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Catalytic Reactions of Living Polymers: Density Functional Study of Reactivity of Phenol and Phenoxides with the Cyclic Tetramer of Polycarbonate

P. Ballone,[†] B. Montanari,[‡] and R. O. Jones*

Institut für Festkörperforschung, Forschungszentrum Jülich, D-52425 Jülich, Germany

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The reactivity of phenol, lithium phenoxide (LiOPh), and sodium phenoxide (NaOPh) with the cyclic tetramer of bisphenol A polycarbonate (BPA-PC) has been investigated using density functional calculations. The potential energy of the system is computed using a suitable reaction coordinate and relaxing all other degrees of freedom by Car–Parrinello molecular dynamics. Both LiOPh and NaOPh catalyze ring opening with small energy barriers ($\Delta E = 4.0, 2.5$ kcal/mol, respectively) to a chain with a phenyl carbonate at one end and a phenoxide at the other, a “living polymer”. The barrier is large for phenol ($\Delta E > 40$ kcal/mol), but the total energy differences between the reactants and the chain are very small in all three molecules. We discuss the balance between changes in entropy and energy, and we compute the vibrational properties of metastable intermediate species.

1. Introduction

The wide range of industrial applications of polycarbonates has led to many studies, and bisphenol A polycarbonate (BPA-PC) has been a prototype for both theorists and experimentalists.¹ Information about the geometric arrangement of the atoms is essential for discussing the properties of any material. The most detailed has been provided in the polycarbonate context by X-ray diffraction studies of molecules closely related to BPA-PC, examples being the crystalline forms of DPBC [4,4'-isopropylidenediphenylbis(phenyl carbonate)].² Furthermore, the cyclic dimer, trimer, and tetramer carbonates of BPA have been isolated and analyzed spectroscopically,^{3,4} and the structures of the crystalline forms of dimer and tetramer determined by X-ray diffraction.⁴ The cyclic tetramer of BPA-PC (Figure 1)^{3,4} provides an ideal model for theoretical studies of the microscopic properties of polycarbonate. It has a very large, open cyclic structure, with a cavity of 14–16 Å and outside diameter of 20.7 Å,⁴ and the size of the ring (132 atoms) means that there is no hindrance to ring-flipping or *cis-trans-trans-trans* isomerization about the carbonyl group.⁴ The tetramer should then reproduce much of the local bonding behavior found in a single polymer chain, and cyclic oligomers in general are useful intermediates for the preparation of high molecular weight polycarbonate.⁵ Ring-opening polymerization, ROP,⁶ is often initiated by nucleophilic attack, giving rise to a “living” polymer.⁷

The present work describes density functional calculations of the reaction of the cyclic tetramer with the molecules phenol, lithium phenoxide (LiOPh), and sodium phenoxide (NaOPh). NaOPh has often been used as a reactant, one of the best known examples being its carboxylation to salicylic acid (Kolbe–Schmitt reaction),⁸ and it initiates polymerization in cyclic carbonates.⁶ The crystallization of PC is increased by several orders of magnitude by the presence of alkali metal aryl

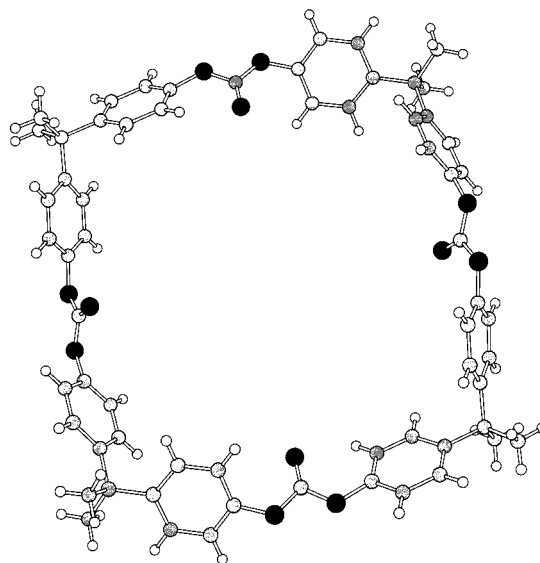


Figure 1. Cyclic tetramer of BPA-PC: C atoms, gray; O, black; H, white.

carbonates such as sodium benzoate, and this has been attributed to the formation of NaOPh chain ends that catalyze a fast transesterification reaction.⁹ The polymerization of macrocyclic oligomers of BPA-PC has been initiated by numerous ions, and the reaction of LiOPh with the cyclic tetramer is unusual, since it has no measurable exotherm (enthalpy less than 0.3 kcal/mol)¹⁰ and must be entropy driven. Experimental studies on a number of functionalized bisphenols using a variety of catalysts showed that the carbonate group provided the site for ring-opening polymerization in all cases.¹¹

These calculations continue a series of applications of the density functional (DF) method¹² to organic molecules, molecular crystals, and polymers, including the cyclic tetramer.¹³ The DF calculations are free of adjustable parameters and, when combined with molecular dynamics (MD),¹⁴ allow us to simulate the microscopic dynamical behavior of systems and to avoid geometrical structures corresponding to unfavorable local

* Corresponding author (e-mail: r.jones@fz-juelich.de).

