**­­­­Development and Testing of Molecular Mechanics Force Fields for Biomedical Research**

Molecular simulations have played essential roles in biochemical and biophysical sciences. A chief application of molecular simulations is to elucidate the molecular interactions between small molecule inhibitors and protein or nucleic acid targets, and then enhance or eradicate the functions of their targets via rationally design of high-potency agonists or antagonists. **However, nowadays it is still very challenging to accurately predict the free energies of such interactions and processes.** A key to the success of molecular simulation studies and structure-based rational drug designs is the quality of molecular mechanics force fields (**MMFFs**), and a major impediment to developing more accurate MMFF is to test them objectively in experiments. **The major goal of this project is to further develop the existing and develop a new generation of the general AMBER force fields (GAFF) for studying biomolecule-ligand interactions.** The new general-purpose MMFF (referred to here and thereafter as **GAFF3**) is based on a new charge model which is efficient, largely conformation-independent, and has high accuracy on solvation free energy calculations. In a long run, **a spectrum of high-quality GAFF3 compatible MMFFs will be developed for all types of biological macromolecules for biomedical research.** The new set of MMFFS will be highly transferable and self-consistent for studying protein-ligand and nucleic acid-ligand interactions, and will enable more accurate calculations of the protein-ligand and nucleic acid-ligand binding free energies and facilitate computer-aided drug discovery. The developed GAFF force fields will be rigorously scrutinized and critically assessed through direct comparisons with experiments in an extensive set of model systems. The large-scale FF assessment will be accelerated by using GPUs in both molecular dynamics simulations and alchemical free energy calculations.

**A. BACKGROUND AND SIGNIFICANCE**

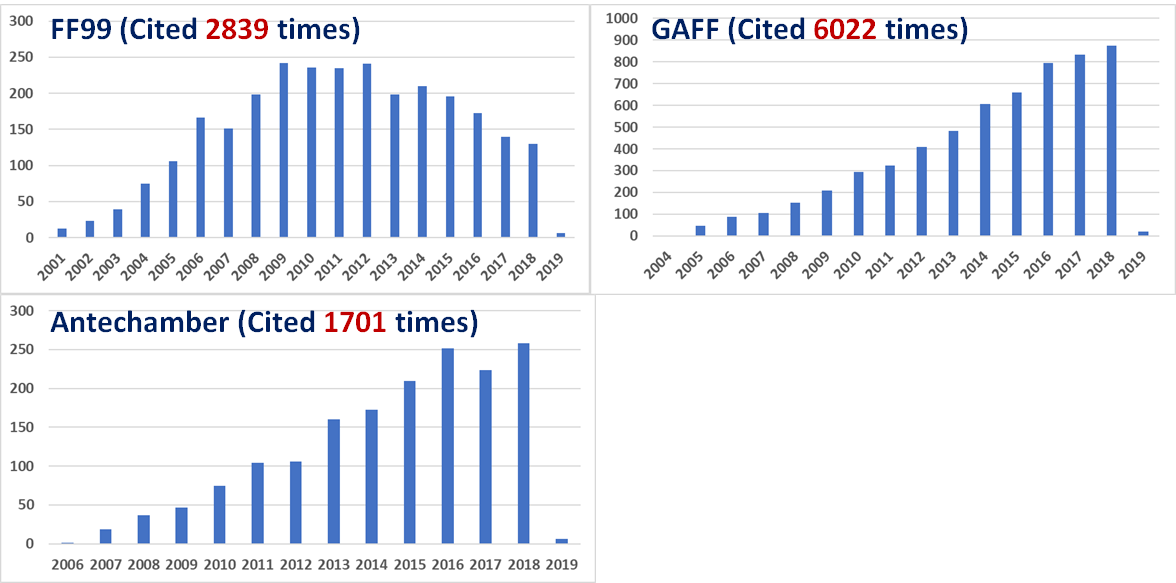
**A.1. Molecular Simulations and Force fields.** Atomistic simulations have become an important tool for studying the structure, dynamics, interactions, and functions of biomolecular systems.[1-8](#_ENREF_1) Critical to their success is the accurate representations of potential energy (referred to as the “force field”). Advancements in computer technology and simulation methodology have enabled molecular dynamics (**MD**) simulations of increasingly complex biomolecular systems over longer times and with larger molecules. With these advances, the need for accurate and efficient force fields that can represent diverse biomolecular systems has become much more critical.

A force field (**FF**) is composed of an energy function and its associated parameters. Because of its simplicity and robustness, the effective functional form (**Eq. 1**) has been extensively used in many biomolecular FFs, which include AMBER FF94,[9](#_ENREF_9) FF99,[10](#_ENREF_10) FF99SB,[11](#_ENREF_11) FF12SB,[12](#_ENREF_12) FF14SB,[13](#_ENREF_13) CHARMM,[3](#_ENREF_3) NAMD,[14](#_ENREF_14) OPLS,[8](#_ENREF_8), [15](#_ENREF_15) and GROMACS,[16](#_ENREF_16) etc.

Where *r*, *θ*, and *Φ* are the instant values of bond lengths, angles, and dihedral angles, *req* and *θeq* are the equilibrium values of bond lengths and angles, *n* and *γ* are the dihedral multiplicities and phases, *Kr*, *Kθ* and *Vn* are the respective force constants, *Rij* is the distance between atom *i* and atom *j*, *qi* and *qj* are their respective atomic charges, and *Aij* and *Bij* are van der Waals parameters.

**A.2. AMBER Force Fields.** “AMBER force fields” have been the primary force fields in the AMBER simulation packages and adapted to run directly in CHARMM and NAMD. Since the first description by Weiner and Kollman in 1981,[17](#_ENREF_17) AMBER-associated force fields have evolved from united-atom[18](#_ENREF_18) to all-atom models,[9-10](#_ENREF_9), [17](#_ENREF_17), [19-20](#_ENREF_19) from additive to polarizable,[21-25](#_ENREF_21) and from special (such as proteins[9](#_ENREF_9) and nucleic acids,[10](#_ENREF_10) lipids[26](#_ENREF_26) and carbohydrates[27](#_ENREF_27)) to general organic molecules.[28](#_ENREF_28) AMBER force fields, which are for simulations of biological systems including proteins, DNAs & RNAs, etc., have been widely used in modeling biomolecules with more than 20,000 citations as of this writing.

**A.3. Qualification of the Research Team.** The principal investigator (**PI**), Dr. Junmei Wang, **is the inventor of the widely-used general AMBER force field (GAFF)[28](#_ENREF_28" \o "Wang, 2004 #3)** for general organic drug-like molecules, and the main developer of the corresponding tools for FF parameter assignment including Antechamber.[29](#_ENREF_29" \o "Wang, 2006 #1996) He is also one of the main contributors of the popular AMBER force field family (especially AMBER FF99[10](#_ENREF_10)). AMBER FF99, GAFF and Antechamber have been popular in numerous research studies with more than 10,500 citations in total (**Fig. 1**). Collaborator Dr. Tai-Sung Lee has a broad and strong background in free energy methodology, molecular simulations, and software development.[19](#_ENREF_19), [30-32](#_ENREF_30) Recently he developed and implemented pmemdGTI, a GPU-accelerated thermodynamic integration module into AMBER software package.[30](#_ENREF_30)[L5] Collaborator Dr. Xibing He is both an AMBER and CHARMM developer and he has extensive expertise and experience in force field development.[33-38](#_ENREF_33) In short, we have assembled a highly competitive team and has sufficient expertise and experience to pursue the proposed research.



