**Atropisomerism in Tertiary Biaryl 2-Amides: A Study of Ar-CO and Ar-Ar**′ **Rotational Barriers**

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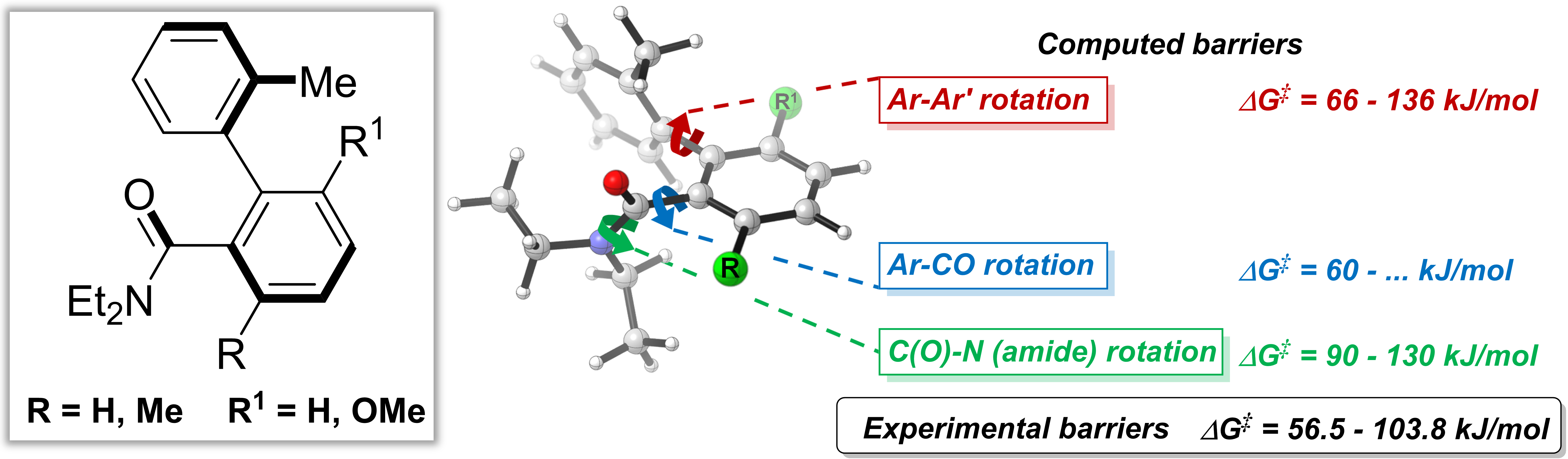
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**Abstract:**

A rotational barrier study was performed on eight tertiary biaryl 2-amides using variable temperature (VT) NMR and exchange (EXSY) spectroscopy experiments. Seven out of the eight 2-amido-2′-methylbiphenyls with additional 3- and 6-substitution patterns (**1-7**) were found to have approximately similar rotational barriers (G‡Tc= 56.5 – 67.5 kJ/mol). However, for both 3- and 6-substitution (**8**), the rotational barrier was found to be significantly higher (G‡ = 102.6-103.8 kJ/mol). Computational studies performed on all eight compounds gave results in good agreement with the experimental rotational barriers. A transition state in which atropisomerism occurs by a cooperative rotation of the Ar-CO and Ar-Ar′ bonds depending on substituent location is proposed.

**Graphical Abstract:**



**INTRODUCTION**

Atropisomeric molecules, discovered by Christie and Kenner[[1]](#endnote-1) in 1922, exhibit restricted rotation (torsion) around a single bond due to excessive nonbonded interactions.[[2]](#endnote-2) Atropisomerism is predominantly associated with biaryls[[3]](#endnote-3) and binaphthyls,[[4]](#endnote-4) in which pivotal bond rotation is prevented by bulky substituents thus allowing the isolation of enantiomeric forms. This type of molecular chirality has had considerable significance in the development of chiral auxiliaries and ligands for asymmetric synthesis.[[5]](#endnote-5) Furthermore, since many biologically active biaryl compounds possess chirality due to restricted rotation,[[6]](#endnote-6) atropisomerism has become important in drug discovery endeavors.[[7]](#endnote-7) Tertiary aryl amides and anilides, known to display atropisomerism,[[8]](#endnote-8) allow generation of atropisomers directly[[9]](#endnote-9) or by dynamic kinetic resolution.[[10]](#endnote-10) These compounds have also been used as effective chiral ligands[[11]](#endnote-11) and auxiliaries.[[12]](#endnote-12) In particular, aryl amides, well known to have high rotational barriers due to the partial double bond character of the C-N bond arising from amide bond resonance,[[13]](#endnote-13) have been shown to exhibit rotational barriers from cooperative C-N/Ar-CO rotations by the elegant and extensive investigations of Clayden and coworkers.[[14]](#endnote-14)

In the course of recent studies, we prepared a series of 2-amido-2ʹ-methyl biaryls (Figure 1) for the development of a general route to phenanthrenes[[15]](#endnote-15),[[16]](#endnote-16) and noted that this biaryl structure type has three bonds which may give rise to high rotational barriers: the CO-N (amide) bond (green), the Ar-CO bond (blue) and the Ar-Arʹ (biaryl) bond (red) and hence may be amenable to a rotational barrier investigation by VT NMR spectroscopy.



Figure 1. Potential rotational barriers of 2-amido-2ʹ-methylbiaryls.

Herein we report a rotational barrier study of a series of substituted 2-amido-2ʹ-methylbiaryls (**1**, **4**-**8**), a 2-(2ʹ-methylphenyl) naphthamide (**2**) and a 2-(1ʹ-methyl-2ʹ-naphthyl) benzamide (**3**) (Figure 2). Thus, three substrates with no other substituents but with variation of phenyl to naphthalene rings (**1**-**3**), three biaryl 2-amides with additional substituents, 6**,** 3ʹ- (**4**), 6- (**5**), 6,5ʹ- (**6**) and 3,5- (**7**), and one biaryl with a 3,6,2ʹ-substitution pattern (**8**) were investigated. Based on previous studies concerning the rotational barriers of substituted biaryls,3 observation of high Ar-Arʹ rotational barriers was expected for compounds **4**-**6** and **8**.



Figure 2. 2-Amidobiaryls in the present rotational barrier study. IUPAC-numbering is given. Biaryls 4-8 follow the numbering of biaryl 1.