[†] Permanent address: Università degli Studi di Messina, Dipartimento di Fisica, Contrada Papardo, I-98166 Messina, Italy.

[‡] Present address: Department of Physics, Queen's University, Belfast BT7 1NN, Northern Ireland.

minima in the energy surface. Our earlier work^{2,15} has showed that DF calculations—with the exchange-correlation energy functional proposed by Perdew, Burke, and Ernzerhof¹⁶—provide a satisfactory description of the intermolecular interactions in crystalline forms of polyethylene (PE) and DPBC. Rotation barriers in small organic molecules are described well by both local density and gradient-corrected functionals,¹⁷ although both tend to underestimate reaction barriers.¹⁸ Nevertheless, DF calculations provide the only viable method at present to identify reaction mechanisms in systems as large and as complex as those studied here, to account for the atomic relaxations during the reaction, and to provide estimates of the energy barriers. Moreover, the structural rearrangements involved are relatively minor, and only the intermediate stages display species with a different chemical configuration. Essential details of the calculations are provided in section II. The properties of the isolated phenol, LiOPh, and NaOPh molecules are discussed in section III, and the reactions of these three molecules with the cyclic tetramer are described in section IV. The vibration frequencies found for various structural units are presented in section V. Our findings are summarized in section VI, where we also discuss the balance between energy and entropy during the reactions.

2. Method of Calculation

The method of calculation was described in detail previously.^{19,20} The electron–ion interaction is represented by ionic pseudopotentials with the (nonlocal) form suggested by Troullier and Martins.²¹ We adopt a plane wave basis with a kinetic energy cutoff of 35 au, and the expansion of the orbitals uses a single point ($k = 0$) in the Brillouin zone. Periodic boundary conditions are assumed, and we use an orthorhombic cell with lattice constants 58.6, 41.0, and 26.5 Å. Density functional calculations presently require an approximation to the exchange–correlation energy. Our previous calculations on organic molecular crystals¹⁵ indicated that the PBE approximation, which is relatively simple and free of empirical input, is suitable in this context. It is used in the present work. All structures for the reactants and products were optimized using a combination of DF calculations with MD,²² and the coordinates of *all* atoms are allowed to relax. The vibration frequencies and eigenvectors of the molecules were calculated for the optimized geometries discussed in section V. The dynamical matrix **D** is evaluated using finite differences with atomic displacements δ of the order of 0.005 Å. The computed frequencies change by at most a few cm^{-1} upon increasing or decreasing δ by a factor of 3, so that anharmonic contributions are very small. The intensity of infrared fundamentals in absorption α for a mode with frequency ν_j is estimated by associating an electrostatic charge q_I with each atom I and evaluating

$$\alpha(\nu_j) = \frac{8\pi^3 N \nu_j}{3ch} \left| \sum_I q_I \delta \mathbf{r}_I^{(j)} \right|^2 \quad (1)$$

where $\delta \mathbf{r}_I^{(j)}$ is the displacement of atom I for the j th vibrational eigenvector and N is the molecular density. The atomic charges q_I are computed by a least-squares fit to the electrostatic potential around each molecule.²³

3. Isolated Phenol, LiOPh, and NaOPh Molecules

The reactions studied here are initiated by the molecules shown in Figure 2: (a) phenol, (b) LiOPh, and (c) NaOPh. Some properties of the isolated molecules that could influence their

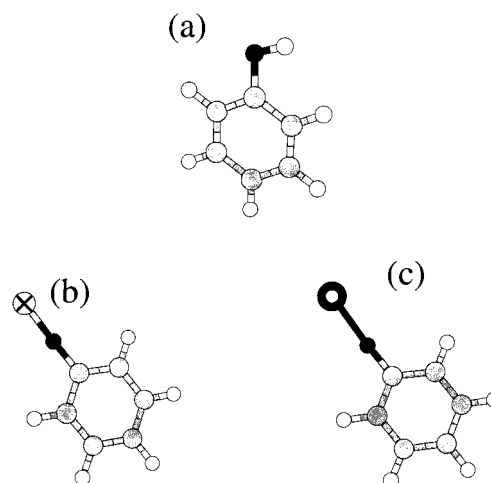


Figure 2. Molecules studied in the present work: (a) LiOPh; (b) NaOPh; (c) phenol. Key: C atoms, gray; O, black; H, white; Li, crosses; Na, double circles.