**Figure 1. Annual citations of FF99, GAFF and Antechamber.** (webofknowledge.com)

**TAISUNG: I would move section D “ADDITIONAL REMARKS on FORCE FIELD DEVELOPMENT” here to justify the need of point-charge FF development**

**A.4. Development of a New, Accurate, Highly Transferable, and Self-Consistent Force fields for Studying Protein-Ligand and Nucleic Acid-Ligand Interactions.** It is a challenging and ambitious goal to develop such kinds of force fields given the extremely difficult nature of MMFF development. MMFF development has long been more of an “art” than a “science”.[39-41](#_ENREF_39) Atomic charge model, the cornerstone of a MMFF, are developed both empirically and physically (such as the RESP[42-43](#_ENREF_42) charge of AMBER FFs). Ideally, a charge model for additive force fields would possess all the following features: efficient, conformation-independent, highly accurate in solvation free energy calculations, and can be applied to both small organic and macromolecules. Unfortunately, no charge method in the current FFs has all of these features. In this proposal, we propose to develop a new, efficient, largely conformation-independent and high-quality atomic charge model for our new AMBER force field development.

Accurate solvation free energy calculation[44-55](#_ENREF_44) plays an essential role for us to achieve the above research goal. Solvation free energy (**ΔGsolv**) calculation is essentially similar to that of receptor-ligand binding free energy (**ΔGbind**) calculation, as both deal with the free energy of transferring a molecule from one environment to another environment (from gas phase to solvent for ΔGsolv, and from solvent (usually water) to the binding site of the receptor for ΔGbind). The simplest, most ubiquitous and most important solvent is water. The calculation of ΔGsolv in water (hydration free energy, **ΔGhyd**) is still much more computationally demanding than that of other physicochemical properties such as density and heat of vaporization. Therefore, usually ΔGsolv (including ΔGhyd) is not used as a main target during FF parameterization, but just as an auxiliary validation test. In such situation, the standard of justifying ΔGhyd is also not as strict as that of justifying density and heat of vaporization. Usually an unsigned error of 1.0 kcal/mol is considered as good, and 2.0 kcal/mol is still acceptable. The general difference between the calculated ΔGhyd with current popular force fields is > 1.2 kcal/mol or even larger,[54](#_ENREF_54) limiting the accuracy of predicting ΔGbind even with the state-of-the-art alchemical free energy calculation methods, such as thermodynamic integration (**TI**)[56](#_ENREF_56) and free energy perturbation (FEP),[57](#_ENREF_57) which are rigorous but very expensive. Loose criteria and poor quality of ΔGsolv prediction will ultimately impair the accurate prediction of ΔGbind no matter how much efforts are put on sampling protein and ligand conformations.

With the growing computer power, facilitated by both inexpensive GPU-based computer clusters[58-59](#_ENREF_58) and access to the local and national supercomputing resources, particularly the GPU-TI[30](#_ENREF_30) code recently released in AMBER 18,[31](#_ENREF_31) it is timely for us to introduce TI reproduction of experimental solvation free energies into our FF parameterization scheme as a primary target. We set off to develop a new and high-quality atomic charge model, and based on that a spectrum of coherent FFs will be developed for modeling both general organic and biological molecules.

**A.5. Critical Assessment of GAFFs in Modeling Protein-Ligand and Nucleic Acid-Ligand Interactions.** Assessing a newly developed force field in realistic biological applications is a challenging task. Improving the accuracy of computational predictions is of critical importance for the drug design process. Binding constants depend exponentially on the binding free energy, thus relatively small errors (0.6 kcal/mol) can have a significant impact on the efficacy of a potential drug. The main factors that determine the accuracy of a free energy simulation are the extent to which important configurational degrees of freedom are sampled,[60-61](#_ENREF_60) and the accuracy and reliability of the underlying force field (FF) that models the molecular interactions. The force fields include large sets of parameters that were derived from quantum-chemical calculations and selected experimental data such as density, heat of vaporization of neat liquids, or hydration free energies. However, to the best of our knowledge, no currently available force field has been tuned to consider quantitative binding free energy data in its parametrization scheme, due to the associated computational burden.[62-63](#_ENREF_62) Given that binding free energies represent one of the most important applications in computational chemistry, it is worthwhile to develop new methods that would enable it to enter into the force field parameterization/refinement process.

Very recently, Dr. Lee et al. has developed a set of tools based on GPU-accelerated free energy methods, i.e., the 𝞴-SAMS approaches, which speedup the convergence of alchemical transformations by roughly three to five folds. Given that the speedup of the AMBE GPU-TI code over the CPU code is over two orders of magnitudes, now it is possible to assess the performance of new force-field in free energy calculations on GPU roughly three orders of magnitudes faster than on CPU. We will utilize the new GPU-accelerated free energy sampling (𝞴-SAMS) to critically evaluate the proposed force fields: 1. Establishing extremely fast free energy simulation pipeline infrastructure; 2. Performing simulations on a current golden-standard of protein-ligand free energy simulation data set; 3. Producing meaningful error bars through multiple independent simulations.

Besides the pathway method, a set of endpoint free energy methods, molecular mechanics Poisson-Boltzmann and Generalized Born Surface Area (MM-PB/GBSA),[32](#_ENREF_32), [64-66](#_ENREF_64) which achieve a good balance between computational efficiency and accuracy, have been widely used in structure-based drug designs. The performance of MM/PBSA and MM/GBSA depends on the quality of both the MMFFs and the implicit solvation models. We have conducted critical evaluations of both methods.[67-70](#_ENREF_67) To improve the performance of the two endpoint methods, a key is to develop force field-consistent solvation models. For the newly developed charge model for GAFF3, we plan to optimize the PBSA and GBSA radius parameters to reproduce measured solvation free energies. We will conduct large scale evaluation for a diverse drug targets that have multiple inhibitors through collaboration.

Moreover, our force field models have been and will continue to be assessed in numerous drug discovery projects where our predictions can be confirmed by experimental biologists. For most of the projects, we intend to develop novel compounds to activate (as agonists) or inhibit (as antagonists) protein targets. We have established a broad collaboration network with experimental scientists. Most collaborations resulted in grant application and publications.[71-76](#_ENREF_71)

**B. INNOVATION**

The novelty of this R01 project is highlighted from the following five aspects:

