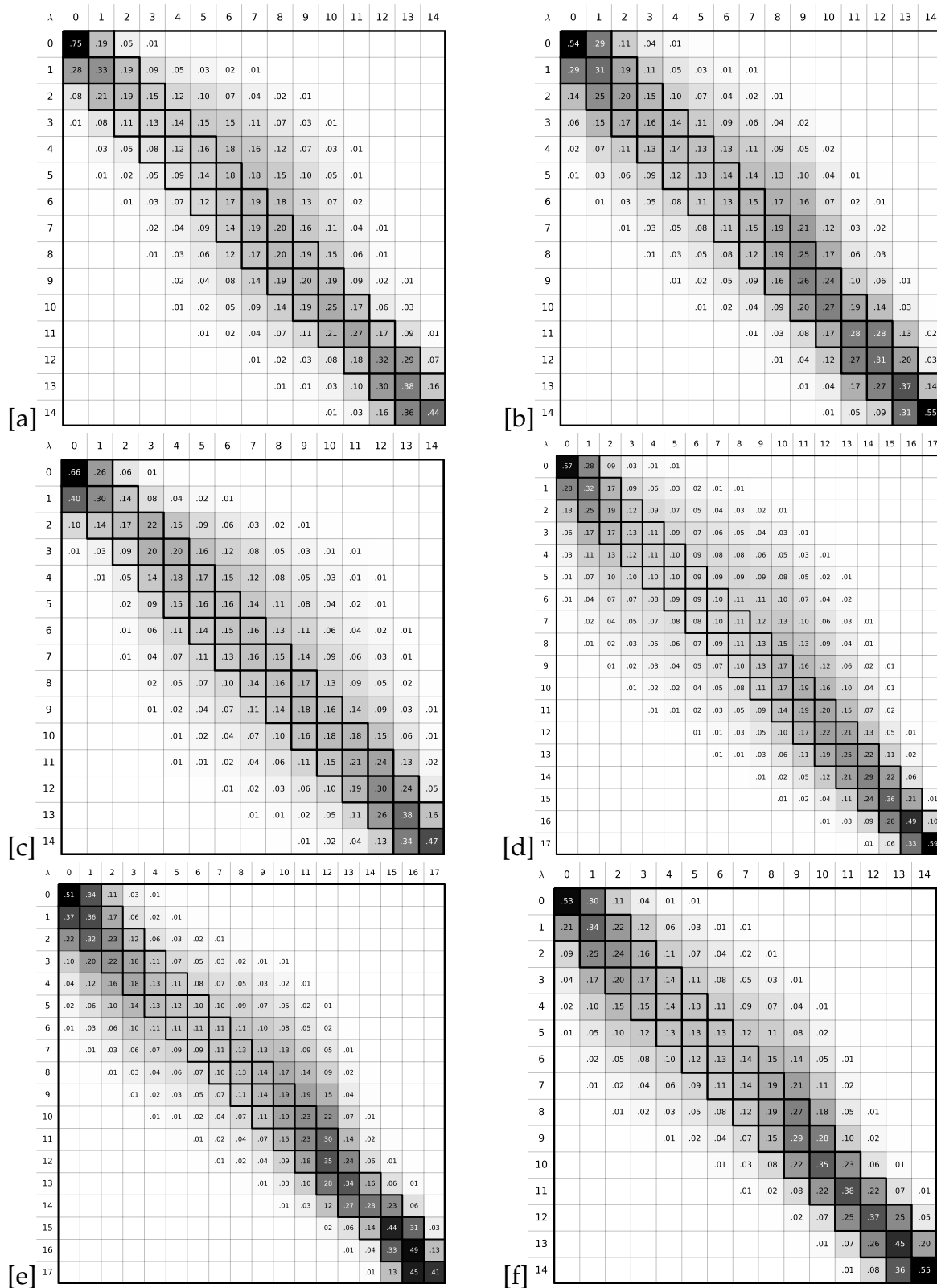


Figure B.0.1 – Overlap matrix for hexane+pyrene [a], hexane+phenanthrene [b], 1-octanol+propane [c], 1-octanol+anthracene [d], 1-octanol+phenanthrene [e], and toluene+pyrene [f].

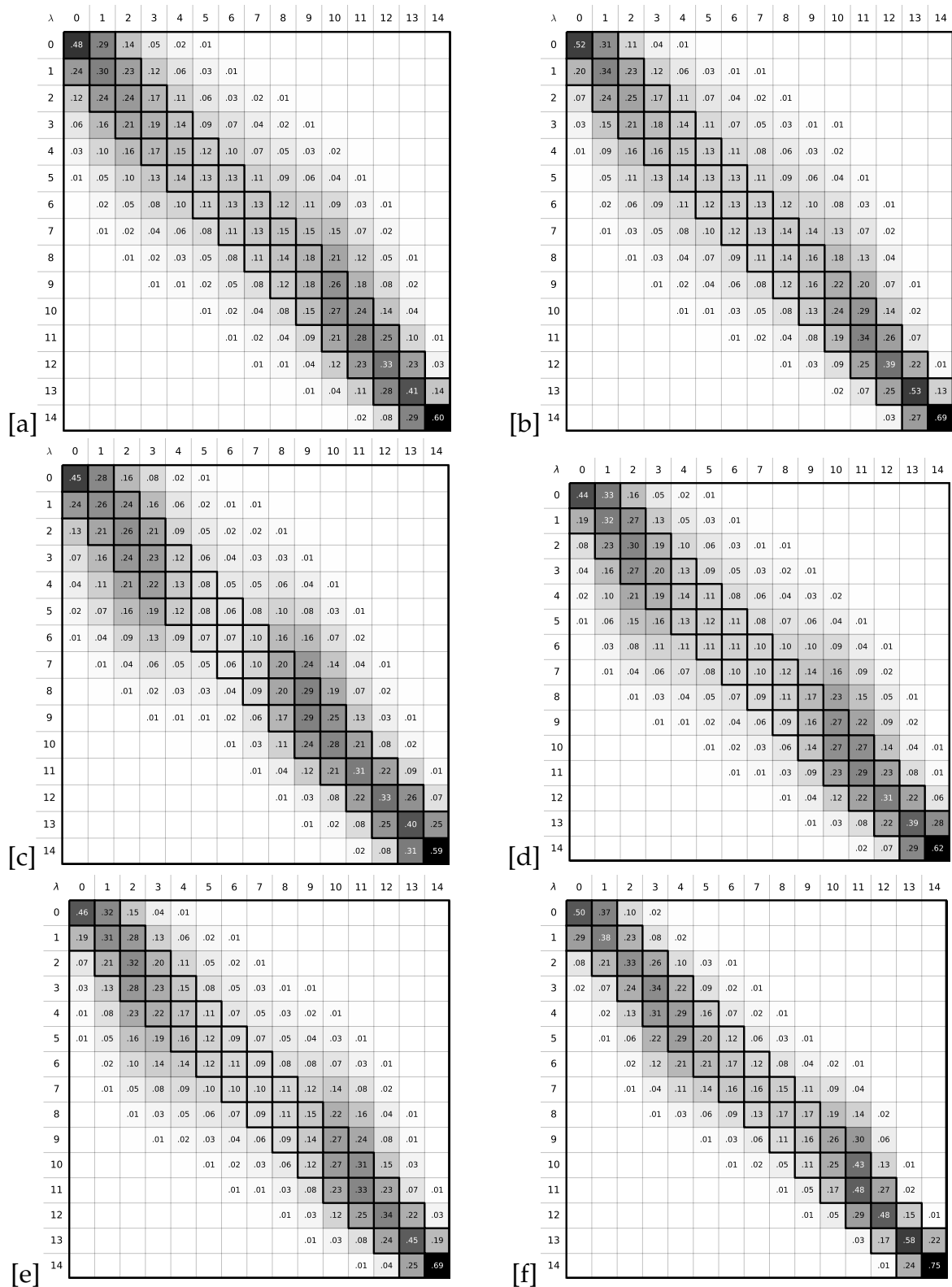


Figure B.0.2 – Overlap matrix for toluene+anthracene [a], toluene+phenanthrene [b], water+propane [c], water+benzene [d], water+toluene [e], and water+phenanthrene [f] .

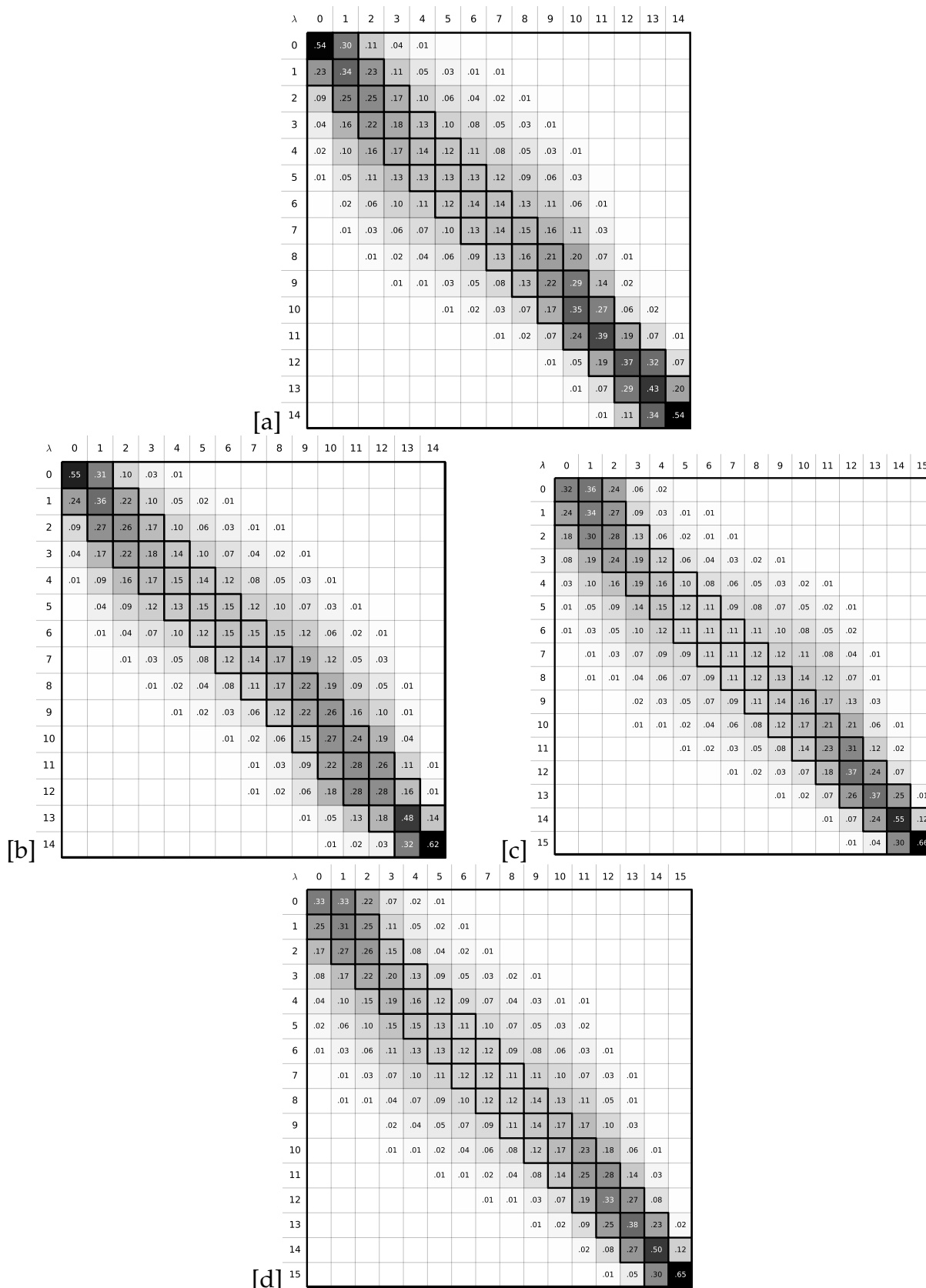


Figure B.0.3 – Overlap matrix for the different  $w_{CO_2}$  of the mixture toluene+ $CO_2$ +phenanthrene.  $w_{CO_2} = 0.087$  [a],  $w_{CO_2} = 0.119$  [b],  $w_{CO_2} = 0.169$  [c], and  $w_{CO_2} = 0.289$  [d].