TABLE 1: Calculated Properties for Molecules $\text{C}_6\text{H}_5\text{OM}$ ($\text{M} = \text{H, Li, Na}$)^a

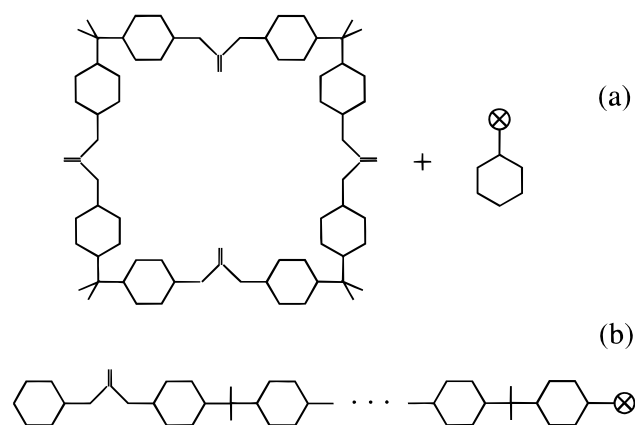
| | M = H | M = Li | M = Na |
|--|-------|--------|--------|
| dipole moment (D) | 1.3 | 7.6 | 12.0 |
| $q(\text{M})$ (e) | +0.35 | +0.83 | +0.91 |
| $q(\text{O})$ (e) | −0.53 | −1.10 | −1.11 |
| $q(\text{C})$ (e) | +0.40 | +0.82 | +0.84 |
| $d(\text{M}–\text{O})$ (Å) | 0.98 | 1.56 | 2.11 |
| $\alpha(\text{C}–\text{O}–\text{M})$ (deg) | 108 | 180 | 180 |
| $\Delta E(\text{H}_2\text{O})$ (kcal/mol) | 6.6 | 16.5 | 10.3 |

^a $\Delta E(\text{H}_2\text{O})$ is the cohesive energy of the complex with a water molecule bound to the terminal atom M.

interaction with polycarbonate chains, particularly the structure and charge distribution, are given in Table 1. LiOPh and NaOPh are rather similar, each having a linear $–\text{C}–\text{O}–\text{M}$ ($\text{M} = \text{Li, Na}$) termination and an $\text{O}–\text{M}$ bond that is predominantly ionic (as revealed by the analysis of the atomic charges and the Mayer bond order²⁴). The dipole moments are large and oriented along the linear termination. In phenol, by contrast, the $\text{O}–\text{H}$ termination has a strong covalent component (the stretching frequency of 3530 cm^{-1} is typical of the hydroxyl group), the $–\text{C}–\text{O}–\text{H}$ angle is bent, both the C and O atoms have a small electrostatic charge, and the dipole moment is much less. As a measure of the ability of these molecules to bind to closed-shell molecules, we have computed their binding energy with a single water molecule. The bond where the oxygen in H_2O is the acceptor for the $–\text{M}$ termination of phenoxide is strongest in LiOPh. The binding energy of the hydrogen bond in phenol (6.6 kcal/mol) is in good agreement with previous theoretical estimates.²⁵ The present calculations do not allow us to make an unambiguous separation into covalent and ionic components. Nevertheless, all these bonds (particularly water–LiOPh) have a covalent component, since the trends cannot be accounted for by the computed atomic charges and standard values of the ionic radii.

4. Reactions with Phenol and Phenoxides

Computer simulations cannot sample all possible reaction paths, and the most favorable found may depend on the initial geometry, the number of points sampled on the path, and the accuracy of the local relaxation procedure. Such effects have been reduced by performing two simulations on each system, with the initiating molecule *outside* and *inside* the tetramer ring, respectively, and not in its average plane. The initial reaction