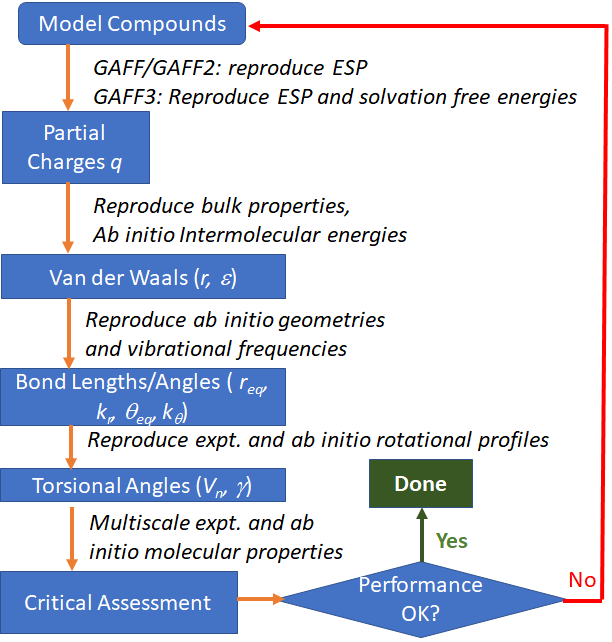
* In contrast to the traditional “bottom-up” strategy in MMFF development, the GAFF2/GAFF3 development is mainly guided by a novel top-down strategy to maximize the transferability and self-consistency.
* A novel charge method will be developed to largely overcome the limitations of the charge models in current mainstream FFs. This charge model is efficient, much less conformation-dependent, and can be applied to arbitrary small molecules and biomolecules. MMFFs based on this charge method will be coherent, self-consistent and have much better transferability. Those MMFFs will be able to much better model the protein-ligand and nucleic acid-ligand interactions, and as such, they can boost the successful rate of structure-based rational drug design.
* Solvation free energy will be applied as the **primary target** for the development of a charge model. As aforementioned, solvation free energy (including hydration free energy) has never been used as the primary target for force field development. This is partly due to the much higher computational cost of calculating solvation free energies comparing to that of calculating other physicochemical properties. This limitation can be alleviated due to the increasing computing power and computer resources nowadays. Furthermore, the high cost at the beginning stages like force field development will be rewarded with more accurate results at the final goal stages of applications on the biological systems, such as the calculation of protein-ligand binding free energies for the purpose of drug design.
* Advances on TI calculations and large-scale FF evaluations accelerated by GPU-TIs. The GPU-accelerated 𝞴-SAMS approach is novel in that it uses a statistically optimal estimation of the free energy difference between alchemical states through Rao-Blackwellization of the joint distribution of conformational and alchemical spaces. Here, we leverage the 𝞴-SAMS approach to run the Gibbs sampler along a fine-grid in 𝞴 space to produce a statistically optimal distribution and collect dU/d𝞴 statistics for thermodynamic integration.
* Applications of the newly developed 𝞴-SAMS approaches represent a major innovative advance in the field that would reach unprecedent milestones for various biological simulations and are perfectly suitable for force field development where large scale of free energy simulations are needed.

**C. APPROACH**

**C.1 General Force Field Development Philosophy and Parameterization Strategy.**

The research goal of this application is to develop and evaluate a set of high-quality force fields that are highly transferable and self-consistent for studying protein-ligand and nucleic acid-ligand interactions. What are the traits of a good force field? We believe **transferability, self-consistency, accuracy, and applicability are the most important attributes of a good force field**. Even though most biomolecular force fields were developed using the effective function form (**Eq. 1**), the parameterization strategies are quite different. AMBER force fields distinguished themselves from others by using physical charges derived to fit the *ab initio* electrostatic potentials. This special feature, which makes AMBER more expandable as proven by the success of GAFF, will be kept in our newly developed FFs. Traditionally, most FFs were developed following a bottom-up strategy: FF parameterizations were performed one compound class after another, with an assumption that the developed FF parameters could be seamlessly transferred to new compound classes. However, this assumption may not always hold. To maximize the transferability and self-consistency, a novel top-down strategy was applied to develop the GAFF2/GAFF3 force fields: (1) the FF parameters were optimized for multiple compound classes simultaneously; and (2) new atom types and new FF parameters were introduced only when they could significantly improve the fitting performance. Both the bottom-up and top-down strategies will be applied in our force field development in the future.

**Figure 2. Strategies of force field development.** The FF parameterization of non-bonded terms, which is the focus of this R01 grant, are shown in dashed-rectangle.



The parameterization for an FF is an iterative procedure as shown in **Fig. 2**. The targets of FF parameterization include the quantum mechanics (**QM**) calculations of geometries, dipole moments, vibrational frequencies, conformational and intermolecular energies, etc., and the experimental data of physicochemical properties, such as crystal geometries, crystal volumes, liquid densities, heats of vaporization, etc. The atomic partial charge assignment is usually the first step of FF parameterization, because the atomic charges are the main contributor to the intra- and inter-molecular energies, and other bulk properties including the solvation free energies.

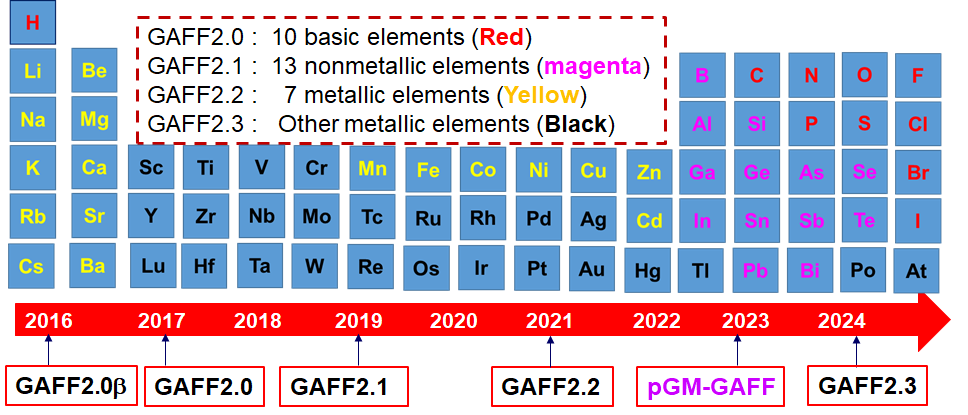
**C.2 Aim 1. Continue Development of the Second Generation of GAFF Force Field (GAFF2)**

**Rational.** It is necessary to develop a general force field that is compatible to the mainstream AMBER biomolecular force fields in biomedical research for studying biomolecule-ligand interactions. As most AMBER biomolecular FFs apply RESP charges,[42-43](#_ENREF_42) GAFF2 has been and will continue be developed using the RESP charge method.

**C2.1 Current GAFF2 Development.**After its release, we received numerous suggestions and comments on GAFF, mainly coming from the AMBER mailing list. We have maintained GAFF and Antechamber for years and released several subversions (1.5, 1.6, 1.7 and 1.8) where recognized problems are remedied. Evidently, GAFF could only become a successful force field with continuous input from its users. Recently, we realized that the GAFF van der Waals parameters, which were adopted from the AMBER FF94/99 force fields, needed to be optimized to bring the performance of GAFF to a new higher level. As such, we decided to move forward to develop the second generation of GAFF – GAFF2. A beta version of GAFF2 – GAFF2β, which covers the chemical space of the common organic molecules, has been released with AMBER14 and 15. Later GAFF2.0 has been released with AMBER 16, 17 and 18. The major improvement achieved by GAFF2 includes: (1) reparameterization of the van der Waals parameters to better reproduce the high-level *ab initio* and high quality experimental data of a variety of molecular properties including intermolecular energy, liquid density, heat of vaporization, and hydration free energy (the RMS errors reduced 20 to 40% for the condensed molecular properties); (2) development of empirical functions and a set of associated parameters to directly calculate the bond stretching and bond bending force constants (Compared to GAFF, the RMS errors of vibration frequencies reduced ~50% in GAFF2); and (3) reparameterization of torsional angle parameters using a much larger set of model compounds for which the rotational profiles were calculated at the MP2/aug-cc-pVTZ level. GAFF2 is expected to be an even more successful general-purpose force field and GAFF2-based docking scoring functions could significantly improve the success rates of virtual screenings in target-based drug design.

