

Optimization and Application of Lithium Parameters for the Reactive Force Field, ReaxFF

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To make a practical molecular dynamics (MD) simulation of the large-scale reactive chemical systems of Li–H and Li–C, we have optimized parameters of the reactive force field (ReaxFF) for these systems. The parameters for this force field were obtained from fitting to the results of density functional theory (DFT) calculations on the structures and energy barriers for a number of Li–H and Li–C molecules, including Li₂, LiH, Li₂H₂, H₃C–Li, H₃C–H₂C–Li, H₂C=C–LiH, HC≡CLi, H₆C₅–Li, and Li₂C₂, and to the equations of state and lattice parameters for condensed phases of Li. The accuracy of the developed ReaxFF was also tested by comparison to the dissociation energies of lithium–benzene sandwich compounds and the collision behavior of lithium atoms with a C₆₀ buckyball.

1. Introduction

Organolithium compounds, which are increasingly important reactive intermediates, are frequently the reagents of choice for a variety of synthetic purposes because they are often more useful than other alkali metal and magnesium compounds.^{1,2} However, reliable experimental data on lithium compounds is scarce. Also, because the application of high-level ab initio calculations demands high computer capacity, the development of an accurate force field (FF) for lithium is very important for carrying out molecular dynamics simulations. Generic FFs such as DREIDING³ and UFF⁴ allow predictions for broad classes of compounds, particularly when coupled with charge equilibration (QEq)⁵ or other methods for predicting charges. However, in general, these force fields do not describe chemical reactivity, whereas the ReaxFF^{6–8} can simulate the breaking and reforming phenomena of bonds during dynamics. Additionally, the ReaxFF is able to predict accurately not only the reactivity of bonds in the polymer systems but also the crystal and mechanical properties of the condensed phases.^{6–8}

In the past decade, the semiempirical MO (molecular orbital) methods of MNDO⁹ and PM3¹⁰ for lithium compounds have been applied to obtain properties such as the heat of formation, bond length, and dipole moment. However, both methods are available to a few gas phases only; in other words, they cannot describe the condensed phases appropriately. Also, Li/MNDO is known to have some severe deficiencies (e.g., overestimation of the C–Li and H–Li interactions⁹).

In this article, we report lithium parameters for the ReaxFF. The parameters are developed by using DFT results of various organolithium compounds, and the applicability of the new parameter set is subsequently demonstrated for two cases: (i) the dissociation energies of lithium–benzene sandwich compounds, giving useful information on graphite anodic electrodes in Li ion batteries and (ii) the collision behavior of lithium atoms

with a C₆₀ buckyball, thereby elucidating the formation mechanism of an endohedral Li@C₆₀ complex. (Here, the notation Li@C₆₀ denotes a LiC₆₀ molecule with one Li atom encapsulated inside a C₆₀ cage.)

2. Computational Details

The ReaxFF framework was initially developed for hydrocarbons.⁶ It was thereafter successfully employed in the study of Si/SiO₂ interfaces,⁷ where the system energy is partitioned into several partial energy contributions in eq 1.

$$E_{\text{system}} = E_{\text{bond}} + E_{\text{over}} + E_{\text{under}} + E_{\text{lp}} + E_{\text{val}} + E_{\text{pen}} + E_{\text{tors}} + E_{\text{conj}} + E_{\text{vdWaals}} + E_{\text{Coulomb}} \quad (1)$$

Note that the explanation for each energy term was reported in detail in a previous work.⁷ Unlike the ReaxFF for hydrocarbons⁶ and Si/SiO₂⁷ and the total energy expression for organolithium in particular, Li–C–H systems can be considered to be a summation of electrostatic (Coulomb), bond, overcoordination, and van der Waals energies:

$$E_{\text{system}} = E_{\text{bond}} + E_{\text{over}} + E_{\text{vdWaals}} + E_{\text{Coulomb}} \quad (2)$$

Because of the relatively large ionic character of the Li–H and Li–C bonds,^{11,12} angle bending and torsion terms in the total energy are not included because they were all set equal to zero. The Coulomb interactions in our ReaxFF were calculated between all atom pairs with the atomic charges. They are determined for each configuration using the electron equilibration method (EEM).¹³ We optimized the EEM parameters (electronegativity χ , chemical hardness η , and shielding radius r) to reproduce the charge distribution of numerous clusters, shown in Figure 1, involving Li, C, and H obtained from DFT calculations using the JAGUAR code¹⁴ with the B3LYP (Becke three-parameters plus Lee–Yang–Parr) functional and a 6-31G** basis set. The EEM parameters for C and H were determined in a previous study.⁶