SCHEME 1: (a) Reactants and (b) Products of Ring-Opening of the Cyclic Tetramer by LiOPh

coordinate is the distance R_C between the C atom of a carbonate group of the ring and the O atom on phenol or phenoxide. R_C is reduced from values >5 Å, where the interaction is very weak, in progressively shorter steps (~ 0.5 , 0.4 , or 0.2 Å) until a structural transformation or a large energy increase occurs. For each value of R_C , we relax *all* other degrees of freedom of the structure until the energy gradients along the unconstrained directions are less than 10^{-5} au. There are qualitative differences between the results for phenol and those for LiOPh and NaOPh, and we provide full details below. The total potential energy E_{tot} in phenol increases rapidly as R_C is decreased. We have discontinued the simulation when ΔE_{tot} reached 40 kcal/mol, without any significant chemical rearrangement occurring. On the other hand, a reaction takes place in both phenoxides that involves small energy barriers and a nontrivial sequence of characteristic steps. The first stage of all reactions is the formation of a bridge between M and the O of the carbonyl group. The reduction in E_{tot} favors a further approach of the reactant molecule to the cyclic tetramer and allows significant changes in the geometry (particularly in the torsion and bending angles) that allow the reaction to proceed. The change from sp^2 to sp^3 of the electronic configuration of the carbonate C atom is crucial, as the compact surroundings of negatively charged O atoms facilitate the passage of the alkali metal ion around the reaction center. The final state is an open structure with a weak bond between the metal and the new carbonyl group. Thermal effects and the interaction with other carbonate molecules should break this bond readily above room temperature, leading to the final configuration shown in Scheme 1b.

4.1. Lithium Phenoxide. Figure 3 shows six snapshots of the trajectory for the reaction of LiOPh initially outside the cyclic tetramer, and the corresponding energy changes as a function of R_C are shown in Figure 4a. The two molecules interact very weakly for $R_C \sim 4$ Å, with minor atomic relaxations and energy changes of less than 1 kcal/mol, and steric forces give rise to an energy barrier of 4 kcal/mol at $R_C = 3.8$ Å. For shorter separations the Li–O attraction produces a rotation of the carbonyl group, the formation of a bond similar to that in the LiOPh–water complex, and a rapid decrease (~ 15 kcal/mol) in E_{tot} . The energy stays below the initial value as R_C is reduced to ~ 1.8 Å, where the C of the original carbonate is surrounded by four O atoms with approximately tetrahedral coordination (C_T).

The formation of this configuration (3(3)) is accompanied by an increase in all C_T –O distances, since the electrophilic Li atom is shared between the original O of LiOPh and the O atom formerly in the carbonyl group. C_T has a large positive

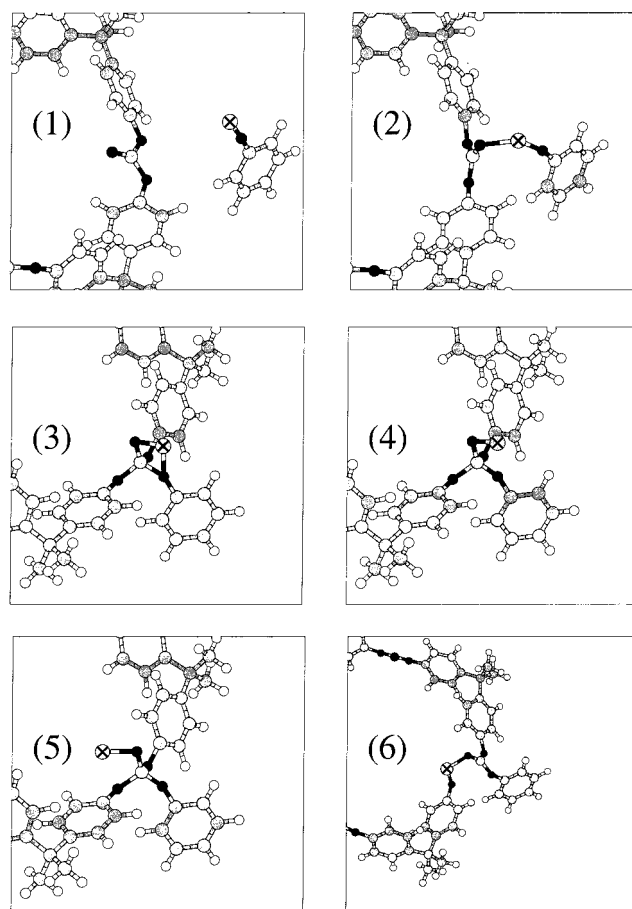


Figure 3. Reaction of LiOPh with cyclic tetramer of BPA-PC. Only part of the ring is shown.

charge, while the four O atoms are nearly equivalent, with charges similar to those of other tetrahedral O atoms in the chain. A reduction of R_C to 1.5 Å (corresponding to a second energy barrier of ~ 4 kcal/mol) is required to symmetrize the tetrahedral geometry of C_T and to break the symmetric bonding of Li. The Li is now attached to the tetrahedral O that is bound to a single C atom (3(4)). The Li–O bond is oriented mainly along one of the other three C_T –O bonds, probably because of the electrostatic attraction between the positive Li and the electron charge in the covalent bonds. Nevertheless, the C_T –O bond provides a 3-fold axis for rotations of the Li–O bond that change only the O– C_T –O–Li torsion angle and give rise to variations of E_{tot} of at most ~ 1 kcal/mol. Under normal thermodynamic conditions, the O–Li bond can then be oriented with equal probability along the three directions specified by the C_T –O bonds at the base of the tetrahedron.

