

Correction to A Toolkit to Fit Nonbonded Parameters from and for Condensed Phase Simulations

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J. Chem. Inf. Model. **2016**, *56* (8), 1479–1489. DOI: [10.1021/acs.jcim.6b00280](https://doi.org/10.1021/acs.jcim.6b00280)

In a recent publication we presented a fitting environment for parametrizing point charge (PC) and multipolar (MTP) force fields for condensed-phase simulations.¹ After publication of this work it came to our attention that one of the scripts contained an error which caused an energy component in the free energy simulations to return incorrect values. This affects the optimization of the parameter l when scaling the Lennard-Jones (LJ) parameters according to $\epsilon^* = l\epsilon$ and $R_{\min}^*/2 = lR_{\min}/2$ but not the MTP terms.

Hence, all compounds considered were reparametrized according to the procedure described in ref 1. The corresponding correlations between experiment and the optimized parametrizations are reported in Figures 1 and 2.

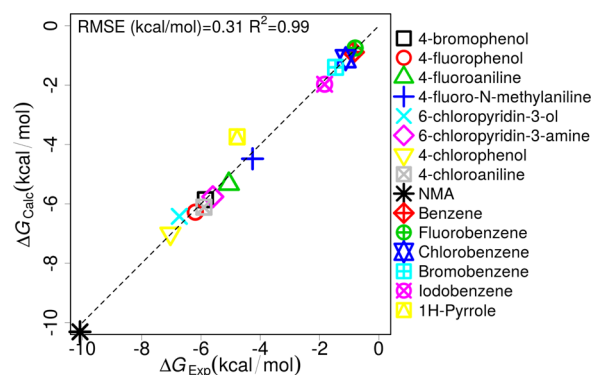


Figure 1. Correlation between experimental and computed solvation free energies ΔG_{hyd} (kcal/mol, respectively, x-axis and y-axis) for a range of compounds of interest. Computed values obtained after optimization of the LJ parameters.

While the best l typically differs by $\Delta l = 0.1$, the average quality of all parametrizations is unchanged. In the published article,¹ the statistical measures for ΔG_{hyd} and ΔH were (RMSE = 0.36 kcal/mol, $R^2 = 0.99$) and (RMSE = 0.53 kcal/mol, $R^2 = 0.97$; see Figures 3 and 4 in ref 1), which changes to RMSE = 0.31 kcal/mol, $R^2 = 0.99$, and RMSE = 0.57 kcal/mol, $R^2 = 0.96$, using the correct script, respectively.

For one example, *N*-methyl-acetamide, the three observables (ρ , ΔH , ΔG_{hyd}) were given explicitly as a function of the scaling l in Table 1 of ref 1. This data has been recomputed and is reported here in Table 1. In this case the same scaling $l = 0.95$ is found to provide the best parametrization, i.e. the one with the lowest score $S = \sum_{i=1}^3 w_i (\text{Obs}_i - \text{Calc}_i)^2$ with $w_\rho = 1$, $w_{\Delta H} = 3$, and $w_{\Delta G} = 5$ which differently weights the three observables.¹ The scores S are now larger in magnitude than in the original

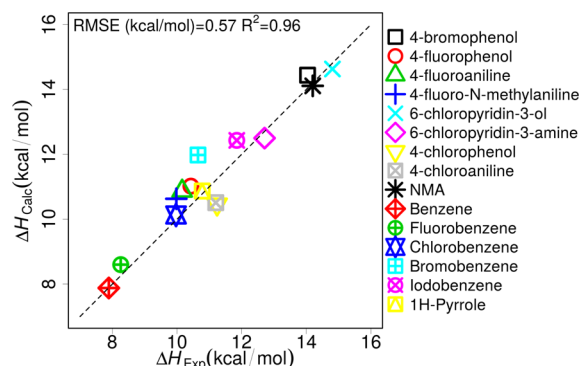


Figure 2. Correlation between experimental and computed enthalpy of vaporization ΔH_{vap} (kcal/mol, respectively, x-axis and y-axis) for a range of compounds of interest. Both, MTP and LJ parameters were optimized.

Table 1. Dependence of ρ (g/cm³), ΔH_{vap} , and ΔG_{hyd} (both in kcal/mol) When Scaling the Lennard-Jones Parameters^a

scaling l	ρ	ΔH_{vap}	ΔG_{hyd}	score S
0.9	1.13	14.24	-10.57	1.2
0.925	1.08	13.95	-10.41	0.8
0.95	1.00	14.11	-10.31	0.3
0.975	0.99	13.84	-10.23	0.5
1	0.95	13.82	-9.78	0.9
1.025	0.92	13.68	-9.27	4.1
1.05	0.88	13.57	-9.01	6.9
1.075	0.84	13.29	-8.21	20.0
1.1	0.81	13.47	-7.98	23.7
exp	0.94 ^{2,3}	14.2 ^{2,4}	-10.08 ⁵	

^aIn bold face is shown the value of l minimizing the score S .

work¹ because the results from the hydration free energy simulations differ.

The current results show that the quality of the parametrizations and all conclusions from the original article remain unchanged. However, the value of the scaling l that is required for a particular quality of a parametrization changes.

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Published: January 5, 2017

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