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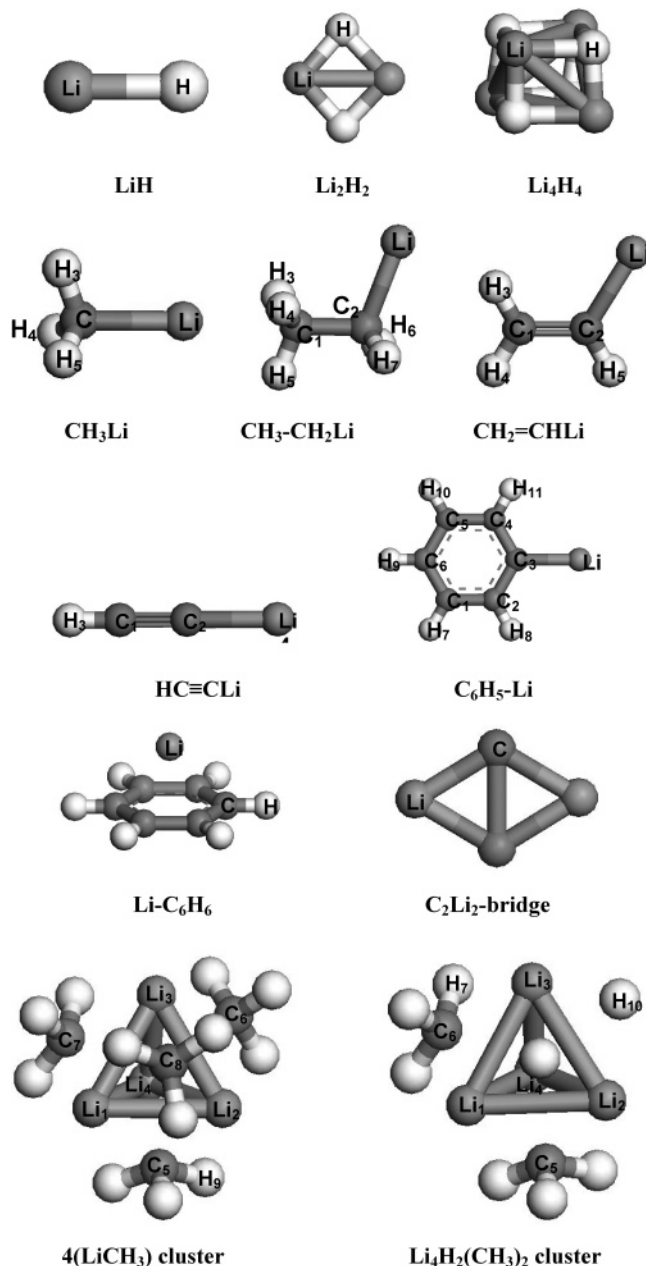


Figure 1. Molecules used to develop the ReaxFF parameters of Li in this study.

For Li crystals, we fitted various phases of Li, including fcc (12), hcp (12), bcc (8), sc (6), and diamond (4), which effectively allowed us to vary the coordination number of Li. In DFT calculations on our periodic systems, we employed the generalized gradient approximation of the Perdew–Burke–Ernzerhof functional¹⁵ for the exchange–correlation potential and ultrasoft pseudopotentials to replace the core electrons, implemented in the CASTEP code.¹⁶ We set a kinetic energy cutoff of 180.0 eV for Li and used the Monkhorst–Pack scheme¹⁷ to generate the k-space grid. We also found that a k-point sampling of 666 was sufficient for convergence.

3. Results and Discussion

3.1. Bond Dissociation. Structures depicted in Figure 1 are determined through full geometry optimizations to find ground-state structures. Dissociation profiles are then constructed through total energy calculations at the modified geometries only

by changing the bond length from its equilibrium value, which is shown in Figure 2.

To optimize the ReaxFF bond energies, we first carried out DFT calculations for the dissociation of single bonds of Li–Li and Li–H. Parts a and b of Figure 2 provide comparisons of the DFT and ReaxFF results for single-bond dissociations of Li–Li and Li–H in Li_2 and LiH systems. We used DFT methods for the singlet state from the equilibrium distance up to the point where it was comparable to the dissociation limit (to the lowest of either the singlet or triplet full bond dissociation energies). In Figure 2a, the equilibrium bond distance and bond dissociation energy of a Li_2 molecule, calculated by DFT, are 2.73 Å and 20.24 kcal/mol, respectively. Fuentealba and Reyes¹⁸ reported that the theoretical (B3PW91 level and 6-311G** basis set) and experimental dissociation energies for Li_2 are 18.54 and 24.42 kcal/mol, respectively, thus verifying that our DFT calculation result is reliable. Using the ReaxFF, the bond distance and bond dissociation energy are 2.72 Å and 33.28 kcal/mol, respectively. According to MNDO and PM3 calculations, the bond distances of Li_2 are 2.48⁹ and 2.05 Å,¹⁰ respectively, which implies that the ReaxFF estimates the bond distance more accurately than the MNDO and PM3 methods. However, it overestimates the dissociation energy of the Li_2 molecule. When the parameters of Li for the ReaxFF were fitted in the present study, gas phases as well as crystal information as reference data calculated by DFT were used. Because the bond energy of Li–Li^+ is 33.7 kcal/mol¹⁹ and the cohesive energy of a Li crystal with bcc structure is 37.7 kcal/mol,²⁰ the bond character between two Li atoms in the solid Li is Li–Li^+ , not Li–Li .¹⁹ The ReaxFF cannot provide an accurate description of the ionic state. Therefore, when the ReaxFF parameters of Li were optimized, the Li–Li single-bond dissociation energy in a Li_2 molecule was overestimated to describe accurately information for the condensed phases of Li, causing a certain degree of inaccuracy for Li_2 molecules.