Relaxation of the two equivalent structures with the Li–O bond oriented along a C_T –O bond different from the original one gives rise to structure \sim (6), which has a weak bond between Li and the new carbonyl O. This geometry is locally stable, with an energy ~ 6 kcal/mol lower than the combined energy of the reactants. The difference in energies between this metastable geometry and 3(2), which is an additional 6 kcal/mol more stable and has a *cis-cis* configuration of the carbonate group in 3(6). At room temperature, the weak bond between Li and O should break easily, leading to the open chain with an active $-C=O-Li$ termination.

The approach of LiOPh from *inside* the tetramer ring yields the same product, but the details of the reaction mechanism

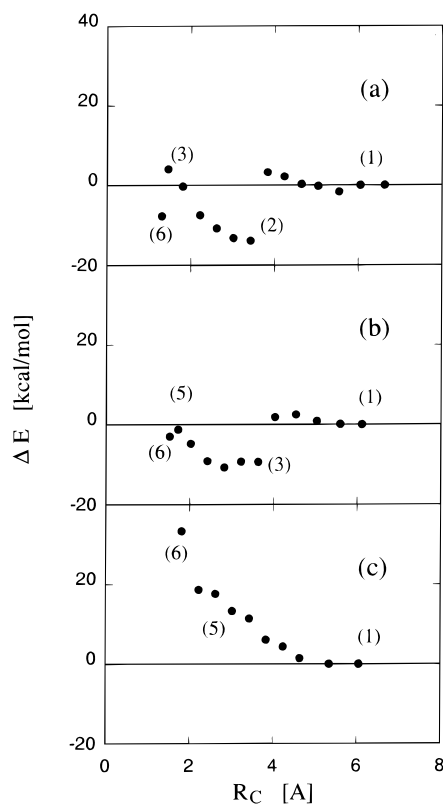


Figure 4. Variation of total potential energy with reaction coordinate R_C during reactions of (a) LiOPh, (b) NaOPh, and (c) phenol. $\Delta E = 0$ corresponds to the sum of the energies of the isolated molecules. The numbers denote the related structures in Figures 3, 5, and 6, respectively.

differ. The interaction of the molecules is very weak for $R_C > 4$ Å, but the formation of a bond between Li and the two tetrahedral O in the carbonate group ($R_C \sim 3.8$ Å) is accompanied by a lowering of E_{tot} of ~ 12 kcal/mol, similar to the first attractive well in the other trajectory. The bonds of Li to the three O atoms are of unequal strengths down to $R_C = 2.1$ Å, where the carbonate C acquires a nearly tetrahedral coordination and an sp^3 electronic configuration. At this point, the Li atom is at the center of the base of the tetrahedron surrounding C_T . A reduction of R_C to 1.61 Å leads to an increase in E_{tot} to 6 kcal above the energy of the reactant molecules, and the tetrahedral coordination of C_T becomes deformed. Further reducing R_C causes one of the O- C_T bonds of the original carbonate group to lengthen, without a further increase in the energy. This leads to a metastable species similar to 3(6). E_{tot} is again determined by the configuration of the new carbonate group, and the deformed *trans-trans* configuration results in a higher energy than in 3(6). Many other trajectories are possible for this reaction, but these two examples show that ring-opening can occur without crossing significant energy barriers.