# Annex

# ANNEX A – Work Published in Scientific Conference



## PARAMETRIZAÇÃO E AVALIAÇÃO DO CAMPO DE FORÇA SAFT- $\gamma$ CG PARA SIMULAÇÃO MOLECULAR DE FENANTRENO

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O fenantreno é um hidrocarboneto policíclico aromático que pode ser usado para estudar moléculas mais complexas como os asfaltenos. Portanto, é interessante desenvolver campos de forças para simulação molecular eficientes computacionalmente que consigam descrever esses tipos de molécula. O campo de força SAFT- $\gamma$  CG, escolhido para o estudo, é do tipo *coarse-grained* e foi desenvolvido com base na equação de estado SAFT-VR Mie. Essa equação modela uma substância subdividindo sua estrutura em segmentos que interagem através de potencial de Mie. Os parâmetros desse campo de força para o fenantreno foram estimados e avaliados em relação à determinação de propriedades de equilíbrio líquido-vapor (ELV). A estratégia de estimação possui duas etapas. A primeira consiste em minimizar o erro quadrático entre a pressão de vapor calculada com a equação de estado (EdE) SAFT-VR Mie e os valores experimentais. Os parâmetros dessa minimização foram, então, usados no cálculo de ELV com o método de *Gibbs Ensemble Monte Carlo* com volume total constante (GEMC-NVT). A segunda etapa consistiu em estimar novamente os parâmetros através do método de mínimos quadrados envolvendo a pressão de vapor e a densidade de líquido saturado calculadas por Monte Carlo (MC) e as calculadas com a EdE SAFT-VR Mie. Essa segunda parte é necessária devido a aproximações teóricas que geram diferenças entre os resultados da simulação molecular e da EdE. Esse campo de força apresentou uma boa descrição da pressão de vapor do fenantreno com método de GEMC quando comparado com os dados experimentais. As densidades de líquido e vapor saturados e as propriedades críticas apresentaram resultados similares aos obtidos com o campo de força atomístico TraPPE-EH, sendo que o SAFT- $\gamma$  CG exige menor esforço computacional. Para continuação do trabalho, pretende-se usar esses parâmetros para calcular a energia de solvatação do fenantreno, em diluição infinita, em tolueno e em solução tolueno + CO<sub>2</sub> com dinâmica molecular.

Palavras-chave: campo de força, SAFT- $\gamma$  CG, simulação molecular, fenantreno.

### Introdução

Os campos de força do tipo *coarse-grained* parametrizados a partir de propriedades experimentais são uma alternativa aos campos de força desenvolvidos com cálculos *ab initio* quando a escala das simulações moleculares precisa ser aumentada. O método *coarse-grained* consiste, basicamente, em dividir a substância em pseudo-átomos representativos de grupos de átomos. Normalmente, a parametrização do campo de força é feita a partir de informações obtidas em uma escala mais detalhada. Quando as informações utilizadas provêm do comportamento em uma escala maior, a estratégia para o modelo *coarse-grained* é dita ser do tipo *top down*. Um dos campos de força que possuem essa estratégia é o SAFT- $\gamma$  CG (Avendaño *et al.*, 2011), o qual foi desenvolvido com base na equação de estado (EdE) SAFT-VR Mie (Lafitte *et al.*, 2013). Esse modelo usa o potencial de Mie para descrever moléculas formadas por segmentos conectados. A vantagem de usar essa equação como base

para o campo de força é a sua capacidade de descrever bem as propriedades do fluido, incluindo derivadas de segunda ordem, de uma variedade de sistemas (Lafitte *et al.*, 2013). Com base nessas ideias, o presente trabalho teve como objetivo parametrizar o campo de força SAFT- $\gamma$  CG para o fenantreno, cuja estrutura é apresentada na Figura 1, utilizando dados de equilíbrio líquido-vapor e considerando que ele é formado por três segmentos esféricos. O fenantreno foi escolhido como forma de testar a capacidade de representação desse campo de força simplificado, já que sua parametrização para moléculas menores tem sido bem sucedida (Lafitte *et al.*, 2012). Outra razão para o estudo do fenantreno é o fato de ele servir como modelo simplificado para moléculas mais complexas, tais como os asfaltenos, já que possui uma estrutura de anéis policondensados e é solúvel em tolueno.

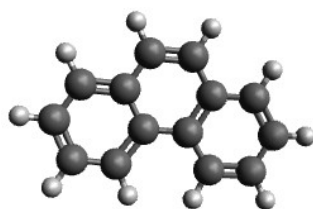


Figura1. Estrutura molecular do fenantreno.

## Metodologia

### *Estimação com a Equação de Estado SAFT-VR Mie*

A EdE SAFT-VR Mie descreve moléculas formadas por segmentos conectados e que interagem através do potencial de Mie, dado por:

$$U_{Mie}(r) = \varepsilon \frac{\lambda_r}{\lambda_r - \lambda_a} \left( \frac{\lambda_r}{\lambda_a} \right)^{\lambda_a / \lambda_r - \lambda_a} \left[ \left( \frac{\sigma}{r} \right)^{\lambda_r} - \left( \frac{\sigma}{r} \right)^{\lambda_a} \right] \quad (1)$$

em que  $\lambda_r$  é o expoente repulsivo,  $\lambda_a$  é o expoente atrativo,  $\sigma$  corresponde à distância entre os centros dos segmentos e  $\varepsilon$  é o parâmetro de energia do segmento. A energia livre de Helmholtz específica para a EdE SAFT-VR Mie para um fluido não-associativo é definida como

$$a = a^{IDEAL} + a^{MONO} + a^{CHAIN}, \quad (2)$$

em que  $a^{IDEAL}$  é a contribuição de gás ideal,  $a^{MONO}$  é contribuição dos monômeros (segmentos desconectados) e  $a^{CHAIN}$  é a contribuição relativa à formação das cadeias de segmentos. Para cadeias que possuem  $m_s$  segmentos do mesmo tipo tangencialmente ligados, a contribuição da cadeia é

$$a^{CHAIN} = -(m_s - 1) \ln^{Mie}(\sigma), \quad (3)$$

onde  $\ln^{Mie}(\sigma)$  é o valor da função de distribuição radial de pares para o fluido monomérico de referência (fluido de Mie). Devido à estrutura aromática do fenantreno, usou-se a seguinte expressão para anéis formados por  $m_s$  segmentos no lugar da contribuição de cadeia:

$$a^{RING} = -m_s \ln g^{Mie}(\sigma). \quad (4)$$

Essa substituição é feita porque a diferença entre uma molécula formada por cadeias e uma formada por anéis, tendo ambas o mesmo número de segmentos, é que a última possui uma ligação a mais (Lafitte *et al.*, 2012). Para o equacionamento completo da EdE, o leitor é referido ao artigo de Lafitte *et al.* (2013). Seguindo essa formulação, os parâmetros  $\varepsilon$ ,  $\lambda_r$  e  $\sigma$  foram estimados e os parâmetros  $m_s$  e  $\lambda_a$  foram fixados em 3 e 6, respectivamente. A razão para fixar o número de segmentos em três deve-se à própria estrutura do fenantreno, que consiste em três anéis aromáticos condensados. Já o parâmetro atrativo foi fixado no valor London para facilitar a estimação, já que é comprovada a alta correlação entre os parâmetros repulsivo e atrativo do potencial (Ramrattan *et al.*, 2015). A minimização foi feita através do método PSO (*Particle Swarm Optimization*) e com apenas dados de pressão de vapor, por uma questão de indisponibilidade de dados experimentais de densidade do fenantreno como líquido saturado. A função objetivo possuiu a seguinte forma:

$$F_{\text{exp}}(\sigma^{SAFT}, \varepsilon^{SAFT}, \lambda_r^{SAFT}) = \sum_{i=1}^{N_p} \left( \frac{P_v^{\text{exp}}(T_i) - P_v^{SAFT}(T_i)}{P_v^{\text{exp}}(T_i)} \right)^2 \quad (5)$$

em que  $N_p$  corresponde ao número de pontos experimentais,  $P_v^{\text{exp}}$  aos pontos de pressão de vapor experimental (Mortimer e Murphy, 1923) e  $P_v^{SAFT}$  à pressão de vapor calculada com a EdE SAFT-VR Mie. Esse cálculo de equilíbrio foi feito usando como base a rotina do ponto de bolha proposta por Smith *et al.* (2007).