For a LiH molecule (Figure 2b), the DFT shows an equilibrium bond distance of 1.62 Å and a bond dissociation energy of 58.95 kcal/mol, and the ReaxFF yields estimates of 1.58 Å and 59.66 kcal/mol, which are in good agreement with the DFT results. The reliability of our results for LiH is verified by comparison with those of Fuentealba and Reyes.¹⁸ They reported that the theoretical and experimental dissociation energies for LiH are 69.97 and 58.06 kcal/mol, respectively. The MNDO and PM3 methods predict the bond distance to be 1.54⁹ and 1.38 Å,¹⁰ respectively, thus demonstrating that the ReaxFF describes the bond distance of a LiH molecule more accurately than these approaches.

Parameters for the Li–C single bond were also determined by DFT calculations on CH_3Li , $\text{CH}_3\text{–CH}_2\text{Li}$, $\text{CH}_2\text{=CHLi}$, $\text{HC}\equiv\text{CLi}$, and $\text{C}_6\text{H}_5\text{Li}$ molecules. Parts c–g of Figure 2 show curves for single-bond dissociations of Li–C for the CH_3Li , $\text{CH}_3\text{–CH}_2\text{Li}$, $\text{CH}_2\text{=CHLi}$, $\text{HC}\equiv\text{CLi}$, and $\text{C}_6\text{H}_5\text{Li}$ systems. The optimized conformational information of each molecule is provided in Table 1. The information in Table 1 shows that the ReaxFF is superior to the MNDO and PM3 methods in terms of a conformational prediction of the above molecules. For a methyllithium (CH_3Li) molecule (Figure 2c and Table 1), the DFT results show that the equilibrium Li–C bond distance is 1.98 Å and the bond dissociation energy is 55.12 kcal/mol. According to the ReaxFF method, the Li–C bond distance is 2.09 Å, and the bond dissociation energy is 52.71 kcal/mol, which is about 3 kcal/mol lower than that obtained through the DFT calculation. For the $\text{CH}_3\text{–CH}_2\text{Li}$ molecule (Figure 2d and Table 1), the DFT calculation indicates that the equilibrium