4.2. Sodium Phenoxide. The similarity of LiOPh and NaOPh is reflected in their behavior during reaction with the cyclic tetramer. Six snapshots illustrating the first of the two simulations are shown in Figure 5, and the corresponding energy variations are shown in Figure 4b. The longer -O-Na bond makes it easier for Na to approach the carbonate (we find no energy barrier) and to form (at $R_C = 3.6$ Å) a bridge between the incoming molecule and the carbonyl O. E_{tot} is lowered by 11 kcal/mol, a similar value to that found in LiOPh. The smaller rotation of the carbonyl required to bond with the incoming Na appears to compensate for the relative weakness of the bond. The energy remains nearly constant, and the structure remains

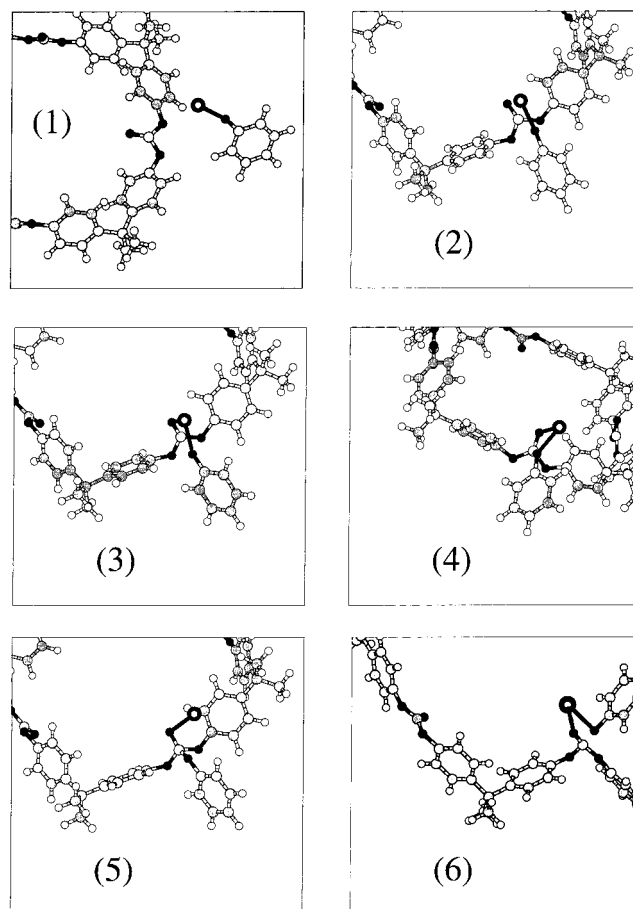


Figure 5. Reaction of NaOPh with cyclic tetramer of BPA-PC. Only part of the ring is shown.

similar (with Na shared nearly symmetrically between two O atoms) on decreasing R_C to 2.01 Å. An increase in energy is required to promote the carbonate C to sp^3 coordination, but E_{tot} remains below the sum of the energies of the isolated reactants. The original Na-O bond is broken at $R_C = 1.5$ Å, and Na remains bound only to the O previously forming the tip of carbonyl. This bond does not align along one of the other C_T -O bonds, and the equilibrium value of the O- C_T -O-Na torsion angle changes from 0 to 41°. This difference from the results for LiOPh appears to be a consequence of the increased size of the metal ion. The related steric hindrance due to the presence of the phenyl rings means that the rotation of the O-Na bond around the 3-fold C_T -O axis requires more energy (~ 4 kcal/mol) than in LiOPh. On the other hand, one of the original C-O single bonds opens when the Na ion approaches it upon a rotation of 120° from the original direction, as found in LiOPh. Figure 4 shows that E_{tot} does not lie above its initial value at any stage of the reaction. The trajectory initiating with NaOPh *inside* the tetramer ring shows somewhat larger differences from the corresponding simulation for LiOPh, with the longest Na-O bond being less favored in forming a bond with the carbonate O atoms. While R_C decreases to 2.6 Å, the C-O-Na termination of NaOPh bends by nearly 60° to maintain the separation of Na from the (positively charged) central C atom above ~ 4 Å. A bond between Na and the two tetrahedral O atoms of the original carbonate appears to be possible only after this C atom has tetrahedral coordination at $R_C = 2$ Å. An additional difference between Li and Na is that the latter prefers attachment to a single O atom, although our simulations show that Na moves from one O to another unpredictably as R_C changes. Despite these differences, presum-