#### *Cálculo com o Método “Gibbs Ensemble Monte Carlo” (GEMC)*

Os parâmetros estimados com equação SAFT-VR Mie foram usados para realizar simulações no GEMC (Panagiotopoulos, 1987) com o simulador CASSANDRA (Shah e Maginn, 2011). O método de GEMC foi desenvolvido com o intuito de estudar a coexistência entre fases através da simulação simultânea de duas caixas com condições de contorno periódicas e que trocam moléculas, energia e volume entre si, mas de forma a manter o volume total constante. O equilíbrio entre elas é obtido através de passos de Monte Carlo (MC) que incluem o deslocamento aleatório das moléculas, mudança de volume e transferências aleatórias de moléculas entre as caixas. Para sistemas com apenas um componente, os cálculos são realizados mantendo-se o volume e o número de partículas total constantes (NVT), mas de maneira a permitir a variação de volume ( $V = V^{liq} + V^{vap}$ ) e partículas ( $N = N^{liq} + N^{vap}$ ) dentro de cada caixa. Essas simulações no GEMC-NVT foram feitas inserindo-se aleatoriamente 400 moléculas na caixa líquida e 100 moléculas na caixa vapor. As densidades iniciais das caixas foram escolhidas ajustando-as às densidades obtidas com a EdE SAFT-VR Mie, para evitar que todas as moléculas migrassem pra uma única caixa ao longo da simulação. A simulação consistiu em, no mínimo, 10000 ciclos de equilibração e 100000 ciclos de produção, sendo que cada ciclo de MC corresponde a 1000 tentativas de rotação, 1000 de translação, 100 de inserção, 100 de exclusão e 10 de variação de volume. A distância de corte usada foi igual a quatro vezes o valor do diâmetro do segmento estimado e as interações de van der Waals foram calculadas através do potencial Mie com correção de longa distância (*tail correction*). As propriedades densidade de vapor ( $\rho_{vap}$ ), densidade de líquido ( $\rho_{liq}$ ) e pressão ( $P_v^{sim}$ ) foram amostradas a cada 100 ciclos de MC e essas amostragens foram divididas em cinco blocos para os cálculos da média e do desvio padrão. Os resultados



obtidos nessas simulações foram usados para estimar coeficientes de correção para os parâmetros conformacionais do campo de força ( $c_\sigma$  e  $c_\epsilon$ ), que são relacionados aos parâmetros provenientes da EdE SAFT-VR Mie através de parâmetros oriundos de ajuste,

$$\sigma^{ajuste} = c_\sigma \sigma^{SAFT} \quad (6)$$

$$\epsilon^{ajuste} = c_\epsilon \epsilon^{SAFT} \quad (7)$$

Os parâmetros de ajuste são necessários porque as aproximações teóricas feitas na EdE geram discrepâncias entre os resultados da simulação molecular e os da EdE (Lafitte *et al.*, 2012). Por isso, a estimação desses parâmetros foi feita de maneira a diminuir as diferenças entre a pressão de vapor e a densidade de líquido saturado obtidas com a equação de estado e as obtidas via simulação molecular. A função objetivo, minimizada através do método PSO, possuiu a seguinte forma:

$$F_{\text{sim}}(c_\sigma, c_\epsilon) = \sum_{i=1}^{N_P} \left( \frac{P_v^{\text{sim}}(T_i, \sigma^{SAFT}, \epsilon^{SAFT}) - P_v^{\text{SAFT}}(T_i, \sigma^{ajuste}, \epsilon^{ajuste})}{P_v^{\text{sim}}(T_i, \sigma^{SAFT}, \epsilon^{SAFT})} \right)^2 + \sum_{i=1}^{N_P} \left( \frac{\rho_{\text{liq}}^{\text{sim}}(T_i, \sigma^{SAFT}, \epsilon^{SAFT}) - \rho_{\text{liq}}^{\text{SAFT}}(T_i, \sigma^{ajuste}, \epsilon^{ajuste})}{\rho_{\text{liq}}^{\text{sim}}(T_i, \sigma^{SAFT}, \epsilon^{SAFT})} \right)^2 \quad (8)$$

Com os parâmetros atrativo e repulsivo fixados nos valores encontrados com a EdE SAFT-VR Mie, o espaço paramétrico pode ser redefinido de maneira a se encontrar um conjunto de parâmetros refinado para o campo de força SAFT- $\gamma$  CG (Lafitte *et al.*, 2012), que são:

$$\sigma^{\text{sim}} = \sigma^{SAFT} / c_\sigma \quad (9)$$

$$\epsilon^{\text{sim}} = \epsilon^{SAFT} / c_\epsilon \quad (10)$$

As simulações pelo método GEMC-NVT foram refeitas seguindo-se a mesma metodologia e os resultados foram comparados com os dados de referência. A determinação do ponto crítico não foi realizada por meio do método GEMC-NVT, pois ele apresenta grandes flutuações perto do ponto crítico, que fazem as caixas de simulação mudarem de identidade durante a simulação. A temperatura crítica ( $T_C$ ) foi então ajustada através do método PSO com a seguinte equação:

$$(\rho_{\text{liq}} - \rho_{\text{vap}}) = A(T_C - T)^\eta \quad (11)$$

As densidades de líquido e vapor no equilíbrio na equação acima foram provenientes de simulações feitas em uma faixa de temperatura entre 476,75 e 825 K. O expoente crítico ( $\eta$ ) foi fixado no valor correspondente ao seu valor universal (0,325) (Hansen e McDonald, 2006) e a constante A foi determinada pelo ajuste. A densidade crítica ( $\rho_c$ ) foi obtida através da lei linear dos retângulos (Equação 12) na mesma faixa de temperatura usada para obter  $T_C$ .

$$\frac{(\rho_{\text{liq}} - \rho_{\text{vap}})}{2} = \rho_c + D(T_C - T) \quad (12)$$

A curva de coexistência líquido-vapor do campo de força SAFT- $\gamma$  CG foi comparada com os resultados disponíveis na literatura para o campo de força TraPPE-EH (Rai e Siepmann, 2013), e as demais propriedades foram avaliadas por comparação com dados experimentais (Mortimer e Murphy, 1923; Nelson e Senseman, 1922; Linstrom e Mallard, 2017). A equação de desvio relativo absoluto médio usada possui a seguinte forma:

$$\Delta X = \frac{1}{N_p} \sum_{i=1}^{N_p} \left| \frac{X_{i^{ref}} - X_{i^{calc}}}{X_{i^{ref}}} \right| 100\%, \quad (13)$$

$X^{ref}$  é o valor da propriedade de referência e  $X^{calc}$  é o valor da propriedade calculado.

## Resultados e Discussão

Os parâmetros do fenantreno com três segmentos para o campo de força SAFT- $\gamma$  CG estimados com a metodologia descrita estão expostos na Tabela 1.

Tabela 1. Parâmetros do campo de força SAFT- $\gamma$  CG para o fenantreno.

$\sigma^{sim} / \text{\AA}$	$(\varepsilon^{sim} / \kappa_b) / \text{K}$	$\lambda_r^{sim}$
4,008	529,646	14,339

A Figura 2 mostra a pressão de vapor do fenantreno em função de temperatura. O campo de força com parâmetros estimados conseguiu apresentar um comportamento similar ao observado nos dados experimentais, considerando a sua simplicidade e eficiência computacional. O valor da pressão de vapor para temperaturas mais próximas do ponto normal de ebulição foi subestimado, enquanto houve uma superestimação dos valores de pressão de vapor para temperaturas mais baixas. Esse mesmo comportamento e o desvio relativo da pressão de vapor, exposto na Tabela 2, foram similares aos resultados observados para os cálculos de propriedades de equilíbrio e transporte do dióxido de carbono e metano com o campo de força SAFT- $\gamma$  CG em trabalhos anteriores (Aimoli *et al.*, 2014; Aimoli, 2015). Observa-se, também, que os resultados obtidos ficaram próximos dos resultados do campo de força TraPPE-EH disponíveis na literatura, o que indica que a simplificação do modelo acarreta somente em pequenos desvios no cálculo da pressão de vapor. O desvio relativo médio está disponível na Tabela 2.

Tabela 2. Desvio relativo médio entre os resultados obtidos com o campo de força SAFT- $\gamma$  CG e os dados experimentais.

	$\Delta / \%$
$P_v$	9,01

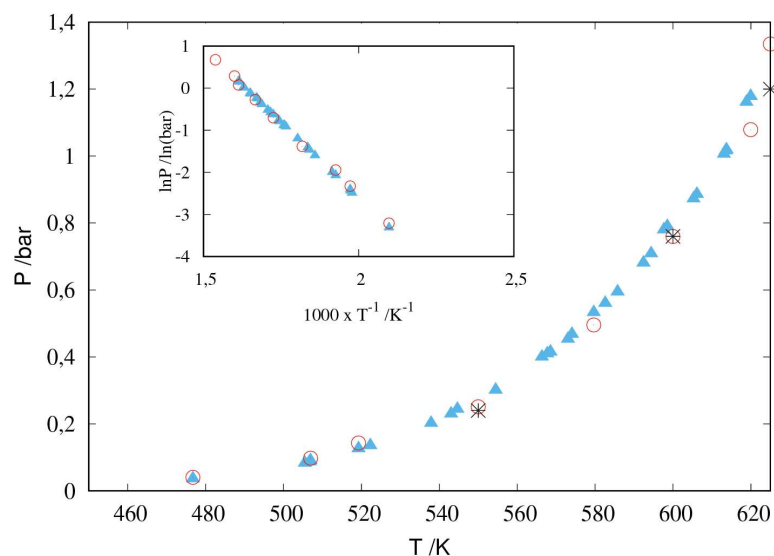


Figura 2. Pressão de vapor em função da temperatura calculada com o campo de força SAFT- $\gamma$  CG (círculos vermelhos), dados experimentais (triângulos azuis) e resultados do campo de força TraPPE-EH (asteriscos pretos). A figura inserida é o mesmo gráfico, porém no formato de Clausius-Clapeyron.

Na Figura 3, as curvas de densidade de líquido e vapor saturados obtidas para o campo de força SAFT- $\gamma$  CG foram comparadas com os dados disponíveis do TraPPE-EH. O campo de força estimado apresentou comportamento razoavelmente similar, mas com discrepâncias maiores do que aquelas observadas para os resultados de pressão de vapor. Adicionalmente, a densidade do fenantreno foi calculada fora do ELV no ensemble NPT-MC a uma temperatura de 298 K e a uma pressão de 1 bar como forma de comparar os resultados com o valor experimental disponível que é igual 1180 kg/m<sup>3</sup> (Linstrom e Mallard, 2017). O campo de força TraPPE-EH forneceu um valor de densidade igual 1110 kg/m<sup>3</sup> e o campo de força SAFT- $\gamma$  CG forneceu um valor igual a 1455 kg/m<sup>3</sup>. Esses resultados mostram que o campo de força SAFT- $\gamma$  CG tem mais dificuldade em prever propriedades em estados diferentes daquele que foi estimado do que o TraPPE-EH por ser um modelo coarse-grained top down. Então, essa diferença na densidade pode ter sido causada pelo uso de apenas a pressão de vapor na estimação dos parâmetros da equação de estado tida como base.

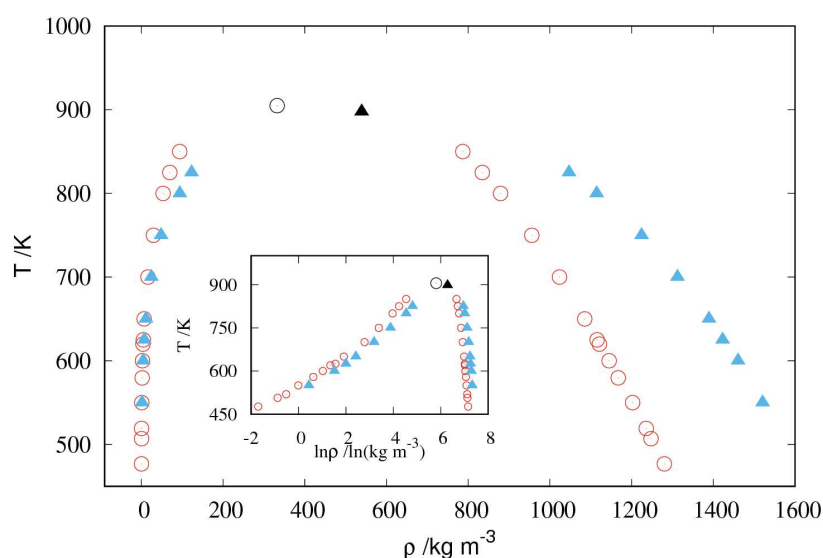


Figura 3. Curva de coexistência líquido vapor calculada com campo de força SAFT- $\gamma$  CG (círculos vermelhos) e com o campo de força TraPPE-EH (triângulos azuis). O círculo e o triângulo pretos representam as propriedades críticas calculadas com o ajuste.