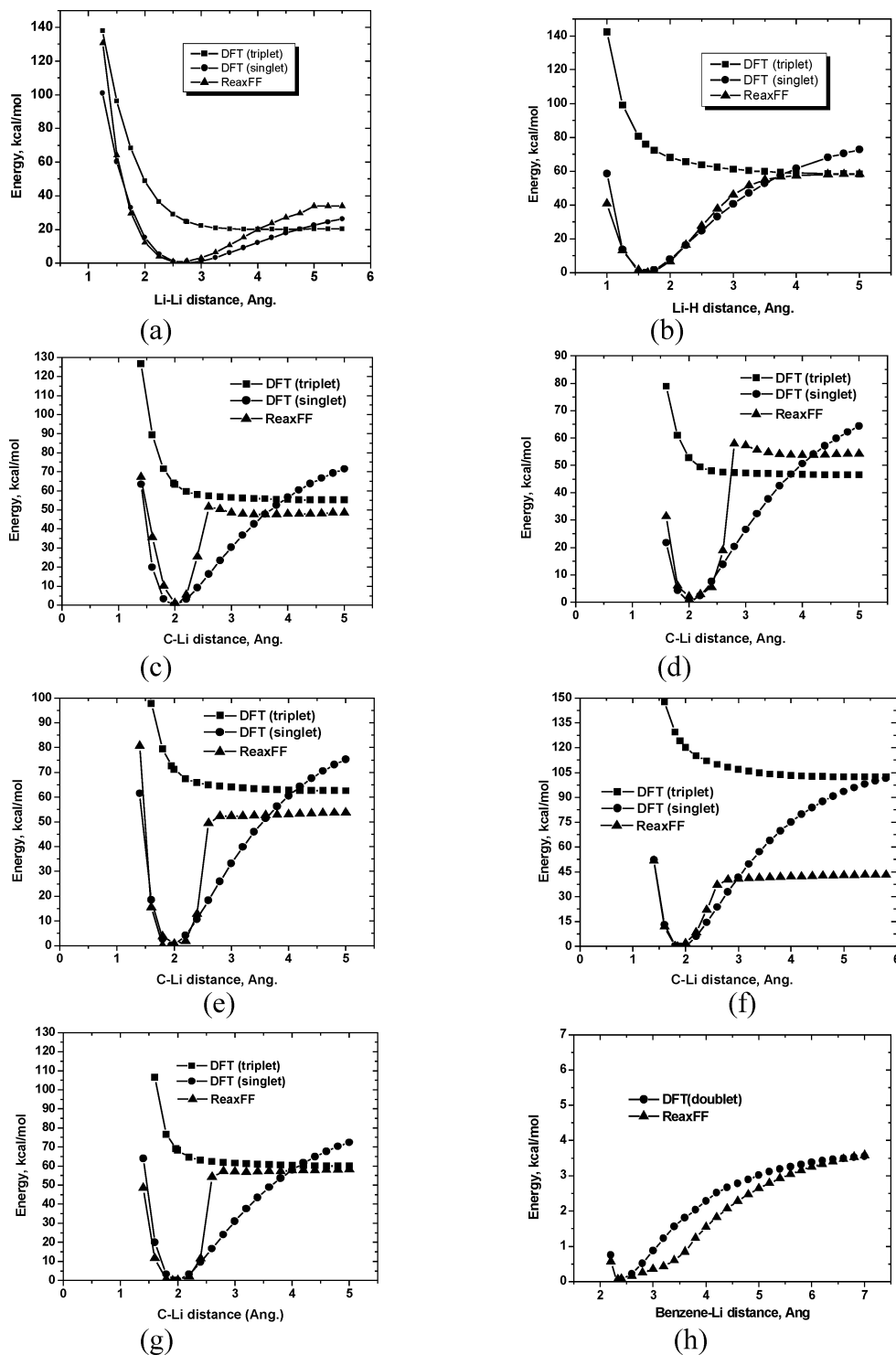


Figure 2. DFT and ReaxFF data for bond dissociations of various Li-C-H systems: (a) the Li-Li single bond in Li_2 , (b) the Li-H bond in LiH , (c) the Li-C bond in CH_3Li , (d) the Li-C bond in $\text{CH}_3\text{-CH}_2\text{Li}$, (e) the Li-C bond in $\text{CH}_2\text{=CHLi}$, (f) the Li-C bond in $\text{HC}\equiv\text{CLi}$, (g) the Li-C bond in $\text{C}_6\text{H}_5\text{Li}$, and (h) the van der Waals interaction of a Li atom on a benzene ring.

Li-C bond distance is 2.01 Å and the bond dissociation energy is 46.51 kcal/mol. The ReaxFF gives an equilibrium Li-C bond distance of 2.01 Å, which is very similar to the DFT value, and a bond dissociation energy of 55.15 kcal/mol, which is higher than the DFT value. One C-C bond in the $\text{CH}_3\text{-CH}_2\text{Li}$ molecule is a single bond, and the Li-C bonding character may be affected by the C-C bond order near the Li-C bond. To investigate this, we additionally considered $\text{CH}_2\text{=CHLi}$ and $\text{HC}\equiv\text{CLi}$ molecules. The bond order between two carbon atoms in $\text{CH}_2\text{=CHLi}$ is 2 (a double bond—one σ bond and one π

bond), and that in $\text{HC}\equiv\text{CLi}$ is 3 (a triple bond—one σ bond and two π bonds). As shown in Figure 2e and Table 1, the DFT calculation reveals that the equilibrium distance of the Li-C bond in the $\text{CH}_2\text{=CHLi}$ molecule (C-C bond order of 2) is 1.95 Å, which is shorter than that (2.01 Å) of the $\text{CH}_3\text{-CH}_2\text{Li}$ molecule. The dissociation energy of the Li-C bond is 62.53 kcal/mol, which is larger than that (46.51 kcal/mol) of $\text{CH}_3\text{-CH}_2\text{Li}$. The ReaxFF indicates that the bond distance of Li-C is 1.77 Å, which is 0.18 Å shorter than the DFT result, and the dissociation energy of the bond is estimated to be 58.57