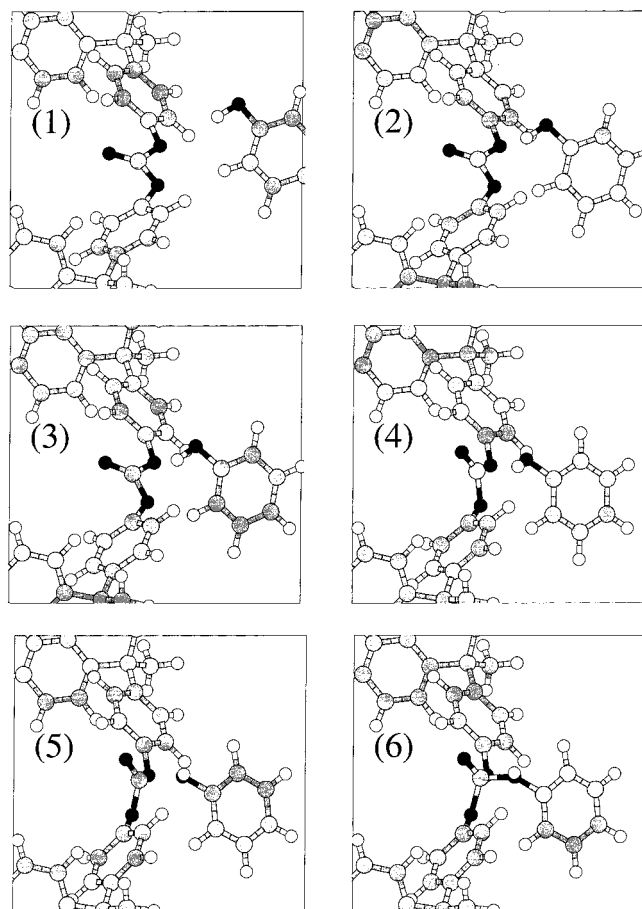


Figure 6. Reaction of phenol with cyclic tetramer of BPA-PC. Only part of the ring is shown.

ably related to the different sizes of Li^+ and Na^+ , the structures resulting when R_C is less than 1.5 Å are very similar.

4.3. Phenol. We have noted that the approach of phenol to a carbonate group in the cyclic tetramer results in a purely repulsive potential energy curve, and the energy barrier (>40 kcal/mol, Figure 4c) is too large for practical reactions. However, the simulation trajectories (Figure 6) show that the reduction of R_C leads to structural changes similar to those found in LiOPh and NaOPh. The change in E_{tot} is initially small, the carbonyl group eventually rotates toward the terminal atom of phenol, the central C atom acquires a tetrahedral coordination, and the nearest phenyl rings rotate to accommodate the incoming molecule. However, we do *not* observe bond formation between the termination of phenol and the carbonate O atoms, and there is no potential energy well that allows the molecules to approach and to change the roles of the C_T —O bonds. Differences in phenol are evident in its structure (the bent —C—O—H bond hinders the approach toward the carbonate O atoms) and other bonding properties evident in the weak hydrogen bond with water. The same result is also found when the phenol molecule approaches the ring from *inside*.

5. Vibrational Properties of Metastable Species

The reactions of LiOPh and NaOPh with the cyclic tetramer take place as a well-defined sequence between transient species whose chemical properties (e.g., the tetrahedral coordination of the carbonyl C) differ from those of both the reactants and product. To aid the identification of these species in future experiments, we have analyzed the vibrations of metastable structures obtained by relaxing structures associated with

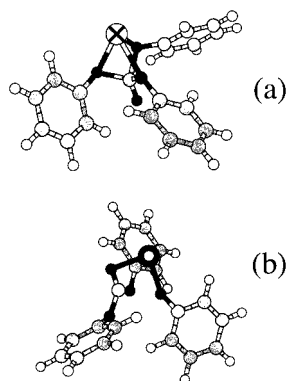


Figure 7. Metastable structures for which vibration frequencies have been determined: (a) structure observed in reaction of LiOPh approaching from *inside* the ring; (b) structure corresponding to 5(3).

shallow potential energy wells along the reaction path. All such structures relax to metastable systems that retain the features of the starting configuration and are stable against dissociation by at least 6 kcal/mol.

To optimize the use of computer resources, we have isolated the active part of the tetramer by (a) cutting the ring beyond the two phenyl rings nearest to the reacting carbonate and (b) saturating the dangling bonds with H atoms. Our previous computations for a series of carbonate systems¹³ have shown that vibrational and chemical properties are remarkably local and transferable within this family of molecules. Moreover, we describe below a very large computation of the vibrational properties of the entire tetramer that demonstrates this transferability for the metastable species as well. The vibrational properties of two metastable molecules (Figure 7) are determined by the method used in our earlier work on carbonates¹³ and outlined in section VI.

The intermediate species reflect the constraints used in the simulations and differ somewhat from the relaxed structures for which we calculate the vibration frequencies, e.g., the tetrahedral C atom always relaxes to a highly deformed sp^2 configuration. Nevertheless, the significant changes produced by Li^+ or Na^+ in some vibrational modes should aid the identification of the intermediate state. This should be possible if the lifetime of these species is long enough in the reaction environment or by using more initiating molecules or short-time (femtosecond) spectroscopic techniques.