Os desvios relativos da densidade de líquido e de vapor saturados e das propriedades críticas encontradas com o ajuste das Equações 12-13 estão resumidos na Tabela 3. Os desvios deixam clara a existência de uma diferença significativa para densidade no equilíbrio ocasionados pela diferença de rigor teórico entre os modelos. Essas discrepâncias talvez possam ser reduzidas com uso dos dados da densidade do campo de força TraPPE-EH para estimação inicial, o que permitiria um melhor ajuste do expoente atrativo do potencial de Mie. Já com relação à previsão da temperatura crítica, os resultados entre os campos de força foram mais próximos.

Tabela 3. Propriedades críticas e desvios relativos entre o campo de força SAFT- $\gamma$  CG e os dados de referência para o fenantreno.

$\Delta / \%$			
$T_c$	$\rho_c$	$\rho_{liq}$	$\rho_{vap}$
4,08	38,17	21,42	38,64

## Conclusão

O campo de força SAFT- $\gamma$  CG foi obtido para o fenantreno com uma metodologia em que se usa a equação de estado SAFT-VR Mie como base para parametrização desse campo de força. Os parâmetros encontrados foram avaliados através de cálculos das propriedades de equilíbrio com o método GEMC-NVT. Os resultados para pressão de vapor tiveram concordância com os dados experimentais e com o resultados da literatura obtidos para o campo de força atomístico TraPPE-EH. As densidades de líquido e vapor no equilíbrio apresentaram maiores desvios em relação ao TraPPE-EH do que a pressão de vapor. Isso mostra que a simplificação da molécula não representou uma grande perda na representação da pressão de vapor do fenantreno e que esse modelo de campo de força pode ser uma alternativa a modelos atomísticos. Com esse campo de força avaliado, pretende-se fazer cálculos de energia de solvatação do fenantreno em tolueno e em soluções tolueno+CO<sub>2</sub> utilizando-se dinâmica molecular.

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# ANNEX B – Paper for Publication in Scientific Journal

# Evaluation of the SAFT- $\gamma$ Mie force field with solvation free energy calculations

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## Abstract

Solvation free energies can be essential in the process of evaluating and developing force fields. In addition to that, it is possible to obtain a diversity of thermodynamic informations using solvation free energies. Hence, we aim the solvation free energies of molecules mimicking asphaltenes in aqueous and non-aqueous solvents with a coarse-grained model known as SAFT- $\gamma$  Mie force field. This model uses a top-down parameterization in which the force field parameters are obtained using a equation of state. The use of solvation free energy calculations to evaluate this force field can then help to improve this force field and increase the scale of these simulations. The solvation free energies presented in this article were obtained by carrying out molecular dynamics simulations using the expanded ensemble method. The output of these simulations was then used to estimate the free energy differences. For this, we employed the MBAR method. The results with solvents other than water had low absolute deviations from experimental data. In turn, hydration free energy calculations required a binary interaction parameter estimated with output data from molecular dynamics in order to obtain accurate free energy differences. These results indicated problems on the description of the water molecule by the SAFT- $\gamma$  Mie force field, but, generally, proved that this coarse-grained model could represent the solvation free energies of the studied solute-solvent pairs.

**Keywords:** solvation free energies, SAFT- $\gamma$  Mie

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## 1. Introduction

Solvation free energy calculations with molecular dynamics (MD) have a variety of applications ranging from drug design in the pharmaceutical industry to the development of separation technologies in the chemical industry. Solvation free energy is, more specifically, the difference in free energy related to the process of transferring a solute from an ideal gas phase into a liquid solution [1]. Through the study of the solvation phenomenon, it is possible to obtain information about the behavior of the solvent in different thermodynamic conditions and the influence of the solute's molecular geometry. It is also possible to calculate other important properties with the solvation

free energy, namely the activity coefficient at infinite dilution, Henry constant, and partition coefficients. Additionally, solvation free energy calculations can be part of the methodology of calculating solubility from molecular dynamics.

The solvation free energy calculations described above are intrinsically complex due to the many competing forces interfering in the behavior of the solute-solvent interaction. In addition, free energy simulations are susceptible to sampling problems in low energy regions, and simulation results need to be correctly post-processed in order to yield free energy differences with small uncertainties. Various simulation methodologies were developed to enable estimations of free energy differences such as the expanded ensemble [2], thermodynamic integration [3], free energy perturbation (FEP) [4, 5, 6], and umbrella sampling [7]. Utilizing FEP methodologies, recent

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articles [8, 9] made available a large database of hydration free energies of small molecules using the GAFF force field. Beckstein et al. [10] also calculated the hydration free energies for fifty-two compounds with the OPLS-AA force field. They obtained an overall root mean square deviation to the experimental data of 1.75 kcal/mol. Izairi and Kamberaj [11] studied hydration free energies but with the intention of comparing the polar and nonpolar contributions. Garrido et al. [12, 13] calculated the free energy of solvation of large alkanes in 1-octanol and water with three different force fields (TraPPE, GROMOS, and OPLS-AA/TraPPE). These authors also estimated the solvation free energy of propane and benzene in non-aqueous solvents like n-hexadecane, n-hexane, ethylbenzene, and acetone with the TraPPE-UA and TraPPE-AA force fields. Roy et al. [14] addressed the choice of the Lennard-Jones parameters for predicting solvation free energy of different solutes in 1-octanol. They calculated the solvation free energy of a set of 205 small organic molecules in 1-octanol and found that the force field parametrization of n-octanol proposed by Kobryn and Kovalenko [15] provided the best agreement to the experimental data. Gonçalves and Stassen [16] calculated the free energy of solvation using the polarizable continuum model coupled to molecular dynamics simulation with the GROMOS force field. These calculations were done with a representative set of solutes and with the solvents tetrachloride, chloroform, and benzene. Using the GAFF and the polarizable AMOEBA force fields, Mohamed et al. [17] evaluated the solvation free energy of small molecules in toluene, chloroform, and acetonitrile, and obtained a mean unsigned error of 1.22 kcal/mol for AMOEBA and 0.66 kcal/mol for GAFF. Genheden [18] expanded the ELBA coarse-grained force field to calculate solvation free energies of more than 150 solutes taken from the Minnesota solvation database in polar (water, hexanol, octanol, and nonanol) and apolar (hexane, octane, and nonane) solvents. He obtained mean absolute deviations of 1 kcal/mol for water and 1.5 kcal/mol for hexane. In this model, three carbons are represented by a single bead and water is represented by a single bead with point-dipole.

As can be seen in the previous paragraph, solvation free energy simulation is performed in the literature using a variety of force field since the choice of force field can be another influencing factor in the output of these calculations.

Hence, we, in this study, assess the efficiencies and shortcomings of the SAFT- $\gamma$  Mie coarse-grained force field [19] with solvation free energy calculations for a variety of pairs solute-solvent. We chose a coarse-grained force field because these kind of models generally reproduce free energy differences since the effects of reducing degrees of freedom in the entropy are counterbalanced by the reduction of enthalpic terms [20]. This fact makes these models a viable option to decrease the computational time of solvation free energy calculations. Additionally, deficiencies in the description of small molecules by coarse-grained models can be revealed by free energy calculations [21, 1]. We specifically picked the SAFT- $\gamma$  Mie force field because it uses, unlike the majority of the force fields, the Mie potential and because its method of obtaining parameters is more straightforward than other coarse-grained models. the SAFT- $\gamma$  Mie force field was initially parameterized with pure component equilibrium and interfacial tension data [19], and this strategy has provided satisfactory results. Examples include the prediction of phase equilibrium of aromatic compounds [22], alkanes, light gases [23], and water [24], thermodynamic properties of carbon dioxide and methane [25], multiphase equilibrium of mixtures of water, carbon dioxide, and n-alkanes [26], and water/oil interfacial tension [27].

We selected the solvents and solutes in our free energy calculations with the intention of testing the force field with standard sets used as a benchmark in solvation free energy calculations and with polycyclic aromatic substances used as models to asphaltenes. Asphaltenes are complicated to characterize by determining their composition on a molecular basis, but the literature broadly accepts that they can be described as a fraction of crude oil soluble in toluene and insoluble in n-alkanes (pentane, hexane, heptane) [28]. They have motivated many studies with interest in developing models for their structure and behavior due to all the problems they can cause during their transportation and refining such as precipitation during the oil processing [29]. This precipitation issue is a recurrent problem due to the growing market of the production of crude oil in deep waters, whose conditions are favorable to precipitation [30]. As an example, asphaltene precipitation due to pressure drop can clog oil production equipment and cause a growth in the cost of production [31]. All these factors make the understanding of the behavior of asphaltenes in different chemical



and physical environments relevant to the oil industry. As we said, asphaltene characterization still faces some issues. Hence, we choose to use polycyclic aromatic hydrocarbons (PAHs), which have well-defined characteristics, to initially test the efficiency of the SAFT- $\gamma$  Mie force field in describing the solvation phenomenon. PAHs are a group of organic compounds that have fused rings, carbon and hydrogen in their structure [32]. The ones utilized in this work were phenanthrene, anthracene, and pyrene since they share similarities with asphaltenes regarding their solubility. In this context, we selected compounds that are used to characterize asphaltenes (toluene, hexane) as solvents in our free energy calculations. We also tested the anti-solvent/solvent effect of carbon dioxide due to its influence in asphaltene precipitation during the oil processing [33]. With these calculations of solvation free energies with the SAFT- $\gamma$  Mie model, we intend to improve this force field and provide satisfactory solvation free energy estimates of PAHs with a coarse-grained model. The success of the description of small asphaltene-like compounds by this force field can then open up the possibility of obtaining satisfactory results for more complex asphaltene models with a force field with a low computational cost.

## 2. Computational Methods

### 2.1. SAFT- $\gamma$ Mie Force Field

The SAFT- $\gamma$  Mie force field uses a top-down coarse-graining methodology in its parameterization. This methodology aims to obtain the intermolecular parameters from macroscopic experimental data such as fluid-phase equilibrium or interfacial tension data. The idea is that the force field parameters estimated with the SAFT-VR Mie EoS [34] can be used in molecular simulations since both the equation of state and the force field use the Mie potential as the intermolecular potential model:

$$U_{Mie}(r) = \epsilon \frac{\lambda_r}{\lambda_r - \lambda_a} \left( \frac{\lambda_r}{\lambda_a} \right)^{\left( \frac{\lambda_a}{\lambda_r - \lambda_a} \right)} \left[ \left( \frac{\sigma}{r} \right)^{\lambda_r} - \left( \frac{\sigma}{r} \right)^{\lambda_a} \right]. \quad (1)$$

The parameter  $\epsilon$  is the potential well depth,  $\sigma$  is the segment diameter,  $r$  is the distance between the spherical segments,  $\lambda_r$  is the repulsive exponent, and  $\lambda_a$  is the attractive

exponent. This force field has the advantage of incorporating the degrees of freedom provided by the use of the Mie Potential [23]. This flexibility offers the exploration of a vast parameter space without using an iterative simulation scheme [19]. Despite these advantages, the force field can be restricted by the shortcomings of the equation of state. As an example, the lack of an association term in the equation can cause an inadequate representation of the properties of hydrogen bonding compounds.