**C2.2 Further Development of GAFF2.** We propose to develop force field parameters for broader coverage of the chemical space of small molecules, going beyond the ten basic elements covered by GAFF and GAFF2.0. The road map of GAFF2 development is shown in **Fig. 3**. Eventually, GAFF2 will become a universal force field covering most elements in the periodic table except lanthanoids and actinoids. All the developed FFs will be released with AMBER. The chemical space coverages are summarized in the dashed rectangle of **Fig. 3**, and the tentative release dates of GAFF2 force fields are shown at the bottom of **Fig. 3**. The significant expansion of the chemical space coverage enables GAFF2 to study very complicated biomolecular systems consisting of different types of entities. It could also find applications in other research fields, such as material sciences.

**Figure 3. Roadmap of GAFF development**.



**GAFF2.1.** Frequently, our users raise questions on the missing force field parameters involved by the uncovered nonmetallic elements, such as boron, silicon, selenium, etc. FF parameterization for 13 nonmetallic elements in GAFF2.1 is ongoing. The same FF parameterization strategy of developing GAFF2.0 is utilized, as the liquid property data, such as density and heat of vaporization, are also available for those nonmetallic elements. We plan to release GAFF2.1 at the end of this year according to the roadmap.

**GAFF2.2/2.3**. FF development for metallic compounds is different from that for organic molecules. First, the functional forms are different: (1) a cosine Fourier expansion should be used for metallic bond angles as the harmonic functional form cannot address the multiple equilibrium bond angle problem;[77-78](#_ENREF_77) (2) the Morse function may describe the bond stretching better for metallic bonds;[77](#_ENREF_77) and (3) the 12-6 Lennard-Jones potential describing the van der Waals term may be replaced with the other potential forms.[79-81](#_ENREF_79) Second, metallic FF parameterization heavily depends on high-quality *ab initio* calculations. We will collaborate closely with Dr. Kenneth Merz at Michigan State University on the general metallic force field development.

**GAFF2.** Besides to expand the chemical space covered by GAFF2, we also plan to significantly improve the quality of GAFF2 by reparametrizing the existing and introducing new parameters. We will focus on the problematic molecules reported by our users and/or occurring to other general-purpose force fields.[82-83](#_ENREF_82) We plan to systematically evaluate GAFF2 in reproducing the *ab initio* rotational profiles for a great number of model compounds. In our effort of conducting drug likeness analysis, a brutal force algorithm was applied to iteratively dissect drug molecules into fragments. We will collect all the fragments that have one rotatable bond to generate model compounds. We will then generate rotational profile at the MP2/aug-cc-pVTZ//MP2/6-31G\* level. Next, we will examine how well rotational profiles by GAFF2 reproduce the *ab initio* ones. New torsional angle parameters will be introduced if it is necessary.

The basic FF parameterization strategy is summarized in **Fig. 2**. We plan to perform FF parameterizations using a set of hierarchical data, which include high-level *ab initio* data (esp. interaction energy), and experimental gas and bulk properties including the thermodynamic data.[84](#_ENREF_84) This multiscale optimization, enabled by the rapidly increasing computer power, can facilitate the development of a highly transferrable and general purpose force field.[85](#_ENREF_85)

**C.3 Aim 2. Development of a New, Accurate, Highly Transferable, and Self-Consistent Force Field (GAFF3) for Studying Protein-Ligand and Nucleic Acid-Ligand Interactions**

**Rational.** Considering the essential relationship between the solvation free energy ΔGsolv (including hydration free energy ΔGhyd) and the receptor-ligand binding free energy ΔGbind, the general accuracy of ΔGsolv calculation determines the best results that can be expected for ΔGbind calculation. Hence, a new set of force fields (**FFs**) with high accuracy of ΔGsolv prediction will enable us to improve the accuracy of ΔGbind prediction, and the latter is extremely important for computer-aided drug design.

**C3.1 Background.** Even though all FF parameters in an MMFF (**Eq. 1**) are developed in an iterative manner, the model of assigning atomic partial charges is always the first step due to its major role in determining the calculated intra- and inter-molecular energies. Traditionally, the AMBER force fields and GAFF assign atomic charges utilizing the RESP charge model, which fits the electrostatic potential (ESP) of a molecule calculated by QM at the HF/6-31G\* level of theory.[42-43](#_ENREF_42) While RESP charge model produces suitable charges for condensed simulations, there are inherent problems, including but not limited to the high cost of QM calculations for every new molecule encountered in real projects. As an alternative solution, GAFF also adopts a faster and more efficient method named AM1-BCC[86-87](#_ENREF_86) to emulate the HF/6-31G\* ESP. The basic idea of AM1-BCC is as follows: the AM1 Mulliken charges, although incapable of satisfactorily reproducing the QM ESP, can capture underlying electronic features such as formal charge and electron delocalization; simple additive bond charge corrections (**BCCs**) are then applied based on the general atom and bond types in the molecule so that the resulting charges can emulate the RESP charges more closely than AM1.

While the AM1-BCC method has been widely and successfully applied along with AMBER force fields and GAFF, we’ve found that the calculated ΔGsolv for various organic solutes have a mean unsigned error (**MUE**) > 1 kcal/mol. Such a limitation is actually inherited from RESP charges, which cannot accurately reproduce ΔGsolv for some functional groups of organic molecules (**Table 1 and Fig. 4**).

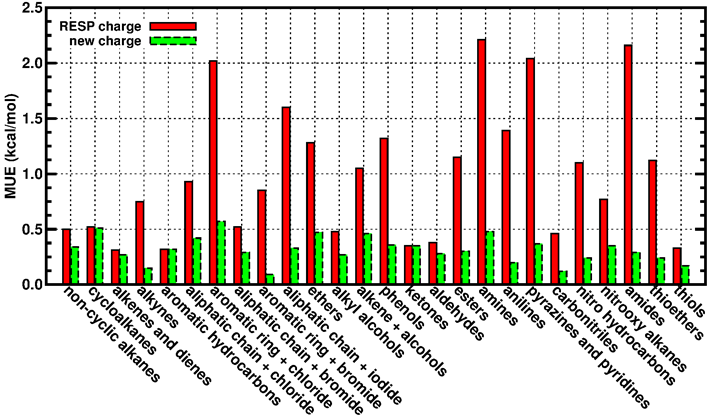
Hence, we propose to re-adjust the BCC values for various atom types and bond types of organic molecules to directly target accurate reproduction of experimental data, which will lead to more accurate ΔGsolv and ΔGbind based on aforementioned discussion. The van der Waals and bonded term parameters will be validated and if necessary, re-adjusted in accordance with newly developed BCC parameters (and corresponding new atomic charges).

**C3.2 - Preliminary Results.** We have calculated the solvation free energies of series of organic molecules in the solvent water (ΔGhyd), using the RESP charges, current (old) AM1-BCC charges and our newly developed AM1-BCC charges. The explicit water molecules are treated with the TIP3P water model[88](#_ENREF_88) which is bounded to AMBER force fields and GAFF. The calculation method is the TI as described in previous literature.[30](#_ENREF_30) As shown in **Table 1**, a lot of calculations using the RESP charges and old AM1-BCC charges have a mean signed error (**MSE**) with the absolute value very close or equal to MUE, which demonstrates a systematic bias, i.e., the calculated ΔGhyd results are systematically more positive or more negative than the experimental data. For example, the calculated ΔGhyd of 17 phenols are systematically more positive than experiment by 1.32 kcal/mol from RESP charges, and systematically more positive by 0.95 kcal/mol from old AM1-BCC charges. With our newly adjusted AM1-BCC model, the systematic error is greatly alleviated, and the MSE is only -0.16 kcal/mol, and MUE is reduced to only 0.36 kcal/mol. Among the 26 types of molecules in **Table 1**, 17 types show systematic errors for calculated ΔGhyd using RESP charges, and 11 of them have a systematic error greater than 1.0 kcal/mol in magnitude. With our new AM1-BCC model, all MSEs are close to 0 kcal/mol and most MUEs are less than 0.4 kcal/mol (**Table 1 and Fig. 4**). In a short summary, the accuracy of calculated ΔGhyd can be significantly improved by adjusting the BCC parameters.