In the course of studies concerned with the preparation of chrysenols (Scheme 1), we observed that treatment of 2-amido-2ʹ-methylbiphenyl **2** under excess LDA conditions resulted in directed remote metalation (DreM) – cyclization to afford 5-chrysenol **9** (Scheme 1).15d,16 However, in some cases,15d,16 fluorenone derivatives were formed, e.g. the conversion of **3** intofluorenone **10** instead of the 6-chrysenol **11**. The 1H NMR spectra of biaryls **2** and **3** displayed evidence for the presence of two rotamers/diastereomers strongly suggesting, as one rationalization, a high kinetic barrier due to hindered rotation about the biaryl bond which prevents attainment of the necessary transition state for the DreM-reaction.[[17]](#endnote-17) As a precedent for this supposition, we have previously observed a case where, of two isolated atropisomers, only one underwent a DreM reaction.[[18]](#endnote-18) These results enhanced further our interest in the present investigation.

**Scheme 1**. Selectivities of the DreM-cyclization of *ortho*-aryl naphthamide **2** and benzamide **3**. Conditions: a) LDA (2.5 equiv), THF, r.t.

**RESULTS AND DISCUSION**

All compounds were prepared either by the directed *ortho*-metalation (DoM)-boronation reactions of *N*,*N*-diethyl benzamides, followed by Suzuki-Miyaura cross coupling reactions of the resulting boronic acids with 2-bromotoluenes, or by reaction of the reversed coupling partners.16 For the VT 1H NMR experiments, all compounds, with the exception of compound **2** (23 °C, 298 K) and **3** (-83 °C, 190 K), were cooled to 250 K before 1H NMR, NOESY and COSY measurements were recorded. At these low temperatures, the maximum chemical shift differences (Δυ in Hz) between the two exchanging 2ʹ-methyl-peaks were obtained. The rate constant, *kc* of the interconversion of the two exchanging rotamers at coalescence temperature, Tc, was estimated by the Gutowsky-Holm equation.[[19]](#endnote-19) From the coalescence temperature, Tc, and the rate constant, *k*c, the activation energy of rotation (ΔG‡Tc) was calculated using the Eyring equation.[[20]](#endnote-20) (see SI).

For compounds **2** and **5** as illustrative cases, the VT 1H NMR data are depicted in Figure 3. At low temperature (298 K and 250 K for compounds **2** and **5**, respectively) the two exchanging 2ʹ-methyl signals are observed as two singlets due to a slow exchange process. Raising the temperature resulted in the expected increased rate of exchange of the two rotamers, observed broadening of the two signals and eventual coalescence. For substrate **2**, coalescence occurred at 332 K, while for substrate **5** at 314 K, which resulted in calculated rotational energy barriers of ΔG‡Tc = 67.5 and ΔG‡Tc = 63.0 kJ/mol, respectively. The results of the VT 1H NMR studies of the 2-amido-2ʹ-methylbiaryls **1-7** are tabulated in Table 1. The 3-Me-6-OMe biaryl amide **8** had a very high energy barrier, and we were unable to approach the coalescence temperature, even up to 380 K[[21]](#endnote-21) in DMSO-*d6*.

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Figure 3. VT 1H NMR of compounds 2 and 5.

**Table 1.** Rotational barriers of compounds 1-**7** from VT-NMR measurements.a,b

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Entry** | **Compound** | **δ**  **(ppm)** | **Δ*vc***  **(Hz)** | ***T*c,**  **°C (K)** | ***k*cd**  **(s-1)** | **ΔG‡Tce**  **(kJ/mol)** |
| **1** | **1** | 2.35, 2.08 | 106.9 | 11 (284) | 237 | 56.5 |
| **2** | **2** | 2.30, 2.12 | 74.4 | 59 (332) | 165 | 67.5 |
| **3** | **3** | 2.46, 2.35 | 42.8 | 10 (283) | 95 | 58.5 |
| **4** | **4** | 2.26, 2.04 | 89.4 | 40 (313) | 198 | 63.0 |
| **5** | **5** | 2.39, 2.15 | 96.6 | 41 (314) | 214 | 63.0 |
| **6** | **6** | 2.24, 2.02 | 88.8 | 28 (301) | 197 | 60.5 |
| **7** | **7** | 2.43, 2.13 | 117.1 | 52 (325) | 260 | 64.8 |

a 400 MHz NMR; b Lowest temperature reached: -83 °C (190 K); cΔ*v* obtained from the methyl peak in exchange at low temperature; d *k*c = 2.22Δ*v*. e Estimated margin of error ± 0.8 kJ/mol.



Figure 4. Diastereotopic and enantiotopic rotamers of biaryl amide 8 resulting from Ar-Arʹ and Ar-CO rotations.

To interpret the VT-NMR data, one must consider that the possibility of both Ar-CO and Ar-Arʹ rotations leads to enantiotopic and diastereotopic rotamers (Figure 4).14 Rotamers due to only Ar-CO rotation or Ar-Arʹ rotation will, due to the diastereotopic rotamer properties, show different distinct signals between atropdiastereomers **A** and **B** in the 1H NMR spectra which thereby also allows an EXSY-NMR study.[[22]](#endnote-22) The **A-B** atropdiastereomeric exchange study of compound **8**, for which a coalescence temperature was unattainable, was carried out in toluene-*d8* at 380 K with mixing times from 1 to 0.01 sec. The rotational barrier was calculated from the volume integrals of the exchange peaks (entry 9, Table 2). (For a complete discussion of these experiments, see SI, Section 2). To verify that these results could be compared with the variable temperature experiments, the EXSY-NMR experiments were also conducted on compound **5** in acetone-*d6* at 273 K with mixing times from 1 to 0.003 sec (entry 6, Table 2). That the rotational energy barrier is almost insensitive to solvents was demonstrated for compound **5** by experiments in toluene-*d8* and CDCl3 which gave the same energy barrier within 1.4 kJ/mol. Atropdiastereomers **A** and **B** of compound **8** were separated by standard flash chromatography. The structure of atropdiastereomer **B** (Figure 4) was assigned by observation of an NOE between the 2ʹ-methyl and the amide methylene protons (SI, Section 2.6). Its isomerization to atropdiastereomer **A** at 232 K (see SI, Section 2.4) was studied by 1H NMR and showed an energy barrier (entry 10, Table 2) which was in good agreement with that measured by EXSY-NMR.