Each substance has initially five parameters to be estimated ( $m_s$ ,  $\sigma$ ,  $\epsilon$ ,  $\lambda_r$ , and  $\lambda_a$ ) according to Eq. 1. The number of segments is usually fixed in an integer value so as this parameters can be used in molecular simulations. The attractive parameter is generally fixed due to its high correlation with the repulsive parameter. Usually, the chosen value for this parameter is 6, corresponding to the London model, which is a good representation of the dispersion scale of most simple fluids that do not have strong polar interactions [35, 23]. There are two strategies to obtain these parameters: one is by fitting the SAFT-VR Mie EoS to experimental data such as vapor pressure and liquid density [36], and the other one is by using corresponding states parametrization [37]. Here, the first strategy was used to find the parameters for phenanthrene with vapor pressure data [38, 39] following the methodology proposed by Müller and Mejía [22]. The parameterization was carried out with the number of segments equal to five and with a geometry such as that in Figure 1, since this level of coarse-graining was also used for a similar molecule (anthracene) in the original paper. We also show in Figure 2 the vapor pressure curve obtained using the EoS with the parameters estimated here.



Figure 1: Coarse-graining level and geometry chosen for phenanthrene.

The parameters for the other compounds were retrieved from the literature, and all these parameters are exposed in Table 1. Only when modeling a mixture with the SAFT- $\gamma$  Mie force field, the use of combining rules is necessary since the segments are equal. The mixing rules for this

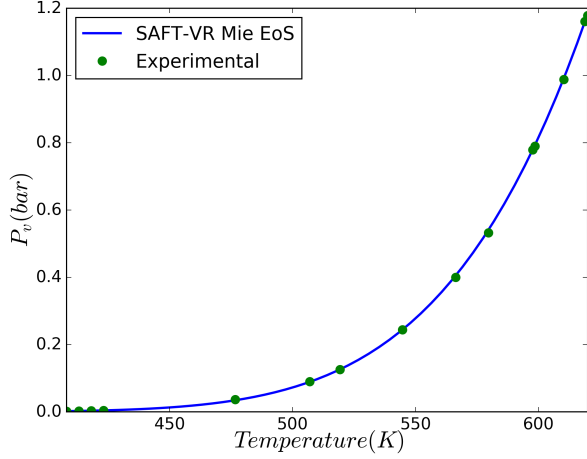


Figure 2: Vapor pressure of phenanthrene ( $P_v$ ) calculated using the SAFT-VR Mie EoS with the parameters estimated here.

force field can be seen in Eqs. 2 to 4 [34].

$$\sigma_{ij} = \frac{\sigma_{ii} + \sigma_{jj}}{2}, \quad (2)$$

$$\lambda_{k,ij} - 3 = \sqrt{(\lambda_{k,ii} - 3)(\lambda_{k,jj} - 3)}, \quad k = r, a, \quad (3)$$

$$\epsilon_{ij} = (1 - k_{ij}) \frac{\sqrt{\sigma_{ii}^3 \sigma_{jj}^3}}{\sigma_{ij}^3} \sqrt{\epsilon_{ii} \epsilon_{jj}}, \quad (4)$$

The  $k_{ij}$  is a binary interaction parameter to correct the deviations of the mixing rule. This parameter can be described as scaling factor. It accounts for the interactions among chemically distinct compounds, which are not explicitly considered by the SAFT-VR Mie EoS. These mixing rules were the ones available in the literature and employed by other papers that used this force field. Therefore, we ended up using Eqs. 2 to 4 in our study. However, the binary interaction parameter was only necessary for aqueous mixtures in our study.

## 2.2. Expanded Ensemble

The strategy chosen in this work to calculate the solvation free energies was to use an alchemical method in which the solute molecule is gradually inserted in the solvent using a thermodynamic path [41]. Each insertion or

Table 1: SAFT- $\gamma$  Mie Force Field for each substance used in this work.

	$m_s$	$\epsilon/\kappa_b$ (K)	$\sigma$ (Å)	$\lambda_r$
Water [26]	1	305.21	2.902	8.0
Propane [23]	1	426.08	4.871	34.29
Carbon dioxide [23]	2	194.94	2.848	14.65
Hexane [23]	2	376.35	4.508	19.57
Octanol [40]	3	495.71	4.341	28.79
Toluene [22]	3	268.24	3.685	11.80
Benzene [22]	3	230.30	3.441	10.45
Pyrene [22]	4	459.04	4.134	15.79
Anthracene [22]	5	259.68	3.631	9.55
Phenanthrene	5	262.74	4.077	9.55

alchemical state is represented by a coupling parameter,  $\lambda$ , that ranges from 0 to 1. When  $\lambda = 0$ , there is no interaction with the solvent and, when  $\lambda = 1$ , the interactions are fully activated. Since the force field used does not explicitly take in consideration the charges, the interactions are only due to the Mie potential. For the coupling of the Mie Potential, we propose a generalized softcore Mie potential based on the softcore potential of Beutler et al. [42]:

$$U_{Mie}^{sc}(r) = \lambda \epsilon \frac{\lambda_r}{\lambda_r - \lambda_a} \left( \frac{\lambda_r}{\lambda_a} \right)^{\left( \frac{\lambda_a}{\lambda_r - \lambda_a} \right)} \cdot \left\{ \frac{1}{[\alpha(1 - \lambda) + (r/\sigma)^{\lambda_a}]^{\lambda_r/\lambda_a}} - \frac{1}{\alpha(1 - \lambda) + (r/\sigma)^{\lambda_a}} \right\}. \quad (5)$$

where  $\alpha$  is a constant whose value is normally assumed to be 0.5. Instead of performing various simulations at each alchemical state to obtain the potential energies of Eq. 5, we decided to use the Expanded Ensemble method [2] since it allows a non-Boltzmann sampling scheme of different states in a single simulation. In this scheme, the sampling is done by biasing the phase space exploration process with weights not related to the statistical ensemble. The partition function of the statistical expanded ensemble,  $Z^{EE}$ , is obtained from the probability distributions corresponding to each  $\lambda$ . Hence,  $Z^{EE}$  is defined as a sum of subensembles  $Z_i$  in different values of  $\lambda$ , that is,

$$Z = \sum_{i=1}^N Z_i \exp(\eta_i), \quad (6)$$

where  $N$  is the number of alchemical states,  $\eta_i$  is the arbitrary weight of the subensemble at each state, and  $Z_i$  is the configurational partition function of state  $i$ . Here, we followed the flat-histogram approach [43, 44, 45] to calculate the weights. This strategy aims to obtain adequate sampling by ensuring that all the states have an equal number of visits, i.e. the ratio of the probability of sampling state  $i$  ( $\pi_i$ ) to the probability of sampling state  $j$  ( $\pi_j$ ) is equal to one. Using this relation, the following equation can be obtained:

$$(\eta_i - \eta_j)_{k+1} = \beta(G_i - G_j)_k. \quad (7)$$

Eq. 7 proposes that the choice of weights is dependent on the free energies that we are attempting to obtain. This equation is then solved iteratively with trial simulations. For the first simulation, the values of  $\eta$  are set to zero, and the histogram of the states visited is obtained. With this histogram, it is possible to estimate the free energy differences and, since the weights are related to the free energies by Eq. 7, the next values of  $\eta$  can be calculated. This iteration goes on until a uniform distribution is attained. The weights found are then used in a longer simulation to obtain the final solvation free energies. The choice of the  $\lambda$  set corresponding to overlapping alchemical states are crucial to acquire accurate free energy differences. In this work, the method chosen to obtain the optimal staging of the  $\lambda$  domain is the one developed by Escobedo and Martinez-Veracoechea [46] with a basis in the study of Katzgraber et al [47]. This method targets "bottlenecks" in the simulation. It does that by optimizing  $\lambda$  through the minimization of the number of round trips per CPU time between the lowest (0) and highest (1) values of  $\lambda$ . The optimization is specifically done by maximizing the steady-state stream  $\phi$  of the simulation, which "walks" among the values of  $\lambda$ . This flow is estimated from a Fick's diffusion type of law:

$$\phi = D(\Lambda)\Pi(\Lambda)\frac{dx(\Lambda)}{d\Lambda}. \quad (8)$$

In the equation above,  $\Lambda$  is the actual continuous value of the coupling parameter. This continuous function of  $\lambda$ 's is obtained by interpolating the  $\lambda$  set linearly.  $D(\Lambda)$  is the diffusivity at state  $\Lambda$  and  $x(\Lambda)$  is the fraction of times that the trial simulation at state  $\Lambda_i$  has most recently visited the state  $\lambda = 1$  as opposed to state  $\lambda = 0$ . The

derivative  $dx(\Lambda)/d\Lambda$  is approximated with the central finite differences method. Finally,  $\Pi(\Lambda)$  is the probability of visiting  $\Lambda$ :

$$\Pi(\Lambda) = \frac{C' \bar{\Pi}(\lambda)}{\Lambda_{i+1} - \Lambda_i}. \quad (9)$$

The  $C'$  term in the equation above represents a constant and  $\bar{\Pi}(\lambda)$  is the arithmetic average of the frequency of visits to the  $\Lambda$  state:

$$\bar{\Pi}_i(\lambda) = \frac{\pi_{i+1} - \pi_i}{2}. \quad (10)$$

The  $\phi$  is maximum when the optimal probability  $\Pi'(\Lambda_i)$  of visiting state  $\Lambda_i$  is proportional to  $1/\sqrt{D(\Lambda)}$  [48]. With that information, it is possible to estimate the diffusivity using one trial simulation with the following equation:

$$D(\Lambda) = \frac{\Lambda_{i+1} - \Lambda_i}{\bar{\Pi}(\lambda)dx(\Lambda)/d\Lambda}. \quad (11)$$

Hence, we can calculate  $\bar{\Pi}$  and, consequently, the cumulative probability, which is used to obtain the new  $\lambda$  state, by

$$\Phi = \int_{\lambda=0}^{\lambda=1} \Pi'(\Lambda_i)d\Lambda = \frac{i}{K}, \quad (12)$$

where  $K$  is the total number of  $\lambda$  states. In order to carry out our solvation free energy simulations, we obtained these cumulative probabilities for every  $\lambda$  set we estimated. A graphical demonstration of the relation between the optimized coupling parameters and the cumulative probability of Eq. 12 is presented in our results in Figure 3.