The modes localized on the carbonate group are the most affected by the formation of the complexes, and their ionic character and infrared activity should make them easy to detect. The most obvious differences in all cases arise from the mixing of modes belonging to different symmetries in the isolated molecules. For instance, “in plane” and “out of plane” bending modes for the carbonyl are no longer separated unambiguously, and mixing of bending and stretching modes is observed for the single C—O bonds.

In the structure of Figure 7a, the C=O stretching mode has a frequency of 1740 cm^{-1} , compared with 1780 cm^{-1} for the isolated cyclic tetramer. Due in part to mixing with other modes, frequency changes in the C=O bending modes are larger, going from the 1139 cm^{-1} computed for the ring to the 910 cm^{-1} for the fragment of Figure 7a. Moreover, two new frequencies appear at 533 and 422 cm^{-1} (stretching and bending of the O—Li bonds, respectively) in a spectral region where the isolated tetramer has very low infrared activity. The carbonate structure is less affected by the approaching molecules in Figure 7b, and the stretching mode of C=O (1750 cm^{-1}) is closer to the original value. The mode with the strongest C=O bending component

(898 cm^{-1}) results from a mixture of the in-plane (1139 cm^{-1}) and out-of-plane (725 cm^{-1}) modes of the cyclic tetramer. The stretching and bending of the Na–O bonds are also mixed, occurring in a frequency range (320–355 cm^{-1}) intermediate between the bending (278 cm^{-1}) and stretching (565 cm^{-1}) of the same bond in NaOPh.

To verify that the results obtained with the fragments are representative of those for larger carbonate molecules, we have repeated the computation for the Na fragment by considering the entire cyclic tetramer adjacent to NaOPh. The differences in the frequencies of the modes discussed above are small (at most 15 cm^{-1}) and show that the use of a truncated molecule is a reliable approximation.

6. Discussion and Concluding Remarks

We have performed density functional calculations on the reactions of the cyclic tetramer of BPA-PC and three molecules (LiOPh, NaOPh, and phenol). Reactions with BPA-PC are known to take place at the carbonate groups,⁵ and simulations have been performed for each molecule approaching this group from *inside* and *outside* the ring. The calculations are performed by choosing an appropriate reaction coordinate and optimizing the positions of *all* other atomic positions. The total energy of the system and the forces and charges on all atoms are monitored throughout the simulations. The approach of the molecules toward the carbonate group results in initial structural changes that are common to all. The carbonyl O in the tetramer rotates toward the terminal atom of the approaching molecule, resulting in a change in the coordination of the carbonyl C from sp^2 to sp^3 . However, the subsequent reaction path and the changes in the total potential energy are very different in phenol, on one hand, and the alkali metal phenoxides, on the other. The energy increases rapidly in phenol, so that a reaction will not take place under normal conditions, while the phenoxides lead to reactions with low energy barriers that result in ring opening. The result is a chain with a phenyl carbonate at one end and a phenoxide at the other (Scheme 1b). The structure and charges of the –C–O–M termination in the phenoxides and in the extended molecules are very similar, so that the “active” termination has been transferred from one molecule to another. Ring-opening may then be viewed as being catalyzed by the phenoxide and giving rise to a “living” polymer.

The driving force toward polymerization can be provided by potential energy or entropy. We have studied this balance by considering the reaction shown in Scheme 1 and computing the total energy of both the separated reactant molecules and the final structure. This reaction does not change the number and type of bonds, and the change in E_{tot} is very small ($\Delta E_{\text{tot}} = +2$, $+1$, and $+1$ kcal/mol for phenol, LiOPh, and NaOPh, respectively). Although the energy differences are negligible in all cases, the energy *barriers* are much higher in phenol. The calculations indicate that enthalpy cannot be the sole driving force for polymerization in the phenoxides, and the absence of an exotherm agrees with experimental findings.¹⁰ Entropy changes are likely to play a major role and could be studied further by measuring the temperature dependence of the reaction rate. The computing demands of the DF-based calculations described above rule out their application to long simulations over a range of temperatures, but we are planning an extensive study using a classical force field.²⁶ First results on a single

ring system (tetramer plus LiOPh) are encouraging. The ring system is very mobile at room temperature and above, and the clear preference for the LiOPh group to be *inside* the ring is consistent with the increased number of van der Waals bonds in this geometry. These first calculations indicate that there is a small entropy difference between the systems a and b in Scheme 1. However, the results for a single ring do not represent the interactions in a melt, for example, where the “living” Li atom can interact with many other molecules. We are extending the classical simulations to cover this case.

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