TABLE 1: Optimized Geometries of Various Organolithium Molecules

parameter ^a	DFT	ReaxFF	PM3 ^b	MNDO ^c	parameter ^a	DFT	ReaxFF	PM3 ^b	MNDO ^c
					CH ₃ Li				
$r(\text{CLi})$	1.978	2.099	2.525	1.821	$\angle(\text{H}_3\text{CLi})$	112.47	110.87	91.74	111.92
$r(\text{CH}_3)$	1.100	1.124	1.063	1.117	$\angle(\text{H}_3\text{CH}_4)$	106.31	108.03	119.91	106.92
					CH ₃ -CH ₂ Li				
$r(\text{C}_2\text{Li})$	2.005	2.029	2.868	1.829	$\angle(\text{H}_6\text{C}_2\text{Li})$	108.28	113.53	95.01	116.05
$r(\text{C}_1\text{C}_2)$	1.542	1.523	1.430	1.528	$\angle(\text{C}_1\text{C}_2\text{Li})$	117.46	95.90	95.35	97.04
$r(\text{C}_2\text{H}_6)$	1.103	1.118	1.075	1.114					
					CH ₂ =CHLi				
$r(\text{C}_2\text{Li})$	1.954	1.797	2.862	1.784	$r(\text{C}_2\text{H}_5)$	1.100	1.162	1.097	1.094
$r(\text{C}_1\text{C}_2)$	1.349	1.274	1.319	1.352	$\angle(\text{H}_5\text{C}_2\text{Li})$	127.49	136.84	125.45	144.70
$r(\text{C}_1\text{H}_3)$	1.095	1.121	1.090	1.112	$\angle(\text{C}_1\text{C}_2\text{Li})$	120.17	101.89	118.56	119.59
					HC≡CLi				
$r(\text{C}_2\text{Li})$	1.897	1.849	2.868	1.744	$\angle(\text{C}_1\text{C}_2\text{Li})$	180.00	180.00	180.00	180.00
$r(\text{C}_1\text{C}_2)$	1.230	1.216	1.216	1.213	$\angle(\text{H}_2\text{C}_1\text{C}_2)$	180.00	180.00	180.00	180.00
$r(\text{C}_1\text{H}_3)$	1.068	1.101	1.057	1.052					
					C ₆ H ₅ Li				
$r(\text{C}_3\text{Li})$	1.973	1.920	2.857	1.797	$\angle(\text{C}_2\text{C}_3\text{C}_4)$	114.60	122.20	116.34	117.85
$r(\text{C}_2\text{C}_3)$	1.416	1.388	1.393	1.413	$\angle(\text{C}_2\text{C}_3\text{Li})$	122.68	118.90	121.86	121.10
$r(\text{C}_2\text{H}_6)$	1.092	1.112	1.095	1.097	$\angle(\text{C}_3\text{C}_4\text{H}_{11})$	119.73	121.63	118.35	120.02

^a See Figure 1. ^b Reference 10. ^c Reference 9.

kcal/mol, which is quite reasonable. Results for HC≡CLi (C-C bond order of 3) are presented in Figure 2f and Table 1. According to the DFT calculation, the bond distance and bond dissociation energy of Li-C in HC≡CLi are 1.90 Å and 102.79 kcal/mol, respectively. The ReaxFF provides a good description of the conformation of the molecule. However, there is a large difference between the dissociation energy of the Li-C bond predicted by the ReaxFF and DFT, revealing the most serious problem in the present ReaxFF for Li. We also considered phenyl-lithium (C₆H₅Li), which has a C-C bond order of 1.5. In Figure 2g and Table 1, the ReaxFF correctly describes the dissociation energy of the Li-C bond and conformation information of phenyl-lithium. From the dissociation curves (parts c-g of Figure 2) of Li-C bonds in various organolithium molecules expressed thus far, we also found that the dissociation curves estimated by the ReaxFF method have a U shape. For instance, in the case of phenyl-lithium (Figure 2g), the dissociation curve appears to be U-shaped up to about 2.8 Å, after which the energy remains constant, thus indicating that a Li-C bond in phenyl-lithium is broken at about 2.8 Å. From the DFT calculation, Li-C bond breaking occurs from 2.6 to 2.8 Å, which is consistent with the ReaxFF result. From the results thus far, we can determine that the Li-C bonding character in an organolithium system is affected by a C-C bond near the Li-C bond. In other words, the higher the bond order of a C-C bond in organolithium molecules, the shorter the bond distance of the Li-C bond near the C-C bond and the higher the bond energy of the Li-C bond.

3.2. van der Waals Interaction. The van der Waals interaction in the ReaxFF is used with a distance-corrected Morse potential including a shielded interaction.^{6,7} Parameters for a Li atom related to the van der Waals interaction are the van der Waals radius (r_{vdW}), van der Waals dissociation energy (ϵ), van der Waals parameter (α), and van der Waals shielding (γ_{vdW}). To determine these parameters, we considered the interaction of a Li atom on a benzene ring (Figure 1). Figure 2h indicates the change in the van der Waals energy of a Li atom on a benzene ring with respect to the distance from the Li atom to the center of the ring. The ReaxFF can quite accurately explain the van der Waals interaction of a Li atom on a benzene ring. The DFT calculation indicates that 2.33 Å is the distance between the Li atom and the center of the benzene ring in

optimized structure, whereas the ReaxFF yields a very similar prediction of 2.32 Å. The DFT calculation in this study was performed with the B3LYP functional. The DFT method may not provide a good explanation of the van der Waals interaction.²¹ To assess the disadvantage of the B3LYP, which may occur in the van der Waals interaction of the Li atom upon the benzene ring, we also used the X3LYP functional²² for the system. The X3LYP leads to an accurate description of van der Waals.²³ For example, previously reported high-quality ab initio calculations [MP4/6-311G(d,p)/MP2/6-311G(d,p)] indicated that a Li cation binds six H₂ molecules at zero Kelvin with enthalpies for adding successive H₂ of -5.39, -4.30, -4.07, -3.65, -1.87, and -2.30 kcal/mol;²³ these values agree well with the X3LYP results of -5.12, -4.47, -3.9, -3.63, -1.55, and -1.52 kcal/mol for the same system.²⁴ The X3LYP result for the system considered in this study is very similar to the B3LYP result. Therefore, we confirmed that the B3LYP functional can appropriately describe the van der Waals interaction of a Li atom on a benzene ring.