As expected, biaryls without amide- and biaryl axis-obstructing 3- and 6-substituents, **1** and **3** respectively (entries 1 and 3, Table 1) showed the lowest rotational barriers. The 6-OMe bearing biaryls **4, 5** and **6** showed somewhat higher activation energies (entry 4, 5 and 6), but less enlarged than expected by a hindered biaryl rotation. Of the methods for assessing the steric bulk of substituents,[[23]](#endnote-23) we compared our results with those for structurally related biaryl systems. In biaryl rotations,27 the methoxy group has a smaller effective van der Waals radii (1.52 Å) than the methyl group (1.80 Å), but should still give a substantial increase in the rotational barrier as observed in the comparison of biaryls **12** with **13** (Figure 5a).[[24]](#endnote-24) On the other hand, benzamide **16** (Figure 5c), which closely resembles **4-6**,shows a ΔG‡Tc = 63.2 kJ/mol that must be attributed to the Ar-CO rotational barrier in view of the symmetry of the other aryl ring.[[25]](#endnote-25) Consequently, the **A-B** diastereoisomer interconversions for biaryls **4**, **5** and **6** occur through Ar-CO rotation. Furthermore, the additional 3ʹ-Me substituent (**4**)had little or no effect. For comparison, although electronically different to **4**, a similarly positioned 3ʹ-ethyl group in a 2,2ʹ-ditrifluoromethyl-biphenyl (not shown) provides an approximately 16 kJ/mol buttressing effect compared to the desethyl system.[[26]](#endnote-26)

The 5ʹ-F substituted biaryl **6,** showed a slightly lower activation energy than the non-fluorinated analogue **5** (compare entries 5 and 6). A higher rotational activation energy was found for naphthamide **2** (entry 2, ΔG‡Tc = 67.5 kJ/mol), which corresponds to a 11.0 kJ/mol buttressing effect of the *peri*-hydrogen (C-8) in **2** compared to biaryl **1**. For comparison with these results, in an investigation of benzene and naphthalene systems (Figure 5b), compound **15** was shown to have a ΔG‡ = 7.6 kJ/mol higher than that of compound **14**, which may be also attributed to a *peri*-hydrogen effect in the former compound.3a In our study, incorporation of a 3-methyl group as in **7** (entry 7, Table 1) results in a significantly higher activation energy (ΔΔG‡Tc= 8.3 kJ/mol) compared to that of the prototype **1** but slightly lower than that observed for the compound with a *peri*-hydrogen effect (**2**). Thus, these observations offer rationalization of both Ar-Ar’ and Ar-CO rotations. Compound **8** in which the Ar-CO rotation is hindered by the 3-Me substituent while the Ar-Arʹ rotation is hindered by the 6-OMe substituent shows a much higher rotational barrier than that observed for other compounds lacking this double hindrance effect.

Our results of very similar ΔG‡Tc values for the series of compounds **1-7** made it difficult to distinguish between the individual Ar-CO and Ar-Arʹ rotations in several studied cases. To shed more light on the dynamics of these molecules, a series of computational studies was performed.



**Figure 5.** Rotational barriers for relevant biaryls: a) effect of an OMe group on Ar-Ar rotation;24 b) *peri*-H effect;3a c) isolated Ar-CO rotation;25 d) isolated Ar-Ar rotation.[[27]](#endnote-27)

**Computational Studies**

A computational study at the CPCM (Toluene) M06L/6-311++G(d,p) // CPCM (Toluene) B97XD/6-31+G(d) level of theory was undertaken to further understand the experimentally measured barriers for compounds **1**-**8**.[[28]](#endnote-28) Five different bond rotations were considered: the biaryl bond (Ar-Arʹ), the aryl-carbonyl bond (Ar-CO), and the amide bond (Et2N-CO, via *syn* and *anti* transition states), and a combined rotation of the Ar-CO bond and the amide bond[[29]](#endnote-29) (Ar-CO / Et2N-CO; *anti* TS only). The corresponding transition state structures are depicted in Figure 6. All barriers are reported relative to the corresponding lowest energy atropisomer in kJ/mol. Table 2 shows the comparison of the results from the computational study with the corresponding experimentally determined values.

The DFT calculations suggest that for compounds **1, 2**, and **4-8**, the atropdiastereomer **B** is on average ΔGfƟ ≈ 1.3 kJ/mol lower in energy than **A**. (The exception, biaryl amide **3**, has atropdiastereomer **A** calculated to be ΔGfƟ = 3.3 kJ/mol more stable). These results concur with the EXSY-NMR experiments which show that the energy barriers of the **B-A** interconversions of **5** and **8** were slightly larger than those of the **A-B** interconversions (see SI, Sections 2.2, 2.3). Studies of energy differences between atropdiastereomers in biaryl amides have been previously reported by Clayden.[[30]](#endnote-30)

|  |  |  |
| --- | --- | --- |
| a) | b) | c) |
| **1** | **1** | **1** |
| d) | e) | f) |
| *Syn*-**1** | *Anti*-**2** | **1** |

**Figure 6.** Calculated transition states for the different bond rotations of biaryl amides **1** and **2.** a) Ar-Arʹ bond rotation, b) Ar-CO bond rotation, c) Ar-CO / Et2N-CO combined rotation with Arʹ and O eclipsed in the transition state, d) C=O – N bond rotation with *syn*-transition state, e) C=O – N bond rotation with *anti*-transition state, f) Ar-CO / Et2N – CO combined rotation with Arʹ and N eclipsed in the transition state.

**Table 2.** Calculateda and experimental (VT 1H NMR) rotational barriers in kJ/mol.

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **Entry** | **Compound** | **Measured**  **(ΔG‡Tc)** | **Ar-Ar′**  **(ΔG‡)** | **Ar-CO**  **(ΔG‡)** | **Et2N-CO**  **(*syn*)**  **(ΔG‡)** | **Et2N-CO**  **(*anti*)**  **(ΔG‡)** | **Ar-CO /**  **Et2N-CO**  **(ΔG‡)** |
| 1 | **1** | 56.5 | 66.2 | 62.1 | 115.1 | - | 91.0e |
| 2 | **2** | 67.5 | 68.0 | - | 124.3 | 105.2 | 109.5f |
| 3 | **3** | 58.5 | 81.4 | 63.2 | 90.3 | - | 91.3f |
| 4 | **4** | 63.0 | 135.6 | 68.9 | 112.7 | - | 93.1e,f |
| 5 | **5** | 63.0 | 126.8 | 68.6 | 112.8 | - | 91.8e,f |
| 6 | **5** | 63.6-64.5b |  |  |  |  |  |
| 7 | **6** | 60.5 | 123.0 | 60.5 | 109.0 | - | 84.5e |
| 8 | **7** | 64.8 | 69.4 | - | 130.8 | 104.0 | 95.5f |
| 9 | **8** | 102.6c | 126.7 | - | 122.0 (B)  130.2 (A) | 107.9 (B)  104.1 (A) | 108.1f |
| 10 | **8** | 103.8d |  |  |  |  |  |

a Calculations were performed at CPCM (Toluene) M06L/6-311++G(d,p) // CPCM (Toluene) B97XD/6-31+G(d) level of theory. b EXSY-NMR experiment in acetone-*d6* at 273 K (see SI, Section 2.3). c EXSY-NMR experiment in toluene-*d8* at 380 K (see SI, Section 2.2). d Isomerization of pure atropdiastereomer **8B** at 232 K (see SI, Section 2.4). e Arʹ – O transition state. f Arʹ – N transition state. “-“: The transition state could not be located.