**C3.3 Experimental Design.** The similar FF development philosophy and FF parameterization strategy for developing GAFF2 will be seamlessly introduced to GAFF3 force field development. GAFF3 will first be a general-purpose FF covering the 10 basic elements as GAFF2.0. In a long run, GAFF3 may be expanded to cover other elements given high-quality experimental solvation free energies are available for deriving BCC parameters.

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| --- | --- | --- | --- | --- | --- | --- | --- |
| **Table 1.** **The mean signed errors (MSE) and mean unsigned errors (MUE) of calculated hydration free energies** (in kcal/mol) of various molecules using atomic charges from RESP, old AM1-BCC and new AM1-BCC methods. ‘\*’: related BCC values are not changed; ‘/’: MSE & MUE are not calculated yet for this type of molecules. | | | | | | | |
| **Molecule Types** | **Mol #** | **RESP charges** | | **Old AM1-BCC charges** | | **New AM1-BCC charges** | |
| **MSE** | **MUE** | **MSE** | **MUE** | **MSE** | **MUE** |
| Non-cyclic alkanes \* | 27 | -0.45 | 0.50 | -0.27 | 0.34 | -0.27 | 0.34 |
| Cycloalkanes \* | 9 | -0.36 | 0.52 | -0.29 | 0.51 | -0.29 | 0.51 |
| Alkenes & dienes | 22 | 0.11 | 0.31 | 0.69 | 0.69 | 0.11 | 0.27 |
| Alkynes | 6 | -0.75 | 0.75 | 0.59 | 0.59 | 0.01 | 0.15 |
| Aromatic hydrocarbons | 38 | -0.08 | 0.32 | -0.80 | 0.81 | -0.18 | 0.32 |
| Aliphatic chain + chloride | 31 | 0.84 | 0.93 | / | / | 0.22 | 0.42 |
| Aromatic ring + chloride | 23 | 2.02 | 2.02 | / | / | 0.22 | 0.57 |
| Aliphatic chain + bromide | 13 | 0.35 | 0.52 | / | / | 0.01 | 0.29 |
| Aromatic ring + bromide | 3 | -0.56 | 0.85 | / | / | -0.09 | 0.09 |
| Aliphatic chain + iodide | 9 | 1.60 | 1.60 | / | / | -0.01 | 0.33 |
| Ethers | 26 | 0.91 | 1.28 | 0.64 | 0.93 | 0.12 | 0.47 |
| Alkyl alcohols | 27 | 0.36 | 0.48 | 1.67 | 1.67 | 0.01 | 0.27 |
| Alkene + alcohols | 4 | 1.05 | 1.05 | / | / | 0.09 | 0.46 |
| Phenols | 17 | 1.32 | 1.32 | 0.95 | 0.95 | -0.16 | 0.36 |
| Ketones | 18 | -0.06 | 0.35 | / | / | -0.02 | 0.35 |
| Aldehydes | 13 | 0.13 | 0.38 | / | / | -0.10 | 0.28 |
| Esters | 37 | -1.11 | 1.15 | / | / | -0.14 | 0.30 |
| Amines | 25 | 2.21 | 2.21 | 1.52 | 1.52 | 0.16 | 0.48 |
| Anilines | 8 | 1.39 | 1.39 | 0.34 | 0.47 | -0.08 | 0.20 |
| Pyrazines & pyridines | 17 | 2.07 | 2.04 | / | / | 0.33 | 0.37 |
| Carbonitriles | 5 | -0.43 | 0.46 | 1.44 | 1.44 | 0.01 | 0.12 |
| Nitro hydrocarbons | 9 | -1.10 | 1.10 | 2.02 | 2.02 | -0.11 | 0.24 |
| Nitrooxy alkanes | 9 | -0.77 | 0.77 | 1.94 | 1.94 | -0.09 | 0.35 |
| Amides | 8 | 2.16 | 2.16 | / | / | 0.25 | 0.29 |
| Thioethers | 7 | 1.01 | 1.12 | 1.22 | 1.22 | -0.05 | 0.24 |
| Thiols | 4 | 0.14 | 0.33 | 0.45 | 0.45 | -0.15 | 0.17 |

**New Charge Model Development for GAFF3.** We plan to adjust the BCC parameters of the new charge model against ΔGsolv and partition coefficients in various solvents. We are going to calculate the ΔGsolv of various molecules in organic solvents, such as octanol, cyclohexane, benzene, etc.[89-90](#_ENREF_89) These solvents are important medium for studying chemical and biological processes, and mimicking the hydrophobic or amphiphilic environments inside of or on the surfaces of biological systems, such as membranes, proteins, DNAs and RNAs. The difference of ΔGsolv of a molecule in two immiscible solvents can be used to calculate an important property – partition coefficient, which is a measure of a solute’s hydrophobicity and a proxy for its membrane permeability in drug design. Especially, partition coefficients are relatively easy to measure experimentally compared to ΔGsolv, and more experimental data of partition coefficients are available. [89-90](#_ENREF_89) The newly developed charge model will be tested and adjusted against ΔGsolv in or partition coefficients between various solvents.



**Figure 4. The mean unsigned errors (MUE) of calculated hydration free energies** for various types of organic solutes using atomic charges from the RESP model and our new AM1-BCC model.

**van der Waals Parameterization for GAFF3.** In a MMFF,van der Waals (**VDW**) interactions are strongly coupled to the employed charge method. We plan to validate, and adjust if necessary, the VDW parameters in the current GAFF. The target data include the interaction energies of dimers and the experimentally measured densities of liquids and heats of vaporization, and solvation free energies. We have finished calculating interaction energies at the MP2/aug-cc-pVTZ//MP2/6-311++G(d,p) level with counterpose corrections for 2954 dimers. We have collected a set of 241 diverse molecules that have measured densities and 308 molecules that have measured heats of vaporization. An efficient VDW parameterization scheme developed by us and has been successfully applied in the development of polarizable GAFF[23](#_ENREF_23) will be utilized to tune VDW parameters to reproduce both the *ab initio* interaction energies and the liquid properties.

**Bonded Term Parameterization for GAFF3.** After finishing the development of the new charge model and the VDW parameters, we will focus on the development of the bonded term parameters. We are going to test, and adjust if necessary, the current parameters of bonds, angle, and dihedrals in GAFF2, to make sure they are consistent with the new charge model and the new VDW parameter set. The target data include experimentally measured (such as crystal data) or QM calculated geometries, dipole moments, conformational energies and rotational profiles of model compounds.

**C.4 Aim 3. Critical Evaluation of GAFF2/GAFF3 in Studying Biomolecule-Ligand Interactions**

**Rational.** Force field validation is an on-going process. We choose different model systems to calibrate different properties for different force fields. We utilize protein-ligand and nucleic acid-ligand systems to evaluate both GAFF2/GAFF3 and the biomolecular force fields. The force field models will also be extensively assessed with real drug discovery projects. We plan to integrate the force field development and validation into drug discovery projects through collaborations.