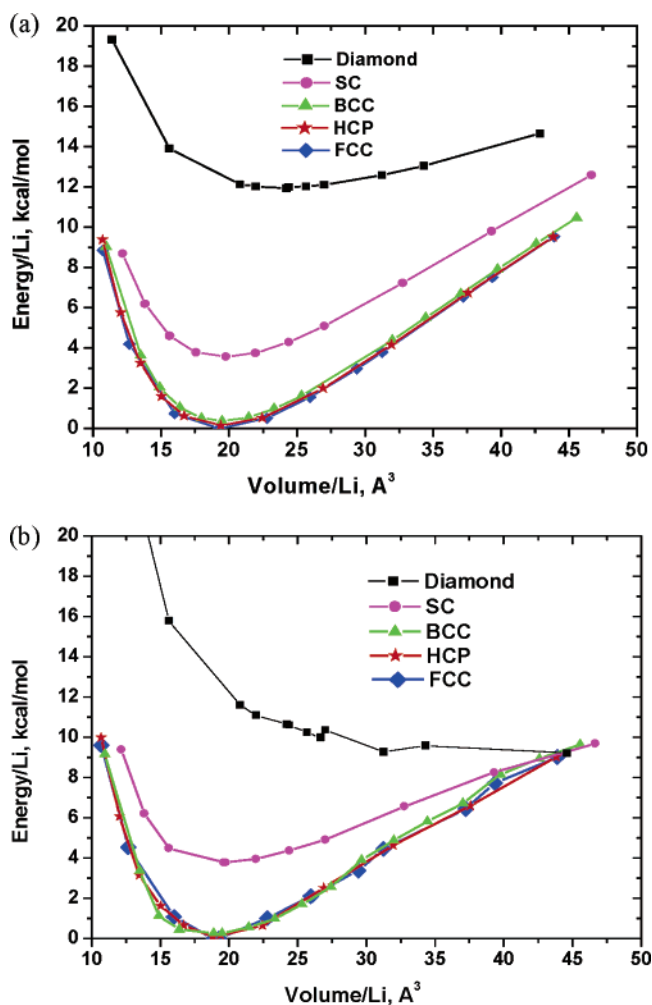
3.3. Charge Distribution. As already mentioned, in the case of lithium, the ionic bond character is larger than the covalent bond character.^{11,12} Hence, we did not consider valence angle terms such as C-Li-C and Li-C-Li. Instead, we focused on atomic charge terms such as EEM electronegativity (χ), EEM hardness (η), and EEM shielding (γ) in optimizing the ReaxFF parameters. Shown in Figure 1 are all of the molecules that we considered to determine the three atomic charge terms of Li. Charge distributions in the ReaxFF are calculated using the EEM method.^{6,7} The EEM parameters were optimized against the Mulliken charge distribution obtained from DFT calculations. The ReaxFF results for the atomic charges of all molecules depicted in Figure 1 are summarized in Table 2. For a Li₂H₂ molecule with a bridged structure, the DFT/Mulliken partial charges for Li and H atoms are 0.1670 and -0.1670, respectively, whereas the ReaxFF calculates partial charges of 0.1696 and -0.1696 for the Li and H atoms, respectively. In the case of methyl-lithium (CH₃Li), the ReaxFF assigns partial charges to C, H, and Li of -0.4649, 0.0771, and 0.2318, respectively, compared to DFT/Mulliken values of -0.5507, 0.0715, and 0.3362, respectively. Overall, the ReaxFF gives a good prediction of the partial atomic charges of various organolithium molecules except for Li on a benzene ring.