The DFT calculations for the Ar-Arʹ rotation give ΔG‡ = 66.2-69.4 kJ/mol range for molecules without a 6-substituent (Table 2, compounds **1**, **2**, **7**). In the transition state of **1** (Figure 6a), the amide group is almost orthogonal to the biaryl plane and twisted away from the bond of rotation. This rotational barrier is comparable to that observed for biaryl **17** (Figure 5d) which bears only 2-methoxy and 2ʹ-methyl substituents (ΔG‡ =66.7 kJ/mol).27 For compound **3**, involving an aryl-naphthyl rotational barrier (entry 3, Table 2), a slight buttressing effect may be responsible for the higher energy barrier as discussed in the section on the VT-NMR experiments above. For compounds bearing 6-OMe substituents (**4**, **5**, **6**, **8**),the Ar-Arʹ rotation experiences considerable hindrance as calculated and observed in the increased energy barrier range of ΔG‡ =123.0-135.6 kJ/mol, comparable to that observed for compound **13** (Figure 5a).

The transition stateof **1** (Figure 6b) depicts that the Ar-CO rotation is facilitated by the presence of a twisted out of plane biaryl which provides space for amide carbonyl rotation without interference. Thus, for 3-unsubstituted biaryls **1**, **3**-**6**, calculations of the Ar-CO bond rotation gave a range ΔG‡ =60.5-68.9 kJ/mol, resembling the observed Ar-CO bond rotation barrier of **16** (Figure 5c). For naphthamide **2**, exhibiting a *peri*-H interaction, and biaryl amides **7** and **8** experiencing rotational hindrance from the 3-methyl group, it was not possible to locate a transition state for the Ar-CO rotation. Instead, a transition state was located which shows the NEt2 group rotated away from the otherwise favored planar amide structure. This, in turn, corresponds to a concerted Ar-CO / Et2N-CO rotation transition state (similar to those depicted in Figures 6c and 6f) which could also be located for all other structures. This concerted rotation was also visible in the EXSY-NMR experiments of **8** where a faster Et2N-CO rotation (ΔG‡EXSY = 98.0 kJ/mol)) (see SI, Section 2.2) was observed together with the concerted rotation (entry 9, Table 2) by the individual amide ethyl – 2ʹ -CH3 signals.

The concerted rotation may be represented by two transition states, with either the C=O or the Et2N group oriented towards the Arʹ-ring (i.e. Arʹ and O, or Arʹ and N eclipsed as depicted in Figure 6c and 6f respectively). Both transition states have very similar energies (G < 4 kJ/mol) and only compounds **2**, **6** and **8** exhibit preference for the Arʹ-N-type transition state, most likely due to increased steric bulk from the 6-substituent.

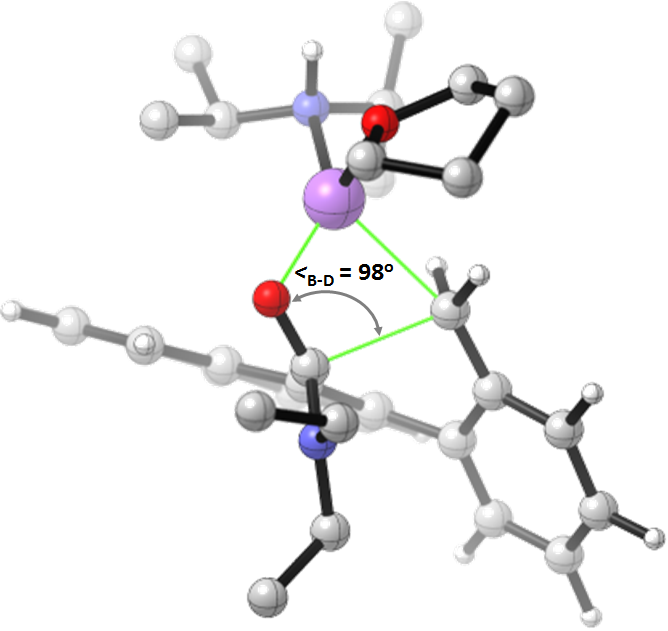
Consideration of the possible atropdiasteromers (Figure 4) in context of the VT-NMR results suggests that either the Ar-Arʹ or the Ar-CO rotation is observed depending on which process has the lowest rotational barrier. On the average, our calculations over-estimate the experimentally measured values by ~3.8 kJ/mol, with the largest deviation being 5.4 kJ/mol for compounds **1**, **5** and **8**. However, due to the consistency of over-estimation, the predicted barriers still correlate quite well with the experimental data.

Atropdiastereomer **8** which displays both Ar-Arʹ and Ar-CO hindered rotations, shows an energy barrier of ΔG‡ = 102.6 kJ/mol by EXSY-NMR (entry 9, Table 2) which is in good agreement with kinetic measurements for the atropisomerization of **8B** to **8A/B** by 1H NMR which gave ΔG‡ = 103.8 kJ/mol (entry 10, Table 2). DFT computation of concerted Ar-CO and C-N bond rotations for compound **8** gave ΔG‡ = 108.1 kJ/mol, in reasonable agreement with the experimental result. In contrast, the Ar-Arʹ rotation provided ΔG‡ = 126.7 kJ/mol. For a comparison, concerted Ar-CO and C-N bond rotations reported by Clayden on *N*-(2,5-pyrrolidin-1-yl)-2-methylnaphthalene-1-amide show a ΔG‡ ≈ 104 kJ/mol.[[31]](#endnote-31)