One common problem and task in real drug design projects is to pre-select which compounds to synthesize and further test with experimental assays which are expensive in terms of both time and cost. Physics-based alchemical free energy calculation methods, such as thermodynamic integration (**TI**) and free energy perturbation (FEP), are expected to be very useful in the pre-selecting (screening, lead identification and lead optimization) stages. However, several factors bring hurdles to the routine usages of free energy calculation methods in real drug design projects: (1) availability and accuracy of FF parameters of query compounds; (2) demanding computing resource; (3) difficulty of automatic system setup and posterior analysis. We are working on Aim 1 (continue development of GAFF2) and Aim 2 (development of GAFF3) in this proposal to solve the 1st problem, developing and improving GPU accelerated TI algorithms to alleviate the 2nd problem (**C.3.1**), and we also plan to develop a user-friendly GUI toolkit to overcome the 3rd problem (**C.3.2**) to help non-expert users to set up the simulations and do analysis for various free energy calculations.



**Figure 5. Preliminary results of *λ*-SAMS approach.** The standard errors averaged 𝞓G from 16 independent simulations are shown for the thrombin ligand 1d->7a mutation in water. “No RepEx”: TI simulations without replica exchange; “RepEx at 100 fs”: TI simulations with the replica exchange (between *λ*-windows) with 100 fs trial frequency; “SAMS 100”: *λ* -SAMS with 100 *λ* -windows; “SAMS 200”: *λ* -SAMS with 200 *λ* -windows. TI calculations were performed with 12 *λ* -windows.

**C.3.1 New Sampling Algorithm for Thermodynamic Integration.** We propose to apply a recently developed innovative sampling algorithms to boost the convergence of TI calculations. The proposed *λ-*SAMS approach was applied to the thrombin ligand 1d->7a mutation [L1] in water as proof of principle. Shown in **Fig. 5**, the *λ-*SAMS, both with 100 *λ*-windows or 200 *λ-*windows reach the standard error of 0.01 kcal/mol in about 5 ns total simulation time, compared to 20 ns by the regular TI simulations and Hamiltonian (*λ-*window) replica exchange/TI simulations.

**Experimental Design.** The detailed plan on sampling algorithm development and evaluation is described as follows.

* Testing infrastructure: we will first establish an infrastructure that enables tuning of force field parameters for enhanced predictive capability using fast GPU-accelerated free energy methods based on the *λ-*SAMS algorithm. As the first step, the evaluation of free energy transformations will be performed using the current AMBER/GAFF, the AMBER/GAFF2 and AMBER/GAFF3 force fields. Results will be compared with benchmark results for all ligand force fields.
* Baseline: The second step will involve applying the framework and validated procedures to obtain the very first benchmark quality data and predictions (including the first meaningful statistical error estimates) of ligand binding potency against the current “gold standard” drug discovery dataset [L2]. This dataset encompasses a broad range of target classes and a diverse set of 200 ligands. Toward this end, we will use the most commonly used publicly available force fields for ligand-protein binding in explicit solvent, including CHARMM/CGenFF, AMBER/GAFF and GAFF2, and OPLS-AA. Not only will the results of this work provide a valuable assessment of the quality of current publicly available force fields for ligand-protein binding, but it will create an extremely powerful dataset from which next-generation versions of these force fields for prospective drug discovery can evolve efficiently.
* Simulation Protocols: Individual alchemical transformations for compound libraries that have a common protein target will be determined using a “thermodynamic map”. Following the same thermodynamic mapping procedure as Schrödinger Inc. which was obtained by the LOMAP algorithm [L3,L4] (which uses the maximum common substructure) and has been reported in the literature for the entire dataset [L2].

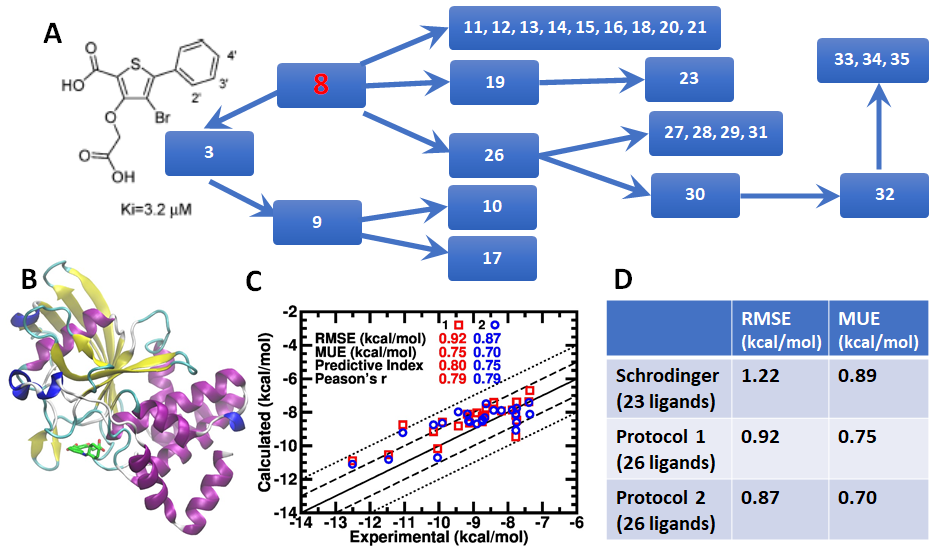
The GPU-accelerated 𝞴-SAMS approaches will be utilized to obtain statistical optimal estimation of relative free energies. Multiple independent sets (16 to 32) of simulations will be performed to establish meaningful error bars.

* Assessment of the to-be-developed GAFF3 force-field: We will perform binding free energy simulations on the data set mentioned above using GAFF3. Since the baseline results developed have benchmark quality and have rigorous statistical error estimates based on multiple independent sets of simulations, we will be able to assess the quality of -GAFF3. Trajectory snapshots from alchemical free energy simulations for these transformations will be saved at 1 ps intervals creating an incredibly rich dataset that can be efficiently post-processed to obtain both TI and FEP/MBAR metadata for further free energy analysis.

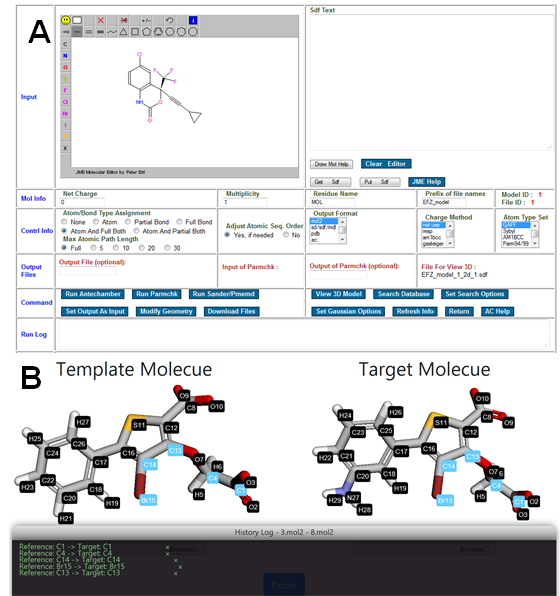
**C.3.2 Web-Based Toolkit for TI Preparation and Analysis.** According to our experience of talking to both academic researchers and industrial R&D personnel, such an automatic or semi-automatic GUI toolkit is in great demand.