TABLE 2: Optimized Charge Distributions of Various Organolithium Molecules

molecule ^a	DFT	ReaxFF	molecule ^a	DFT	ReaxFF
Li	0.3853	0.1398	LiH	-0.3853	-0.1398
Li	0.1670	0.1696	Li ₂ H ₂	-0.1670	-0.1696
Li	0.1993	0.2001	Li ₄ H ₄	-0.1993	-0.2001
C	-0.5507	-0.4649	CH ₃ Li	0.0715	0.0771
Li	0.3362	0.2318	H ₃	0.0631	0.0795
C ₁	-0.2895	-0.2888	CH ₃ -CH ₂ Li	0.0685	0.0904
C ₂	-0.3951	-0.3365	H ₅	0.0631	0.0795
H ₃	0.0776	0.0624	H ₆	0.0631	0.0795
H ₄	0.0776	0.0624	H ₇	0.0631	0.0795
C ₁	-0.2252	-0.2143	CH ₂ =CHLi	0.0592	0.0933
C ₂	-0.2543	-0.3256	H ₄	0.0355	0.0972
H ₃	0.0489	0.0487	H ₅	0.0355	0.0972
C ₁	-0.2720	-0.1131	Li	0.3359	0.3006
C ₂	-0.2033	-0.2651	HC≡CLi	0.0938	0.0881
C ₁	-0.1347	-0.0894	H ₃	0.3815	0.2901
C ₃	-0.1233	-0.1798	Li	0.3815	0.2901
Li	-0.2472	0.2966	C ₆ H ₅ Li	0.0376	0.0625
C	-0.0821	-0.1290	H ₈	0.3767	0.2672
Li	0.1695	0.2404	Li	0.3767	0.2672
Li ₁	0.2876	0.3900	C ₂ Li ₂ Bridge	-0.1695	-0.2404
Li ₂	0.2849	0.3912	C	-0.1695	-0.2404
Li ₃	0.2857	0.3908	4(LiCH ₃) Cluster	-0.6082	-0.5211
Li ₄	0.2850	0.3904	C ₆	-0.6081	-0.5213
C ₅	-0.6078	-0.5213	C ₇	-0.6080	-0.5212
Li ₁	0.3814	0.2334	C ₈	0.1074	0.0435
Li ₂	0.1034	0.3221	C ₉	-0.6155	-0.5329
Li ₃	0.1047	0.3220	C ₆	-0.6165	-0.5329
Li ₄	0.3719	0.2334	H ₇	0.1074	0.0588
			H ₁₀	-0.1905	-0.1894

^a See Figure 1.

3.4. Crystal. Simulations of the condensed-phase Li can be a key application for the ReaxFF. The ability of the ReaxFF potential to predict condensed-phase stabilities was tested against a variety of crystal structures for Li. For the five lithium crystal structures of fcc, hcp, bcc, sc, and diamond, the DFT energies were obtained for a broad range of compression and expansion and then compared against the ReaxFF in Figure 3. Also, for each phase of Li, the lattice parameters and phase stabilities were compared against DFT data, which is expressed in Tables 3 and 4. The ReaxFF correctly describes equations of state of fcc, hcp, bcc, and sc; however, for the diamond-structured Li crystal, the ReaxFF overestimates the lattice parameter of the diamond structure. Except for the diamond structure case, the ReaxFF predicts the lattice parameters very well and also accurately estimates the phase stability for the five Li crystals. According to several experimental²⁵ and theoretical^{26,27} studies, Li is in a close-packed structure (fcc or hcp) at low pressure and temperature and is in a bcc structure at room temperature, which supports our DFT results for Li crystals. Figure 3a and Table 4 reveal that the fcc structure for Li is most stable at zero Kelvin. Moreover, we also discovered that equations of

**Figure 3.** Equations of state (compression and tension) for five crystal structures (HCP, FCC, BCC, SC, and diamond) of lithium calculated using (a) DFT and (b) ReaxFF methods.**TABLE 3: Lattice Parameters for Five Structures (HCP, FCC, BCC, SC, and Diamond) of Lithium Calculated by DFT and ReaxFF**

crystal structure	lattice parameter (Å)	
	DFT	ReaxFF
HCP	a: 3.048, c: 4.819	a: 3.052, c: 4.827
FCC	4.257	4.291
BCC	3.390	3.363
simple cubic (SC)	2.704	2.642
diamond	5.785	7.179

TABLE 4: Phase Stabilities (kcal/mol of Li) for Five Lithium Structures Relative to BCC-Structured Li

crystal structure	ΔE_{DFT}	ΔE_{ReaxFF}
HCP	-0.2325	-0.2483
FCC	-0.3724	-0.3392
SC	3.1997	2.7238
diamond	11.5681	11.0107

state for fcc, hcp, and bcc structures of Li almost overlapped with volume change. For this reason, the crystal structure of Li may be transformed relatively easily by pressure and/or temperature. Young and Ross²⁶ predicted two transitions below 1 Mbar on the 0 K isotherm: hcp-fcc at about 4 kbar and fcc-bcc at about 86 kbar. Also, we compared the cohesive energy of bcc Li, calculated by DFT and ReaxFF. The cohesive energy by the DFT calculation is -43.00 kcal/mol, whereas the ReaxFF