### 3. Molecular Dynamics Simulations

Using the parameters of Table 1, we carried out molecular dynamics simulations to obtain the potential energies at each alchemical state. The chosen software package to perform the simulations was LAMMPS [49]. In this package, the equations of motion were integrated with the velocity-Verlet algorithm [50] with a time step of 2 fs. As required by the coarse-grained model, molecules with more than one bead were treated as rigid bodies. The thermostat and the barostat were the Nosé-Hoover chains as described in Hoover [51] and Martyna et al. [52] with

damping factors of 100 and 1000 time steps, respectively. For the rigid bodies in our simulations, we used the rigid-body algorithm of Kamberaj et al [53]. The potential cut-off was equal to 20 Å [22] with a neighbor list skin of 2 Å. The initial configurations of the solvated systems were also generated using the Playmol package [54], which is integrated with the Packmol package [55]. For the binary mixtures, one molecule of solute and a varying number of solvent molecules- 700 molecules of toluene, 700 molecules of octanol, 1024 molecules of hexane, 3000 molecules of water - were randomly added to a cubic box. Besides the systems with pure substances acting as solvents, we performed simulations to study the solvation free energy of phenanthrene in a mixture of toluene and carbon dioxide with different weight fractions ( $w_{CO_2}$ ). The system consisted of one molecule of phenanthrene for all the cases and 123 molecules of  $CO_2$  and 618 molecules of toluene ( $w_{CO_2} = 0.087$ ); 166 molecules of  $CO_2$  and 589 molecules of toluene ( $w_{CO_2} = 0.119$ ); 232 molecules of  $CO_2$  and 545 molecules of toluene ( $w_{CO_2} = 0.169$ ); 380 molecules of  $CO_2$  and 446 molecules of toluene ( $w_{CO_2} = 0.289$ ). These substances used in our study were selected with the intention of testing the force field with standard sets used as a benchmark in solvation free energy calculations, with aromatic substances used as models to asphaltenes and with water, which probably is the most used solvent in computational studies.

All simulations were performed with the constant temperature and pressure values of 298 K and 1 bar, except the ones containing carbon dioxide. These had the temperature of 298 K and the pressure of the experimental liquid-phase equilibrium corresponding to each composition of the system  $CO_2$ +toluene [56]. For all simulations, the initial box was equilibrated at the NPT ensemble for 2 ns, and the resulting configurations were used as the initial configuration of the expanded ensemble simulations. These were carried out with the LAMMPS user package for expanded ensemble simulations with the Mie potential developed by our research group, available at <https://github.com/atoms-ufjrj/USER-ALCHEMICAL>.

During these expanded ensemble simulations, the sampling of a new alchemical state was tried at every 10 MD steps. To define the optimal values of  $\lambda$  and  $\eta$  corresponding to each state, trial simulations, having around 9 ns of production time, were carried out. In the first simulation, we chose the group of  $\lambda$  values arbitrarily, and we either

set all  $\eta$ 's to zero or assigned values previously found for similar solute-solvent pairs. The subsequent group of  $\eta$ 's were estimated with the flat histogram approach (Eq. 7). We then performed another trial simulation with the new weights. The results of this simulation were used to optimize the group of  $\lambda$ 's by minimizing the number of round trips, as described in the preceding section. The  $\eta$ 's corresponding to the newest group of  $\lambda$ 's were interpolated linearly from the free energy differences. With the final values of  $\eta$  and  $\lambda$  defined for each mixture, larger simulations with a production time of 20 ns were carried out.

Since the employed force field considers that the beads do not have charges, there are no Coulombic interactions, and the only contribution to the total potential energy is due to the softcore potential of Eq. 5. The post-processing method used to effectively calculate free energy differences with the potential energies obtained from the expanded ensemble simulations was the Multistate Bennett Acceptance Ratio (MBAR) method. The software alchemical-analysis [41] was utilized to obtain the  $\Delta G_{solv}$  with MBAR and to assess the quality of the results. After the first estimations, we realized that the binary interaction parameter of Eq. 4 was necessary for systems containing water. Hence, we estimated  $k_{ij}$  for these pairs and, for all the other pairs, we set  $k_{ij}$  to zero. The estimation was done by performing trial expanded ensemble simulations in three values of  $k_{ij}$ , as suggested by Ervik et al. [57]. With the  $\Delta G_{solv}$  obtained with these simulations, we did a linear fit to obtain the refined value of the parameter. We used this strategy because the estimation with SAFT-VR Mie EoS gave poor results for the hydration free energies.

## 4. Results and discussion

### 4.1. Solvation free energies

Our primary intention with this study is to assess the capability of the SAFT- $\gamma$  Mie force field to represent solvation free energies. Hence, we chose benchmark solutes used in the literature (benzene, propane) and aromatic solutes (benzene, pyrene, phenanthrene, anthracene), and, for the solvents, we picked non-polar (hexane), aromatic (toluene), and hydrogen bonding (1-octanol, water) substances. The solvation free energy simulations for the pairs chosen were carried out with binary interaction parameters equal to zero since these parameters were not

necessary according to our preliminary studies. Since the force field does not account for charges, we only calculated the Mie contribution (Eq. 5) to the solvation free energy. A total of 15 to 18  $\lambda$ 's, depending on the solute-solvent pairs, and their respective  $\eta$ 's were estimated as described in the previous sections. The simulations carried out using these optimized weights deviated from the flat-histogram requirement of equal number visits by an average of 5% for all of the pairs solvent+solute. The final  $\lambda$  set for all the pairs was found using the cumulative probability distribution (Eq. (12)). For the hexane(solvent)+benzene(solute) pair, the probability distribution can be seen in Figure 3. From now on we are going to use the terminology solvent+solute. The optimized values of  $\lambda$  and  $\eta$  for this pair and all the other pairs are exposed in Tables A.6 to A.9, available in the Appendix. By observing the coupling parameters found for all the pairs, we can see that they are concentrated on the region with a steeper slope as it is expected in this method. After the expanded ensemble simulations with the optimized intermediate states and weights, we calculated the solvation free energies with MBAR. The results obtained and the absolute deviations to experimental data [58] are available in Table 2. The numerical values for solvation free energies in hexane had overall smaller absolute deviations from experimental data than the deviations in the other solvents. Additionally, this force field presented better results for the pair hexane+benzene than the TraPPE force field ( $-4.35 \pm 0.05$  kcal/mol) [13] and the ELBA coarse-grained force field ( $-2.92 \pm 0.01$  kcal/mol) [18]. It's also important to point out that the solvation free energies with these force fields were obtained with thermodynamic integration, which is different from the method used here. TraPPE is a force field parametrized with fluid-phase equilibria data that uses the Lennard-Jones potential to describe the non-bonded interactions. In the cited paper, they used the united-atom description of the TraPPE force field for the alkyl group, the all-atom description for the polar groups and the explicit-hydrogen approach for the aromatic groups. In the explicit-hydrogen approach, the interaction sites for all hydrogen atoms, some lone pair electrons, and bond centers are accounted for [59]. In turn, the ELBA force field is a coarse-grained model that comprises six independent parameters. This force field models three carbons as one Lennard-Jones site and one water molecule as a single Lennard Jones site with a

point-dipole. We also present the solvation free energies corresponding to each alchemical state ( $\lambda$ ) for all the pairs studied here in Figures 4 to 6. Specifically observing the solvation free energy in hexane (Figure 4), we can see the effect of the molecule's size on the entropic region of the free energy curve. This region is corresponding to the first values of  $\lambda$  where the cavity required for the insertion of the solute is formed.

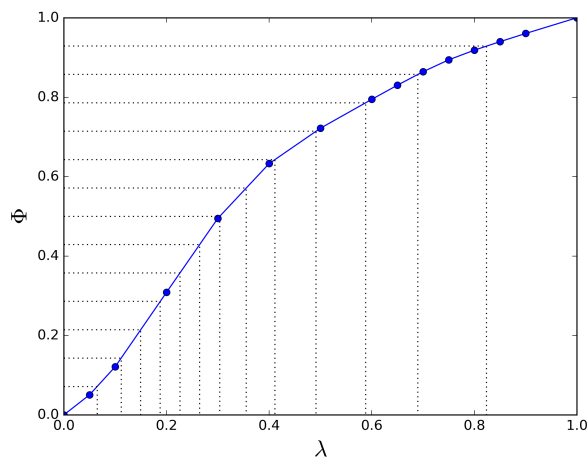


Figure 3: Cumulative probability used to obtain the optimized values of  $\lambda$ 's for the pair hexane+benzene.

We expected that a force field based on an EoS that does not explicitly account for hydrogen bond would not perform well for 1-octanol in mixtures since the parameterization of this molecule did not explicitly account for the interactions of association. All the beads representing 1-octanol have the same intermolecular parameter, and there is no distinction between the polar and apolar groups. Despite this, the solvation free energies of propane and phenanthrene in 1-octanol lied in the desired deviation range of 1-2 kcal/mol [60]. For propane, the observed deviation in solvation free energies was much smaller when compared to the other solutes, which can be attributed to the non polarity of propane and its smoother free energy curve, presented in Figure 5. Such solvation free energy of propane in 1-octanol also had a smaller deviation than the prediction of the ELBA force field ( $-0.92 \pm 0.01$ ) [18]. The absolute deviation of the solvation free energy computed for anthracene in 1-octanol is much higher than the

Table 2: Calculated and experimental values for the solvation free energies (kcal/mol) of solutes in non-aqueous solvents.

Solute	Solvent	$\Delta G_{solv}^{exp}$	$\Delta G_{solv}^{Mie}$	Absolute Deviation
benzene	hexane	-3.96	-3.76 $\pm$ 0.01	0.20
pyrene	hexane	-11.53	-10.82 $\pm$ 0.02	0.71
phenanthrene	hexane	-10.01	-9.16 $\pm$ 0.01	0.85
propane	1-octanol	-1.32	-1.36 $\pm$ 0.03	0.04
anthracene	1-octanol	-11.72	-8.12 $\pm$ 0.03	3.61
phenanthrene	1-octanol	-10.22	-8.34 $\pm$ 0.03	1.47
pyrene	toluene	-12.86	-11.74 $\pm$ 0.01	1.11
anthracene	toluene	-11.31	-9.90 $\pm$ 0.01	1.41

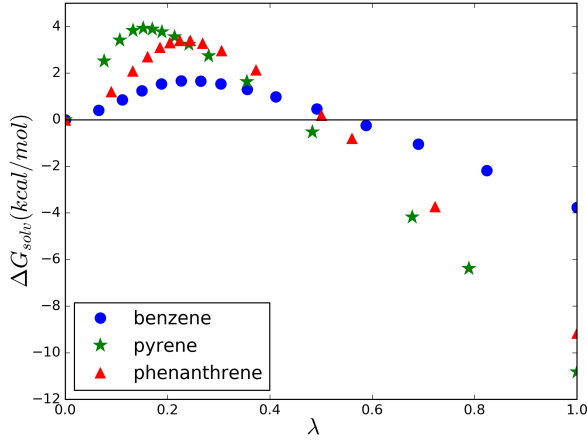


Figure 4: Representation of solvation free energies of different solutes in hexane estimated at each alchemical state.