**TI Setup** We have been developing a series of in-house scripts and programs for setting up TI calculations in AMBER and we have explored protocols of selecting TI 𝛌 windows, mutation paths, integration methods, and GAFF2/GAFF3 parameters to get accurate results. We finished testing a set of systems with the Protein Tyrosine Phosphatase 1B (PTP1B) as the receptor (**Fig. 6B**) and a series of 26 compounds (**Fig. 6A**) as the ligands[91](#_ENREF_91). The experimental values of *Ki* measurements range over 4 orders of magnitude (from several µM to less than 1 nM). We achieved lower root mean square error (RMSE) and MUE than the reported results from the commercial FEP module in Schrodinger Suite[57](#_ENREF_57) (**Fig. 6C** and **6D**). Next, we need to integrate the in-house scripts and programs into the GUI toolkit.

**Figure 6**. **TI setup and results for PTP1B receptor with a series of 26 ligands.** (A) mutation paths of 26 ligands. (B) 3D structure of PTP1B with a ligand. (C) Calculated binding free energies VS experimental data. (D) RMSEs and MUEs from our calculations compared to Schrodinger FEP.



**A GUI-Toolkit for TI Setup and Analysis**. TI calculations are often be used to calculate the relative binding free energy, where an atom or a group of atoms in ligand A is/are mutated to another atom or another group of atoms in ligand B. The designed GUI toolkit will have the following functions: (1) Open a graphic window and load the reference ligand A with 3D coordinates (sdf, mol, mol2, pdb, restart files from various packages). (2) Load the target ligand B in the format of SMILES/SMARTS string or 2D/3D coordinates, automatically generate a 3D structure if the input is not 3D, and show it on a second graphic window. (3) Automatically calculate the maximum common substructure (MCS) between ligands A and B, and map the MCS atoms. The algorithms behind the manipulations are based on the subroutines in the Antechamber program which can precisely describe an atom’s chemical environment (including the atomic paths, atomic connectivity and atomic property) and automatically define complicated force field atom types, bond types and discriminate subtle local chemical environments. Sometimes multiple mapping is possible, such the 2,6 positions or 3,5 positions on a benzene ring; in such cases, users are allowed to click several atoms on ligands A and B to reduce the mapping possibility to what they want. (4) Re-set the atom coordinates of ligand B automatically so that the positions of MCS atoms in ligand B are the same of corresponding ones in ligand A, and the non-MCS atoms in ligand B obtain reasonable new coordinates. (5) Save the new coordinates of ligand B, superpose ligand B to ligand A and show them in one graphic window to let the users have a quick visual check. (6) Write the topology file, parameter file, configuration file and control file which are needed for TI calculations for A→B mutation using the AMBER package. (7) Upon user’s request, convert all necessary files from AMBER style to the style of another simulation package.



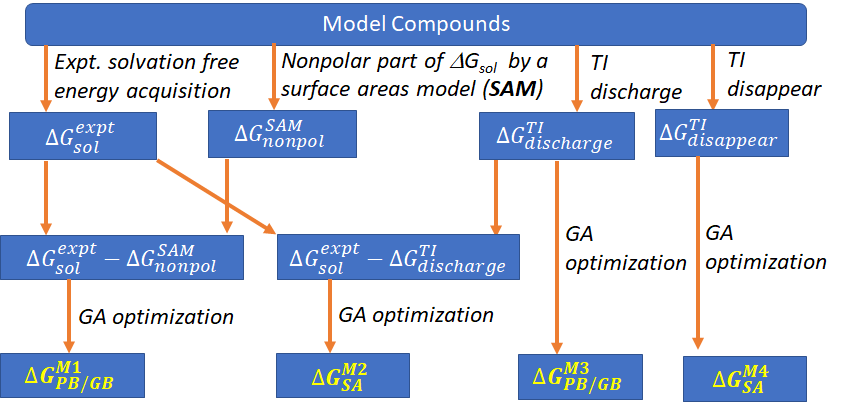
**Figure 7**. **GUIs for Graphic Antechamber (Panel A) and atom pair definition (Panel B)**

We have developed a set of GUIs for some Antechamber programs using PHP (https://mulan.pharmacy.pitt.edu) to automatically get atomic charges and GAFF parameters for arbitrary organic molecules. The graphic antechamber program itself takes the input from JME Molecular Editor (**Fig. 7A**). Occasionally, users may want to define the match atom pairs themselves. We have developed a graphic interface to facilitate users to define atom pairs (**Fig. 7B**). To use this tool, a molecule as the template and another molecule as the target need to load first, then one can define matched atom pairs by clicking atoms. The defined atom pairs are shown in the bottom window and one can delete a pair by click the appended “×”.

**C.3.3 Parameterization of GAFF3 Charge Model-Consistent Implicit Solvation Models.**

MM-PB/GBSA (Molecular Mechanics Poisson-Boltzmann/Generalized Born Surface Area) are two popular endpoint free energy methods in structure-based drug design.[64](#_ENREF_64), [66](#_ENREF_66), [92-95](#_ENREF_92) The performance of MM-PB/GBSA strongly depends on how well calculation errors can cancel out.[66](#_ENREF_66) Therefore, it is necessary to develop a set of solvation models to be consistent with the GAFF3 charge method. The strategy of PBSA and GBSA model development is summarized in **Fig. 8**. In detail, the polar part of a solvation free energy may come from either experimental solvation free energy () or TI free energy of the discharge procedure (). The nonpolar part has also two sources: the TI free energy of atom disappearing procedure () and the difference between the experimental and the TI free energy of the discharge procedure (). A genetic algorithm will be applied to optimize the radius parameters to reproduce the above reference data. Four model sets will be developed: Model Sets M1 and M3 are for the polar part of the solvation free energy, while Model Sets M2 and M4 are for the nonpolar part of solvation free energy.