**Mechanistic Implications**

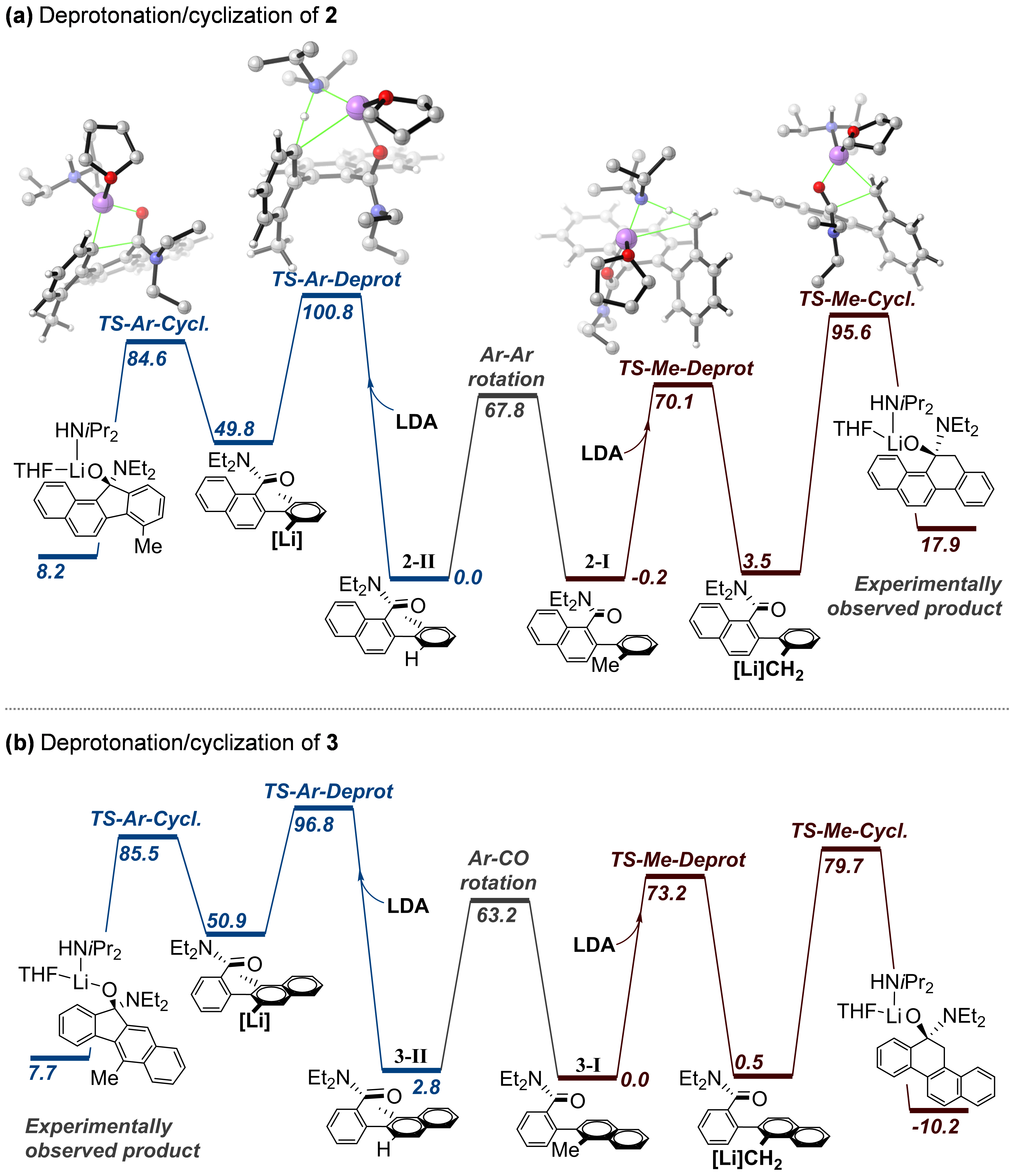
In order to enhance mechanistic understanding of the directed remote metalation (DreM) reaction,[[32]](#endnote-32) biaryls with DMG = 2-COOH[[33]](#endnote-33) and DMG = 2-CONEt2[[34]](#endnote-34) have been investigated for which a CIPE-induced mechanism has been implicated.15c The DreM reaction of 2-*N*,*N*-diethyl carbamoyl-2ʹ-methyl biaryls which undergo 2ʹ-methyl deprotonation and cyclization to phenanthrenes has seen extensive use in synthesis.15,32 For this series as well, the DreM reaction has been postulated to occur by initial coordination of the alkyllithium or lithiumdialkylamide base to the amide carbonyl (CIPE15c), followed by 2ʹ-Me deprotonation and attack of the resulting tolyl anion on the 2-amide C=O bond followed by aromatization to the corresponding 9-phenanthrol derivative. In the present study, we observed that the biaryl amides, all bearing 2ʹ-methyl groups, may be categorized in two series: compounds **1, 2, 4, 5, 7,** and **8** which undergo formation to phenanthrol derivatives16 (e.g., **9**,Scheme 1) and compounds **3** and **6**16which afford fluorenones (e.g. **10**).The favorable DMG effect of the fluoro substituent in **6** is evident in the regioselective formation of the corresponding fluorenone17 and in other cases.[[35]](#endnote-35)

Based on a considerable body of studies,[[36]](#endnote-36) nucleophilic reactions on amides involves an initial nucleophilic attack on the carbonyl to form a tetrahedral intermediate. In our systems, this demands that the biaryl *ortho*-amide adopt a rotational position in the transition state in accord with the Bürgi-Dunitz postulate[[37]](#endnote-37) which states that, in the trajectory of approach empirically developed from a large body of X-ray structures, a nucleophile-C-O bond angle of 100-110 ° (modified to 93-102 ° for O- and N-nucleophiles by Cieplak[[38]](#endnote-38)) is achieved in the last 2-3 Å before bond formation. Thus, as shown in Figure 7, the amide group must undergo rotation orienting the NR2 group partly over the biaryl system while the Ar-Arʹ bond undergoes concurrent rotation into a position for the tolyl carbanion to attack the carbonyl to achieve ring closure.



**Figure 7:** Computed transition state for the formation of the tetrahedral intermediate from 2ʹ-methyl anion approach to the recipient C=O of **3** along the Bürgi-Dunitz angle. Hydrogen atoms of the THF and *i*Pr2NH ligands and of the ethyl groups of the amide have been omitted for clarity.