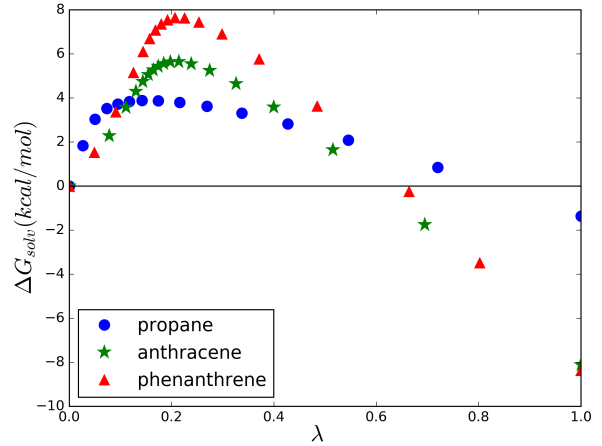


Figure 5: Representation of solvation free energies of different solutes in 1-octanol estimated at each alchemical state.

one calculated for phenanthrene in 1-octanol. The anthracene and phenanthrene molecules have the same geometry (Figure 1) in the SAFT- $\gamma$  Mie model, although anthracene is a linear molecule and phenanthrene is not, and also similar physical properties. Hence, this high deviation of the solvation free energy of anthracene in 1-octanol may indicate a problem in the geometry chosen for anthracene in the SAFT- $\gamma$  Mie force field and the importance of the geometry in modeling the molecules with this force field.

The results also indicate a reasonable capability of the force field for predicting the solvation free energies of

polyaromatic solutes in aromatic solvents. The influence of the molecular geometry on the solvation free energy curves was the same as the one observed for other solvents, as can be seen in Figure 6. We also calculated the  $\Delta G_{solv}$  for phenanthrene in pure toluene and in toluene+ $CO_2$  mixtures. To the best of our knowledge, there are no available experimental data for these solvation free energies, but the previous results for phenanthrene in other solvents showed that the force field is adequate to describe the solvation phenomenon of phenanthrene in a pure aromatic solvent. Hence, we decided to carry out a qualitative study of the influence of  $CO_2$  in the

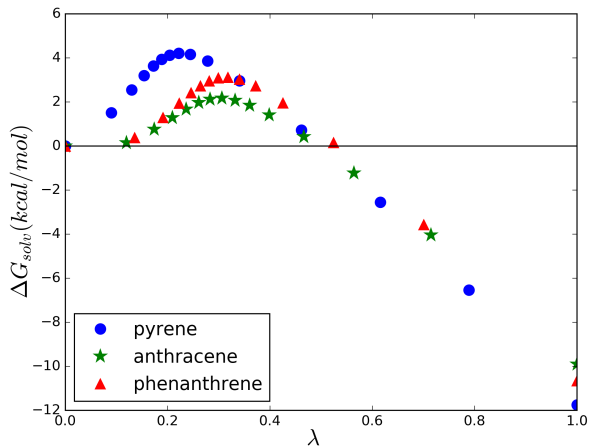


Figure 6: Representation of solvation free energies of different solutes in toluene estimated at each alchemical state.

solvation free energies of phenanthrene in toluene in order to evaluate the description of this system with the SAFT- $\gamma$  Mie force field. The results for these sets are exposed in Table 3. The increase of the mass fraction of  $CO_2$  in toluene caused a small effect on the solvation free energies in the range of weight fractions (0.087-0.289) studied in this dissertation. First, the  $\Delta G_{solv}$  decreased with the increase of  $w_{CO_2}$  up to 0.119. After this, the effect was reversed, and carbon dioxide became an anti-solvent. Soroush et al. [33] reported that asphaltene precipitation occurs when carbon dioxide mass fractions became higher than 0.10 in the system asphaltene+toluene+carbon dioxide, which is in agreement with the anti-solvent effect of carbon dioxide observed in the values calculated here. Although we noticed the anti-solvent effect, the differences observed are pretty small. These minor differences may indicate that the effect of  $CO_2$  is negligible in the solvation of phenanthrene in toluene when using the SAFT- $\gamma$  Mie force field to model the molecules. Nevertheless, more studies need to be done to make a safe assertion about it. It is also worth remarking that this is a qualitative study due to the lack of experimental data. Overall, the methodology proposed by the SAFT- $\gamma$  Mie force field was satisfactory in predicting the solvation free energies of the pairs solvent-solute studied here.

Table 3: Calculated values for the solvation free energies (kcal/mol) of phenanthrene in toluene+ $CO_2$ .

$w_{CO_2}$	$\Delta G_{solv}^{Mie}$
0.0	$-10.65 \pm 0.02$
0.087	$-10.73 \pm 0.02$
0.119	$-10.78 \pm 0.02$
0.169	$-10.71 \pm 0.02$
0.289	$-10.69 \pm 0.02$

#### 4.2. Hydration free energies

Water is a solvent extensively used in experimental and computational studies. Because of this importance and the fact that water has unique properties, such as density maximum at 277 K and increased diffusivity upon compression, developing an accurate computational model for water is an ongoing quest [61]. Hence, we also calculated the solvation free energies in water (hydration free energies) with the SAFT- $\gamma$  Mie force field. With these calculations, we intend to verify if this coarse-grained model would represent the water molecule correctly and would be a good alternative to decrease the computational cost of solvation studies with asphaltene models. The simulations with water as a solvent were carried out using widely studied solutes (propane) and aromatic solutes (benzene, toluene, and phenanthrene) with a set of fifteen intermediate states. We obtained these sets of  $\lambda$  and  $\eta$  with the same methodology used to acquire the sets for the solvation free energies with non-aqueous solvents, and they are exposed in Table A.10, available in the Appendix. At first in our simulations, the binary interaction parameters of all aqueous mixtures were set to zero, but preliminary results for hydration free energies, displayed in Table 5, exhibited a high deviation from experimental data [62, 63]. With these results, the need for binary interaction parameters became clear. First, we estimated  $k_{ij}$  with the SAFT-VR Mie EoS and experimental vapor pressure data, but this strategy also provided results that had high absolute deviations from the experimental data. Therefore, we used the approach of estimating  $k_{ij}$  with the output from solvation free energy calculations with molecular dynamics, as described in the last paragraph of Section 3. We initially found individual values for the interaction parameter of each pair, but, since the parameters for aromatic solutes were very similar (0.148, 0.162, 0.152), we aver-

Table 4: Binary interaction parameters employed.

Pair	$k_{ij}$
water+propane	0.067
water+aromatic	0.154

aged these values. By doing that, we obtained a general parameter for the water+aromatic pairs, which is exposed in Table 4. Also in this table, we display the binary interaction parameter for the pair water+propane.

The relatively large  $k_{ij}$  value of the interaction between aromatic solutes and water can be related to the lack of a term in the SAFT- $\gamma$  Mie to account for the interaction associated with a hydrogen bond substance. Actually, the SAFT-VR Mie has an association term [34], but it was not incorporated in the force field. The SAFT- $\gamma$  Mie model for water [26] has two different temperature-dependent sets of parameters. The parameters utilized in this work were those estimated with experimental interfacial tension data. Hence, we tested the only binary interaction parameter for water+toluene estimated with MD interfacial data available in the literature [27]. Nevertheless, the result also had a high absolute deviation, and this parameter could not be transferred to the calculation of the solvation free energy of toluene in water.

These issues faced by SAFT- $\gamma$  Mie model can also be related to the problems of modeling water with a coarse-grained force field. One of the main difficulties is the choice of which water molecules are going to be represented by which specific beads since water molecules move independently and interact by non-bonded interactions [64, 61]. The SAFT- $\gamma$  Mie water considers that one water molecule corresponds to one bead. This strategy only saves a small amount of simulation time, but it can predict properties at physiological temperatures unlike other more aggressive models such as the MARTINI, which considers that one bead represents various water molecules. In light of all these problems related to modeling the water molecule, the SAFT- $\gamma$  Mie force field appears to be a good alternative when working close to room temperatures, but the necessity of additional parameters estimated with molecular simulation indicates severe flaws in the methodology. This estimation of the binary parameter increased significantly the simulation time required to calculate the hydration free energies, since we

had to carry out three additional simulations for every pair water-solute and then three additional simulations for the three water+aromatic solutes in order to test the averaged binary interaction parameter. If these simulations are necessary for every time a new mixture with water is going to be studied with the SAFT- $\gamma$  Mie force field, the use of this model can become impractical. With this idea in mind, a useful investigation to be made is to check how much other pairs of water+aromatic solute can be modeled using the binary interaction parameter estimated here. Using these binary interaction parameters calculated with data from molecular dynamics, we then obtained the final hydration free energies presented in Table 5.

Hydration free energies calculated using the SAFT- $\gamma$  Mie force field with  $k_{ij} \neq 0$  had low absolute deviations from the experimental data, as expected since the parameters were adjusted to fit these experimental data. Hydration free energies have also been calculated for the pairs studied here by Genheden [18] with the ELBA force field and by Mobley and Guthrie [8] with the GAFF force field for the solutes and with the TIP3P model for water. The GAFF (General Amber Force Field) force field is an all-atom model that consists of bonded and non-bonded parameters and is suitable for the study of a significant number of molecules. In turn, the TIP3P model considers that water is a rigid monomer represented by three interacting sites with non-bonded interactions and Coulombic interactions [65]. Both the GAFF and the TIP3P models use the Lennard-Jones potential to calculate the non-bonded interactions.

Comparing the three aforementioned force fields, the root mean square error (RMSE) of all the pairs tested with the SAFT- $\gamma$  Mie model was 0.24, the RMSE for hydration free energies obtained with the GAFF force field was 0.73, and that for the ELBA coarse-grained force field was 0.44. The difference in between the results of GAFF (-5.26 $\pm$ 0.03) and SAFT- $\gamma$  Mie (-3.47  $\pm$  0.02) force fields is significantly high for phenanthrene, hence the coarse-grained force field with a binary parameter is preferred if the application requires a high level of accuracy. The results also indicated that the SAFT- $\gamma$  Mie model with the binary interaction parameter performed better than the ELBA force field in modeling the solvation phenomenon of the pairs studied in this work, but performed worse with the binary parameter set to zero. This difference in performance occurred despite the fact that both the SAFT- $\gamma$  Mie



Table 5: Calculated and experimental hydration free energy differences (kcal/mol) of solutes in water.

Solute	$\Delta G_{solv}^{exp}$	$\Delta G_{solv}^{Mie}$ $k_{ij} = 0$	Absolute Deviation	$\Delta G_{solv}^{Mie}$ $k_{ij} \neq 0$	Absolute Deviation
propane	$2.00 \pm 0.20$	$1.10 \pm 0.01$	0.90	$2.01 \pm 0.01$	0.01
benzene	$-0.86 \pm 0.20$	$-4.45 \pm 0.03$	3.59	$-1.12 \pm 0.01$	0.26
toluene	$-0.83 \pm 0.20$	$-10.98 \pm 0.30$	10.15	$-0.84 \pm 0.01$	0.01
phenanthrene	$-3.88 \pm 0.60$	$-10.90 \pm 0.04$	7.12	$-3.47 \pm 0.02$	0.41

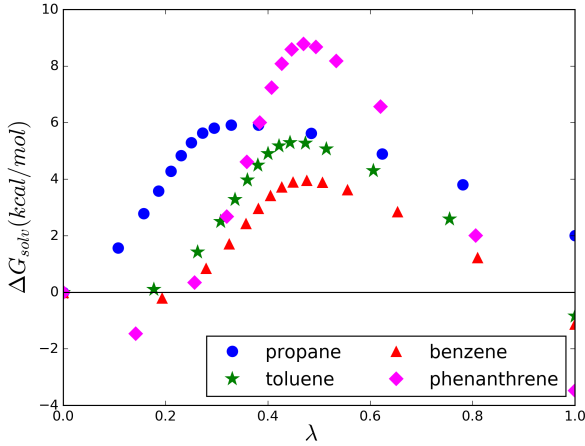


Figure 7: Representation of hydration free energies of different solutes estimated at each alchemical state.

and ELBA models have the same level of coarse-graining for the solvent (one bead represents one water molecule). Therefore, the choice between the two coarse-grained models is dependent on the availability and transferability of binary interaction parameters of the Mie Model. We also present, for the SAFT- $\gamma$  Mie force field, the hydration free energy profiles in Figure 7. Bigger molecules had steeper free energy profiles, as it was for the solvation free energy study in other solvents. We also observe that the hydration free energy for the first non-zero  $\lambda$  is negative for benzene and toluene when a positive value is expected since free energy is required for cavity formation in the solvent for the insertion of the solute. This anomaly can be caused by the fact that the exponential parameters in the Mie potential compensate for cavity formation.