**Figure 8**. **Implicit solvation model development based on GAFF3 charges.**

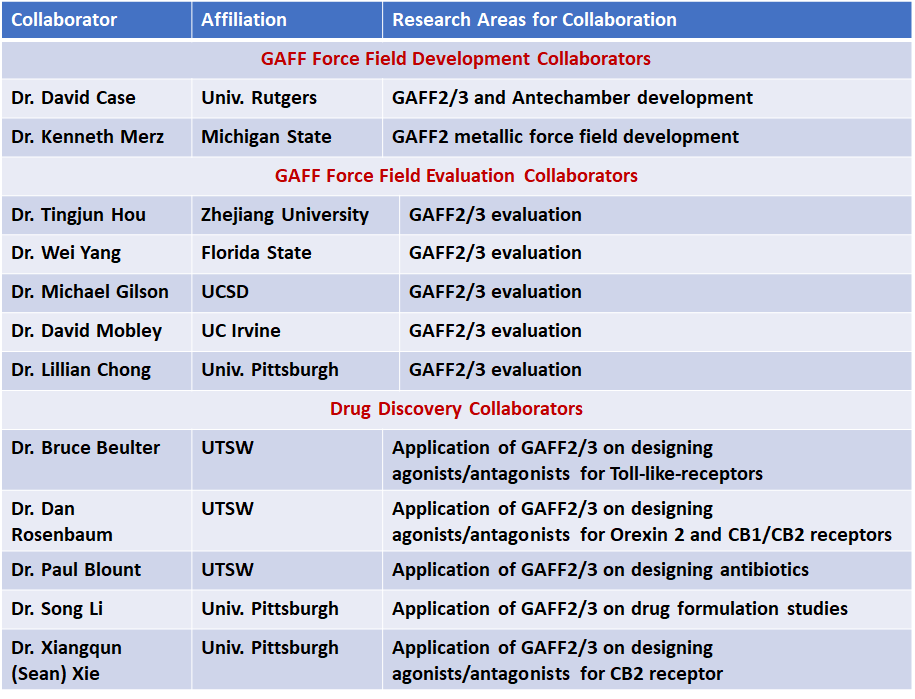


**C3.4 Critical and Systemic Assessment of GAFF2/GAFF3 with TI.**

This most comprehensive data set of relative binding affinities assembled was introduced by Schrödinger Inc. in 2015.[L1] It consists of 200 ligands and 10 different protein targets (Beta-secretase in Alzheimer’s disease, Tyrosine-protein phosphatase non-receptor type 1 in diabetes, thrombin in blood clotting, nonreceptor tyrosine-protein kinase in immunodeficiency, plus the cancer targets Cyclin-dependent kinase 2, Mitogen-activated protein kinase 8, induced myeloid leukemia cell differentiation protein, and P38 mitogenactivated protein kinase). This data set is the current “gold standard” for computational drug design. Another set of 10 to 20 well-studied protein targets and nucleic acid targets (such as DAPI,[96](#_ENREF_96)), which have measured *IC50* or *Ki* for several to tens of diverse small molecule ligands, will be selected to test our GAFF2/GAFF3 force fields.

**C3.5 Systemic Assessment of GAFF2/GAFF3 with MM-PB/GBSA.**

In structure-based drug design, virtual screening becomes an indispensable technique. Endpoint free energy methods, like MM-PB/GBSA, are routinely applied in the last stage to re-rank docking scores. In many cases, the combination of molecular docking and MM/PB(GB)SA rescoring has proven to be a more promising strategy in both the identification of the correct binding poses and the right ranking of the binding affinities of a series of ligands70,72,77,165-171. To evaluate GAFF2/GAFF3 and the newly developed MM-PB/GBSA models, besides the aforementioned systems in C3.4, we plan to study a set of drug targets which have tens, hundreds, even thousands of actives and decoys. We would like to assess both the “Screening Power” and the “Docking Power” of the MM-PB/GBSA models in virtual screenings. We will collaborate closely with Dr. Tingjun Hou at Zhejiang University, who has systematically evaluated the MM/PB(GB)SA methods in virtual screenings for 98 drug targets,[68](#_ENREF_68) to fulfill the task.



**Table 2**. **List of GAFF Development Collaborators**

**C.3.6 Objectively Evaluation of GAFFs Using Drug Discovery Projects.**

Our force field models have been further critically assessed in numerous drug discovery projects where our predictions can be confirmed by experimental biologists. For most of the projects, we intend to develop novel compounds to activate (as agonists) or inhibit (as antagonists) protein targets. As most drug targets are GPCRs and only a few crystal structures are available, protein homology modeling and protein-ligand complex modeling are common tasks. Other common modeling tasks include high-throughput screenings,[97](#_ENREF_97) binding free energy calculations, MD simulations of bioprocesses. The best outcomes for us from those collaboration projects are experimental validations of our molecular modeling prediction (protein-ligand structure models, designed drug candidates, etc.). My collaborators working in wet labs are listed in Table 2.

**D. ADDITIONAL REMARKS on FORCE FIELD DEVELOPMENT**

**Polarizable FF Development.** While additive models will continue to play important roles, polarizable force fields are expected to extend our ability to more adequately study complex biomolecular systems due to their ability to model changing dielectric environments. Atomic polarization effects play a critical role in ligand-receptor interactions, the interactions of ions with nucleic acids, the environmental changes during protein folding, enzymatic mechanism, and low or heterogeneous dielectric environments.[98](#_ENREF_98) Towards the goal of properly including polarization, a great deal of effort has been directed to developing polarizable versions of force fields. A variety of methods have been explored, including induced dipole models,[22-25](#_ENREF_22), [99-101](#_ENREF_99) the fluctuating charge models,[102-106](#_ENREF_102) the Drude oscillator models,[107-111](#_ENREF_107) and the detailed multipole expansion model.[99](#_ENREF_99), [112](#_ENREF_112) In collaboration with Dr. Yong Duan and other AMBER FF consortium PIs, we have developed a set of polarizable FFs based on the framework of Thole’s atomic dipole interaction models. Four papers that document the solid progress on developing the AMBER polarizable FFs have been published.[22-25](#_ENREF_22)  Recently, we have developed a polarizable Gaussian Multipole (**pGM**) model.[113-118](#_ENREF_113) This model represents a novel framework, which applies Gaussian functions to seamlessly treat the multipole polarization and electron penetration effects. We have finished parameterization of atomic polarizability and screening factor parameters for pGM, which outperforms the previous Thole polarizable models in reproducing high-level ab initio polarizability anisotropy. As a member of AMBER consortium team, the PI will develop **pGM-GAFF**, a polarizable GAFF force field based on the framework of pGM.

**Long-Term Goals on GAFF3-Compatible Force Field Development for Biomolecules.** The current approach of studying protein-ligand and nucleic acid-ligand interactions using a combination of the general-purpose and biomolecular force fields may have a consistency problem. Our long-term plan is to develop a set of new GAFF-based biomolecular force fields for proteins, nucleic acids, lipids and carbohydrates. The spectrum of specific force fields will be developed in the spirit of the top-down FF parameterization strategy, i.e. force field parameters are mostly inherited from the new-GAFF and special force field parameters are introduced for biological building blocks only when it is necessary. This development will ultimately lead to a set of highly transferrable, self-consistent, and coherent force fields for studying biomolecule-ligand interactions.

**E. TIMELINE AND MILESTONES**

We have proposed three aims this R01 project. The roadmap and milestones are summarized below:

|  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Objective** | **Force Field Development Aims** | | **Y1** | | **Y2** | | **Y3** | | **Y4** | | **Y5** | |
| **1** | Aim 1a: Additional Torsional Angle Parameterization | |  |  |  | |  | |  | |  | |
| ***2*** | Aim 1b: Extension to all nonmetallic elements | |  | |  |  |  | |  | |  | |
| ***3*** | Aim 1c: Extension to key metallic elements | |  | |  |  |  |  |  | |  | |
| ***4*** | Aim 2a: Charge model development for GAFF3 | |  | |  | |  |  |  |  |  |  |
| ***5*** | Aim 2b: van der Waals parameterization for GAFF3 | |  | |  | |  |  |  |  |  |  |
| ***6*** | Aim 2c: Bonded term parameterization for GAFF3 | |  | |  | |  |  |  |  |  |  |
| ***7*** | Aim 3a: New TI-GPU technique development | |  | |  | |  |  |  |  |  |  |
| ***8*** | Aim 3b: Parameterization for GBSA and PBSA | |  | |  | |  |  |  |  |  |  |
| ***9*** | Aim 3c: Critical evaluation of GAFF2 in studying biomolecule-ligand interactions | |  | |  | |  |  |  |  |  |  |
| ***10*** | Aim 3c: Critical evaluation of GAFF3 in studying biomolecule-ligand interactions | |  | |  | |  |  |  |  |  |  |
| **Preparation** | **First Round Parameterization** | **Refinement** | **Evaluation** | | | | | | | | | |

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