X-ray crystallographic studies of *ortho*-lithiated *N*,*N*-diisopropylbenzamide and *N*,*N*-diisopropylnaphthamide have shown dimeric structures with amide-aryl bond angles of 47° and 65° respectively,[[39]](#endnote-39),[[40]](#endnote-40) indicating that, in the solid state, planarity of Li-C=O interaction is achieved to higher degree than the unmetalated systems (amide-Ar ring bond angle = 90°). Recent studies of laterally metalated *N,N*-diisopropyl 2-propylbenzamide in solution reveal similar structures in which Li is coordinated to both carbonyl and the anion, to the extent that tridentate amine and ether ligands PMDTA **(***N*,*N*,*N*ʹ,*N*ʹʹ,*N*ʹʹ-pentamethyldiethylenetriamine) and DGME (diglyme) prefer di-coordination to the tetra-coordinating Li, leaving one amine free in solution.[[41]](#endnote-41) With these mechanistic considerations in mind, a computational study was conducted for the proposed pathways of the DreM-reaction of **2** and **3** with LDA (Figure 8).[[42]](#endnote-42) The coordination sphere of Li was saturated with an explicitly included molecule of THF.[[43]](#endnote-43) The calculated mechanism included an initial endergonic precomplexation of LDA with the biaryl, followed by deprotonation of either the Arʹ-H or the 2ʹ-Me position, and subsequent nucleophilic attack of the resulting lithiated position on the amide carbonyl, resulting in ring-closure and eventual formation of either the fluorenone **10** or the phenanthrol **11** derivatives respectively. Bond lengths and angles in the coordination sphere of Li were quite similar to those obtained from X-ray structures by Wheatley.39-41 For both substrates **2** and **3**, the metalation of the 2ʹ-methyl group was found to be kinetically and thermodynamically favored over Arʹ-metalation (G**‡** = 24-31 kJ/mol). The Arʹ-metalation was found to be endergonic by ~50 kJ/mol, while the 2ʹ-Me-deprotonation was nearly thermoneutral (G < 4 kJ/mol). With a concerted rotation of Ar-Arʹ, Ar-CO and Et2N-CO bonds, both **2** and **3** show ability to attain transition states complying with the Bürgi-Dunitz angle (Figure 7) (fluorenone: 107 °, phenanthrol: 98-99 °). For Arʹ ring deprotonation-cyclization to the fluorenone (Figure 8), these transition states were found to be lower in energy than the preceding deprotonation step; for 2ʹ-Me-deprotonation-cyclization to the phenanthrols, they were higher in energy. However, since the transition states for the cyclization for both substrates **2** and **3** have lower barriers than the highest transition states in the Arʹ-deprotonation-cyclization pathway (G**‡** = 8-13 kJ/mol in favor of the fluorenone formation pathway), these calculations are not in full agreement with the different outcomes of the reactions of **2** and **3**. In fact, from the calculated model, the formation of the phenanthrol **11** from **3** is both thermodynamically and kinetically more favorable than the formation of **9** from **2**, which is opposite to experimental observation. Thus, the considered reaction mechanism and the employed computational results suggest that both of the substrates should favor 2ʹ-Me deprotonation and a subsequent ring closure to a phenanthrol derivative. Whether there is involvement of dimers, multiple metalations (sometimes 2 equivalents or more of organolithium base is needed for DreM reactions), or different complexation from those based on the CIPE concept15c are current speculations for further investigations.



**Figure 8:** Comparison of the free energy (in kJ/mol) pathways of 2ʹ-Me- and Arʹ ring-deprotonation and the consecutive cyclization reactions of **2** (top) and **3** (bottom). Hydrogen atoms of the THF and *i*Pr2NH ligands and of the ethyl groups of the amide have been omitted for clarity. [Li] = ‑Li(*i*Pr2NH)(THF)42

**Conclusions**

Based on VT 1H NMR measurements, EXSY-NMR experiments, and computational studies, we conclude that 2-amido-2ʹ-methyl biphenyls **1-8** may be described as atropdiastereomers involved in dynamic bond rotation around the Ar-Arʹ- and Ar-CO-bonds, which is dependent on the location of substituents. For biaryls bearing a substituent *ortho* to the amide functional group (**2, 7**) in which the Ar-CO rotation is hindered, the Ar-Arʹ-rotation show the lower energy barrier (ΔG‡Tc = 64.8-67.5 kJ/mol), while compounds with substituents located *ortho* to the biaryl bond (**4**-**6**) which hinder this rotation, show lower energy barrier for the Ar-CO-rotation (ΔG‡Tc = 60.5-63.0 kJ/mol). For 2-amido-2ʹ-methyl biarylsbearing a substituent *ortho* to both the amide and the biaryl bond (**8**), true isolable atropisomers were observed at room temperature, and the measured energy barrier for atropdioastereomeric interconversion (ΔG‡ = 102.6-103.8 kJ/mol) complied with a concerted Ar-CO/Et2N-CO bond rotation. Neither calculated nor measured rotational barriers were sufficiently high to interfere with nor allow rationalization of the different reactivity of these compounds in the cyclization to fluorenones or phenathrols by the directed remote metalation (DreM) reaction.

**EXPERIMENTAL SECTION**

**General**

Tetrahydrofuran (THF) was distilled under nitrogen atmosphere from Na/benzophenone. *N*,*N*,*N*ʹ,*N*ʹ-tetramethylethylenediamine (TMEDA) was distilled and stored over potassium hydroxide (KOH). Glove box was used when necessary. All reactions were carried out under nitrogen atmosphere if not otherwise specified. TLC was performed on Merck silica gel 60 F254 plates, using UV light at 254 nm and 5% alcoholic molybdophosphoric acid for detection. Normalsil 60, 40-63µm silica gel was used for flash chromatography. 1H NMR and 13C NMR was recorded on a Varian Mercury 300 MHz (UiS, Norway), or a BRUKER ADVANCE-400 (automatic sample changer, BB auto tuning; Queen’s University, Kingston, ON, Canada), all at room temperature. Chloroform-*d1* was used as solvent, unless otherwise specified. Chemical shift was reported in ppm compared to TMS (δ 0, singlet, for 1H NMR), or for 13C resonance signal to CDCl3 (δ 77.0, triplet). The splitting pattern was recorded as a singlet, s; doublet, d; triplet, t; double doublet, dd; double triplet, dt; quartet, q; multiplet, m; broad, br. IR was recorded on a Perkin Elmer FT-IR spectrometer, version 3.02.01. HRMS was measured on a LTQ Orbitrap XL ion trap mass spectrometer with electrospray ionization. Melting points was determined on a Stuart Scientific melting point apparatus SMP3 and are uncorrected. Synthesis of compounds **1** and **4-8** were reported previously.16

## *N*,*N*-diethyl-2-(*o*-tolyl)-1-naphthamide (2). *N*,*N*-diethyl-1-naphthoylamide (0.781 g, 3.44 mmol) in THF (10 mL) was added dropwise to a solution of *s*-BuLi (3.7 mL, 5.17 mmol, 1.4 M solution in cyclohexane), TMEDA (0.77 mL, 5.16 mmol) in THF (10 mL) at -78 °C. After stirring for 1 h, triisopropyl borate (1.97 mL, 8.59 mmol) was added, the mixture stirred at -78 °C for 1.5 h and warmed to rt over 18 h. The reaction mixture was quenched with NH4Cl (15 mL) and extracted with diethyl ether (3 x 15 mL). The organic layer was washed with water (2 x 45 mL), dried over MgSO4, subjected to filtration, and concentrated *in vacuo* to give 1-(diethylcarbamoyl)naphthalene-2-yl)boronic acid as a brown oil which was used without further purification in the next experiment.