The results found here for both the solvation free energies and hydration free energies fulfilled the intentions

of this dissertation. We assessed the prediction capability of the SAFT- $\gamma$  Mie force field and provided satisfactory solvation free energy estimates of PAHs using a coarse-grained force field. In addition to that, we found flaws in the methodology used by the SAFT- $\gamma$  Mie force field to model the water molecule. Hence, these shortcomings of this model can now be addressed, and the force field can even be improved by using other mixing rules to avoid the use of a binary parameter or, even, using hydration free energy estimates in the parameterization of water. These results also encourage us to calculate solvation free energies of more complex molecules mimicking asphaltenes in non-aqueous solvents in future studies.

## 5. Conclusions

This study consisted in the study of solvation free energy calculations of aromatic solutes in different solvents by using the SAFT- $\gamma$  Mie coarse-grained force field. Solvation free energy studies are mostly done using water as a solvent and with all-atom force fields based on the Lennard-Jones potential. Therefore, with this study, we were able to provide data about the capability of a coarse-grained force field based on the Mie potential in calculating solvation free energies. Additionally, the solvation free energy estimations carried out here can help improve the SAFT- $\gamma$  Mie force field since these calculations are helpful in identifying errors and shortcomings in the modeling process. The SAFT- $\gamma$  Mie uses the SAFT-VR Mie EoS in its parameterization, which results in a relatively straightforward top-down method of obtaining parameters. Following this strategy, the phenanthrene parameters, which were not available in the original database of this force field, were obtained using vapor-liquid equilibrium data.

To perform our expanded ensemble simulations, we optimized the coupling parameters and their respective simulation weights. The resulting potential energies from the expanded ensemble simulations then served as input to estimate solvation free energies with the MBAR method. The results for non-aqueous solvents had absolute deviations from the experimental data of less than 2.0 kcal/mol, except for the pair 1-octanol+anthracene. We also observed the geometry effect on the free energy curves - larger molecules had steeper curves and more substantial absolute deviations. The influence of carbon dioxide on the solvation free energy of phenanthrene in toluene was found to be negligible according to the SAFT- $\gamma$  Mie force field. Hydration free energy calculations with the SAFT- $\gamma$  Mie model required the use of relatively large values of  $k_{ij}$  to produce satisfactory results. We chose to estimate the parameter with the output from molecular dynamics data since the strategy of using the SAFT-VR Mie EoS provided high absolute deviations from the experimental data. This necessity of one additional parameter probably happens due to the lack of a term to account for the hydrogen bond in the EoS on which this force field is based and due to problems associated with the coarse-graining of water molecules. The results obtained with  $k_{ij}$  estimated with MD output were satisfactory, the absolute deviations from the experimental data found were smaller than the ones for the GAFF and ELBA force fields.

Overall, the SAFT- $\gamma$  Mie force field proved to be a suitable model to represent the solvation phenomenon of non-aqueous solvents. It correctly described solvation free energies of solutes mimicking asphaltenes dissolved in hexane, toluene, 1-octanol. However, the calculation of hydration free energies required the use of a binary interaction parameter estimated with MD output, which increased the simulation time significantly. This fact evidenced flaws in the methodology used by the SAFT- $\gamma$  force field and raised questions about the feasibility of this model for hydration free energy calculations. Nevertheless, the SAFT- $\gamma$  Mie force field for water used here does not predict freezing at room temperature as other force fields do, which is essential for our hydration free energy calculations. Therefore, it would be relevant to test if the binary interaction parameter for our aromatic solutes estimated here can be used in hydration free energy calculations of other aromatic solutes and if we could use MBAR to obtain the  $k_{ij}$  through reweighting. Hence,

we would only need to carry out molecular dynamics simulations with one value of  $k_{ij}$ , and then use this output to estimate with MBAR the results with other  $k'_{ij}$ s. We also have some ideas that could be developed in the future using the results from this dissertation. The SAFT- $\gamma$  Mie force field could be used to model larger asphaltene models and, consequently, increase the scale of the simulations we performed. Additionally, it would be a valid investigation to study new methodologies to calculate solubility with solvation free energies using the SAFT- $\gamma$  Mie force field.

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## Appendix A.

Table A.6: Optimized values of  $\lambda$  and  $\eta$  for the hexane+solute pairs.

benzene		pyrene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000
0.065	0.708	0.076	4.234	0.090	1.981
0.112	1.385	0.107	5.620	0.132	3.461
0.15	1.892	0.132	6.499	0.161	4.494
0.188	2.399	0.152	6.690	0.185	5.185
0.226	2.519	0.170	6.643	0.205	5.552
0.264	2.457	0.189	6.461	0.224	5.725
0.304	2.367	0.213	6.091	0.244	5.722
0.356	1.921	0.242	5.566	0.268	5.523
0.411	1.411	0.280	4.729	0.305	4.975
0.492	0.524	0.355	2.853	0.372	3.576
0.588	-0.663	0.483	-0.778	0.500	0.297
0.69	-2.016	0.678	-6.947	0.560	-1.390
0.824	-3.922	0.788	-10.631	0.722	-6.309
1.000	-6.583	1.000	-18.141	1.000	-15.448

Table A.7: Optimized values of  $\lambda$  and  $\eta$  for the 1-octanol+solute pairs.

propane		anthracene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000
0.027	3.126	0.078	3.932	0.049	2.578
0.050	5.109	0.111	6.178	0.091	5.663
0.073	6.093	0.130	7.426	0.125	8.575
0.095	6.570	0.143	8.201	0.144	10.069
0.117	6.826	0.154	8.717	0.157	10.978
0.142	6.956	0.164	9.085	0.169	11.599
0.174	6.969	0.174	9.357	0.180	12.040
0.215	6.847	0.184	9.556	0.192	12.340
0.269	6.554	0.197	9.676	0.206	12.499
0.337	6.050	0.214	9.681	0.225	12.478
0.427	5.228	0.238	9.490	0.253	12.161
0.545	3.955	0.274	8.958	0.298	11.280
0.720	1.843	0.326	7.906	0.371	9.406
1.000	-1.903	0.399	6.088	0.484	5.891
		0.515	2.777	0.664	-0.516
		0.695	-2.960	0.802	-5.908
		1.000	-13.768	1.000	-14.073

Table A.8: Optimized values of  $\lambda$  and  $\eta$  for the toluene+solute pairs.

pyrene		anthracene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000
0.090	2.563	0.119	0.218	0.136	0.726
0.130	4.338	0.174	1.210	0.191	2.307
0.154	5.439	0.209	2.052	0.223	3.430
0.172	6.181	0.236	2.664	0.246	4.233
0.188	6.670	0.261	3.122	0.264	4.780
0.204	6.986	0.283	3.378	0.281	5.149
0.222	7.121	0.306	3.449	0.299	5.354
0.244	7.025	0.332	3.311	0.318	5.389
0.278	6.520	0.360	2.936	0.340	5.222
0.340	5.010	0.399	2.209	0.372	4.717
0.462	1.247	0.466	0.567	0.425	3.440
0.616	-4.283	0.564	-2.211	0.524	0.444
0.788	-11.032	0.715	-6.983	0.701	-5.814
1.000	-19.814	1.000	-16.923	1.000	-17.803

Table A.9: Optimized values of  $\lambda$  and  $\eta$  for the phenanthrene+ $CO_2$ + toluene mixture with different values of  $w_{CO_2}$ .

$w_{CO_2} = 0.087$		$w_{CO_2} = 0.119$		$w_{CO_2} = 0.169$		$w_{CO_2} = 0.289$	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
0.128	0.604	0.128	0.732	0.064	0.883	0.066	0.806
0.184	2.067	0.186	2.223	0.108	0.764	0.111	0.760
0.217	3.164	0.219	3.319	0.175	1.969	0.172	1.983
0.240	3.940	0.244	4.098	0.214	3.156	0.204	2.967
0.260	4.472	0.267	4.704	0.240	3.974	0.227	3.627
0.277	4.823	0.289	5.031	0.258	4.457	0.245	4.082
0.295	5.035	0.313	5.084	0.273	4.750	0.262	4.395
0.318	5.059	0.339	4.950	0.287	4.921	0.279	4.583
0.347	4.762	0.373	4.371	0.305	4.962	0.299	4.621
0.397	3.753	0.425	3.055	0.326	4.885	0.325	4.423
0.491	1.031	0.488	1.196	0.361	4.401	0.365	3.739
0.670	-5.148	0.525	-0.027	0.419	2.990	0.428	2.198
0.791	-9.713	0.730	-7.185	0.527	-0.299	0.530	-0.842
1.000	-18.098	1.000	-17.769	0.697	-6.180	0.701	-6.763
				1.000	-17.998	1.000	-18.163

Table A.10: Optimized values of  $\lambda$  and  $\eta$  for the water+solute pairs.

propane		benzene		toluene		phenanthrene	
$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$	$\lambda$	$\eta$
0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
0.107	2.673	0.193	-0.295	0.177	0.182	0.142	-2.462
0.157	4.703	0.279	1.468	0.262	2.432	0.256	0.597
0.186	6.047	0.324	2.931	0.307	4.244	0.319	4.504
0.210	7.148	0.357	4.168	0.336	5.552	0.358	7.762
0.230	8.017	0.381	5.091	0.360	6.696	0.384	10.104
0.250	8.883	0.405	5.891	0.380	7.558	0.407	12.185
0.272	9.291	0.427	6.443	0.400	8.233	0.427	13.607
0.294	9.700	0.449	6.770	0.422	8.678	0.446	14.490
0.328	9.900	0.476	6.900	0.443	8.859	0.469	14.834
0.381	9.930	0.506	6.805	0.473	8.810	0.494	14.667
0.484	9.463	0.555	6.392	0.514	8.452	0.533	13.832
0.623	8.195	0.653	5.109	0.606	7.148	0.620	11.069
0.781	6.378	0.810	2.421	0.755	4.273	0.806	3.279
1.000	3.333	1.000	-1.480	1.000	-1.547	1.000	-6.122