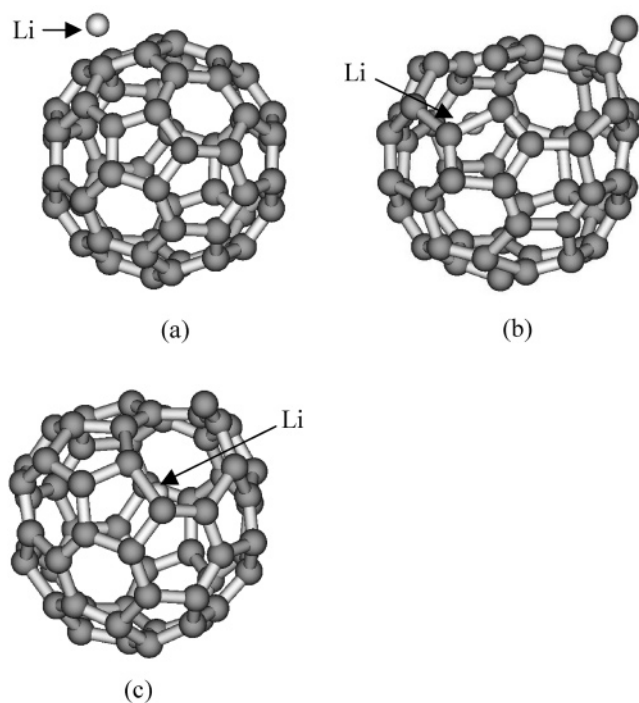


Figure 5. Snapshots of the MD simulation for the formation mechanism of the endohedral [Li@C₆₀] complex where the Li atom hits the center of a six-membered ring of C₆₀ with 13 eV of kinetic energy, taken at simulation times of (a) 0 ps, (b) 0.2 fs, and (c) 10 ps. Here, yellow and gray atoms refer to carbon and hydrogen atoms, respectively.

According to the MD simulation with the ReaxFF, the lowest incident energy needed for a Li atom to penetrate a hexagon ring on the C₆₀ buckyball and form the endohedral [Li@C₆₀] complex is estimated to be around 13 eV (Figure 5), which is approximately twice as large as the ab initio MD simulation result.³³ The difference in the threshold energy is mainly due to the difference in the radii of Li and Li⁺. In the case of the ab initio MD simulation,³³ the interaction between Li⁺ and C₆₀⁻ was investigated while we focused on that between neutral Li and C₆₀. Because the ionic radius of Li⁺ is $\sim 0.6\text{--}0.9$ Å and the hole of a hexagon ring is the same size or slightly smaller,³³ Li⁺ can penetrate into the cage through the center of the hexagon ring relatively easily. However, the atomic radius of Li is ~ 1.5 Å, which is about twice the size of the radius of Li⁺. This results in a higher threshold energy for the formation of the endohedral [Li@C₆₀]. As shown in Figure 12, the Li atom in the formed endohedral [Li@C₆₀] complex is at an off-center position in the C₆₀ cage, which is in good agreement with previous theoretical works.^{34,35} When the incident energy of Li is below 13 eV, the Li atom cannot enter the C₆₀ cage and stays outside the center of the hexagon of the cage. Our ReaxFF simulation also indicates that except for a Li atom hitting a six-membered ring of C₆₀, the Li atom cannot be inserted into C₆₀ although the incident energy of Li is increased up to 30 eV, similar to the ab initio MD simulation.³³ From these results, we can determine that the endohedral [Li@C₆₀] can be formed by the bombardment of a Li atom with a C₆₀.

4. Summary

We have developed the ReaxFF for Li, Li–H, and Li–C systems. The ReaxFF for Li has been tested against a substantial data set derived from DFT calculations on small clusters and condensed systems covering both reactive and nonreactive aspects of organolithium. Also, the developed ReaxFF is similar

or more accurate in expressing organolithium systems than the semiempirical MO methods, PM3 and MNDO. However, the ReaxFF overestimates the van der Waals interaction of Li with benzene rings. Using MD simulations with the ReaxFF, we also found that endohedral [Li@C₆₀] can be formed by the bombardment of a Li atom with a C₆₀ if one Li atom with an incident energy of above 13 eV hits a six-membered ring of C₆₀ perpendicularly. We believe that this ReaxFF for Li can be used extensively to simulate aspects of organolithium reactivity and interactions between Li and CNT and fullerene.

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Appendix

The ReaxFF parameters for Li optimized in this work are as follows. Here, symbols of the parameters are shown in refs 6 and 7.

TABLE A1: Atom Parameters

r_0 (Å)	$p_{ov/un}$	Coulomb parameters			van der Waals parameters				
		η (eV)	χ (eV)	γ (Å)	r_{vdw} (Å)	ϵ (kcal/mol)	α	γ_{vdw} (Å)	
Li	2.17	-3.08	8.72	1.23	4.42	2.57	0.014	13.62	1.08

TABLE A2: 1–3 Bond Order Correction Parameters

	λ_3	λ_4	λ_5
Li	5.39	0.48	0.05

TABLE A3: Bond Energy and Bond Order Parameters

bond	D_e^σ	$p_{be,1}$	$p_{b0,1}$	$p_{b0,2}$
Li–Li	40.7	-0.327	-0.064	4.22
Li–C	57.3	-0.002	-0.009	20.49
Li–H	60.5	-8.065	-0.066	5.01

TABLE A4: van der Waals and Bond Radius Parameters

bond	r^σ (Å)	r_{vdw} (Å)	ee (kcal/mol)	aa
Li–C	1.971	1.066	0.292	29.32
Li–H	1.669	1.026	0.406	14.22

References and Notes

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