All solutions were degassed prior to use. A mixture of PdCl2(dppf) (118 mg, 0.14 mmol, 5 mol%) and 2-bromotoluene (0.35 mL, 2.91 mmol) in DME (6 mL) was stirred at room temperature for 15 min. The solution of (1-(diethylcarbamoyl)naphthalen-2-yl)boronic acid (3.44 mmol,) in DME (4 mL) was added, followed by 2M Na2CO3 (6 mL), at room temperature. The mixture was heated at reflux for 18 h, cooled, and extracted with diethyl ether (3 x 20 mL). The organic layer was dried over MgSO4, subjected to filtration and concentrated *in vacuo*. The crude product was purified by flash column chromatography (petroleum ether: ethyl acetate 2:1) to afford 0.85 g (92%) of **2** as a brown oil as a *ca* 3:2 mixture of rotamers. 1H NMR (CDCl3, 300 MHz): δ 7.87-7.85 (app. m, 3H), 7.57-7.51 (m, 3H), 7.42-7.13 (m, 4H), 3.89-3.82 (m, 1H), 3.26-2.70 (m, 3H), 2.26 and 2.19 (s, 3H), 0.94-0.70 (m, 6H); 13C NMR (CDCl3, 75 MHz): δ 168.9 (**C**O), 139.9 (**C**), 138.6 (**C**), 137.8 (**C**), 134.9 (**C**), 133.9 (**C**), 132.5 (**C**H), 131.2 (**C**H), 130.0 (04) (**C**H), 130.0 (0) (**C**H), 128.2 (**C**H), 128.1 (**C**H), 128.0 (**C**H), 127.7 (**C**Hx2), 127.5 (**C**H), 126.9 (**C**Hx2), 126.2 (**C**H), 125.6 (**C**H), 125.5 (**C**x2), 124.5 (**C**x2), 42.7 (N**C**H2 - minor rotamer), 42.1 (N**C**H2 - major rotamer), 37.7 (N**C**H2 - major rotamer), 37.5 (N**C**H2 - minor rotamer), 20.4 (**C**H3 - minor rotamer), 20.3 (**C**H3 - major rotamer), 13.9 (2) (NCH2**C**H3 - minor rotamer), 13.9 (0) (NCH2**C**H3 - major rotamer), 11.9 (NCH2**C**H3 - major rotamer), 11.6 (NCH2**C**H3 - minor rotamer); IR (KBr): 3055 (w), 2974 (m), 2933 (m), 2873 (w), 2238 (w), 1628 (s), 1492 (m), 1473 (m), 1434 (s), 1381 (m), 1280 (m), 1267 (m), 1221 (m), 1128 (m), 828 (m), 761 (m), 728 (m); Mass spectrum *m/z* (relative intensity %): 340.2 [M + Na]+ (100); HRMS (ESI) Calc. for C22H23ON + Na: 340.1672, Found 340.1671.

## *N*,*N*-diethyl-2-(1-methylnaphthalen-2-yl)benzamide (3). *N,N*-diethylbenzamide (612 mg, 3.45 mmol) in THF (10 mL) was added dropwise to a solution of s-BuLi (3.7 mL, 5.18 mmol, 1.4 M solution in cyclohexane), TMEDA (0.77 mL, 5.16 mmol) and THF (10 mL) at -78 °C. After stirring for 1 h, triisopropyl borate (1.97 mL, 8.59 mmol) was added, and the mixture stirred at -78 °C for 1.5 h and warmed to rt over 18 h. The reaction mixture was quenched with NH4Cl solution (15 mL), and extracted with diethyl ether (3 x 15 mL). The organic layer was washed with water (2 x 45 mL), dried over MgSO4, subjected to filtration and concentrated in vacuo to give 2-(N,N-diethylcarboxamido)phenylboronic acid as a brown oil which was used without further purification in the next experiment.

All solutions were degassed prior to use. A mixture of PdCl2(dppf) (116 mg, 0.14 mmol, 5 mol%) and 2-bromo-1-methylnaphthalene (638 mg, 2.89 mmol) in DME (6 mL) was stirred at room temperature for 15 min under inert atmosphere. The solution 2-(*N*,*N*-diethylcarboxamido)phenylboronic acid (3.45 mmol) in DME (4 mL) followed by 2M Na2CO3-solution (6 mL) was added at room temperature and the mixture was heated at reflux for 18 h. After allowing the mixture to cool down it was extracted with diethyl ether (3 x 20 mL). The organic layer was dried over MgSO4, subjected to filtration and concentrated *in vacuo*. The crude product was purified by flash chromatography (petroleum ether: ethyl acetate 2:1) afforded 862 mg (94%) of product **3** as a viscous, brown/red oil. 1H NMR (CDCl3, 300 MHz): δ 8.02 (app s, 1H), 7.78 (app. s, 1H), 7.50-7.30 (m, 8H), 3.58-2.35 (5 peaks app. s, 4H), 2.71 (s, 3H), 1.28-1.11 (m, 3H), 0.60-0.5 (app. d, *J* = Hz, 3H); 13C NMR (CDCl3, 75 MHz): δ 170.0 (**C**O), 137.2 (**C**), 134.2 (**C**), 131.5 (**C**), 129.0 (**C**H), 128.3 (**C**Hx2), 127.8 (**C**), 127.6 (**C**Hx2), 126.9 (**C**), 126.2 (**C**Hx2), 125.8 (**C**H), 125.5 (**C**Hx2), 124.6 (**C**); IR (KBr): 3064 (w), 2973 (m), 2933 (m), 2872 (w), 1630 (s), 1513 (w), 1458 (m), 1426 (m), 1381 (m), 1288 (m), 1221 (w), 1098 (m), 1078 (w), 871 (w), 836 (w), 784 (w), 764 (m), 480 (m), 471 (m); Mass spectrum *m/z* (relative intensity %):340.2 [M + Na]+ (74): HRMS (ESI) Calc. for C22H23NO + Na: 340.16789. Found 340.16773.

**Variable temperature NMR**

Variable temperature NMR (VT NMR) were carried out with temperatures ranging from 190 Kelvin (-83 °C) to 332 Kelvin (59 °C). All compounds were recorded on a 400 MHz Bruker NMR using toluene-*d8* as solvent. The temperature of the probe used in these experiments was calibrated by an ethylene glycol solution. From Δ*υ*, the rate constant, *k*c (the rate for rotamer interconversions) at coalescence temperature, Tc, was calculated by the Gutowsky-Holm equation (*k*c = πΔυ / √2 = ~2.22Δυ s-1).[[44]](#endnote-44) The temperature was raised from a temperature sufficiently low to observe separate signals for the tolyl methyl group, until the coalescence of the two peaks was reached. From the coalescence temperature, Tc, and the rate constant, *k*c, the activation energy of rotation (ΔG‡Tc) was calculated using the Eyring equation (ΔG‡Tc = *RTc*[23.76-ln(*kc/Tc*)]), where R is the gas constant, 8.3145 kJ/mol.[[45]](#endnote-45) The margin of error was estimated to ± 0.8 kJ/mol by estimating the accuracy of Δ*υ* and Tc from the measurements and applying the extremities in the calculations.

Spectra of the variable temperature experiments of compound **1**-**7** are given in SI, section 1 together with a more comprehensive description. The rotational barriers for compound **8** could not be measured as the coalescence temperature was not reached at 380 K, which was the limit of the NMR probe.

**EXSY NMR experiments**

The EXSY experiments[[46]](#endnote-46) were performed on a Bruker Avance-600 spectrometer using the standard noesygpph pulse program. The spectra window was set to 8 ppm and 2K x 512 data points were acquired and zerofilled to 2K x 2K. Different mixing times were performed from 0.001 s up to 80% of the relaxation time of the appropriate nuclei. The mixing time was changed to obtain EXSY spectra that contained no exchange between peaks, and compare it with EXSY spectra that contained exhange between peaks. Integration of diagonal and cross peaks and comparison with the two mixing times provided the rate of the isomeric interconversion. From the rate of interconversion k at a given temperature T, the rotational energy barrier was then calculated (ΔG≠ = *RT*[23.76-ln(*k/T*)]). The data were processed using MestReNova and the integration of the various signals was analyzed using the EXSYCalc program from MestReC to obtain the rate constant.[[47]](#endnote-47)

**EXSY NMR of 8.** While the coalescence of the two isomeric 2´-methyl peaks of **8** was not observed in neither toluene-d8, DMSO-d6 nor acetone-d6; the rotational barrier could be measured from the volume integration of the exchanging peaks in an EXSY experiment by varying the mixing time from 1 to 0.01 sec at 380 K.[[48]](#endnote-48) At 1 sec, the exchange between the protons could be observed, while at mixing time 0.01 sec, no exchange was observed. By volume integration of the exchanging peaks, the rate of interconversion and the rotational energy barrier was calculated.In this EXSY experiment, two interconversions were observed: a) The atropdiastereomeric **A-B** interconversion from the exchanging peaks of aromatic protons (ΔG‡EXSY = 102.6 kJ/mol), and b) the *syn*/*anti* Et2N-CO **A-A´** amide interconversionfrom the peaks representing the CH3-amide proton exchange (ΔG‡EXSY = 98.0 kJ/mol).

NMR spectra and a more comprehensive discussion of these experiments are given in SI, section 2.

**ASSOCIATED CONTENT**

**Supporting information**

NMR spectra of new compounds, detailed discussion and spectra of VT and EXSY NMR experiments and DFT calculations are given in the supporting information.

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**REFERENCES**

1. Christie, G. H.; Kenner, J. H. *J. Chem. Soc.* **1922**, *121*, 614-620. [↑](#endnote-ref-1)
2. Alkorta, I.; Elguero, J.; Roussel, C.; Vanthuyne, N.; Piras, P. in Advances in Heterocyclic Chemistry; Katritzky, A., Ed.; Academic Press, **2012**, Vol. 105, p. 1-188. [↑](#endnote-ref-2)
3. (a) Peck, T. G.; Lai, Y. H. *Tetrahedron* **2009**, *65*, 3664-3667. (b) Mazzanti, A.; Lunazzi, L.; Minzoni, M.; Anderson, J. E. *J. Org. Chem.* **2006**, *71*, 5474-5481. (c) Leroux, F. *Chem. Bio. Chem*. **2004**, *5*, 644-649. (d) Ceccacci, F.; Mancini, G.; Mencarelli, P.; Villani, C. *Tetrahedron: Asym*. **2003**, *14*, 3117-3122. [↑](#endnote-ref-3)
4. (a) Brunel, J. M. *Chem. Rev*. **2005**, 105, 857-897. (b) Meca, L.; Reha, D.; Havlas, Z. *J. Org. Chem*. **2003**, 68, 5677-5680. [↑](#endnote-ref-4)
5. (a) Kumarasamy, E.; Raghunathan, R.; Sibi, M. P.; Sivaguru, J. *Chem. Rev*. **2015**, *115* (20), 11239–11300. (b) Bringman, G.; Mortimer, A. J. P.; Keller, P. A.; Gresser, M. J.; Garner, J.; Breuning, M. *Angew. Chem. Int. Ed.* **2005**, *44*, 5384-5427. (c) Berthod, M.; Mignani, G.; Woodward, G.; Lemaire, M. *Chem. Rev*. **2005**, *105*, 1801-1836. (d) Pu, L. *Chem. Rev*. **1998**, *98*, 2405-2494. [↑](#endnote-ref-5)
6. (a) Smyth, J. E.; Butler, N. M.; Keller, P. A. *Nat. Prod. Rep.* **2015**, *32*, 1562-1583. (b) Bringmann, G.; Gulder, T.; Gulder, T. A. M.; Breuning, M. *Chem. Rev*. **2011**, *111*, 563-639. (c) Lloyd-Williams, P.; Giralt, E. *Chem. Soc. Rev.* **2001**, *30*, 145-157. (d) Williams, D. H.; Bardsley, B. *Angew. Chem. Int. Ed.* **1999**, *38*, 1172-1193. (e) Nicolaou, K. C.; Boddy, C. N. C.; Bräse, S.; Winssinger, N. *Angew. Chem. Int. Ed.* **1999**, *38*, 2096-2152. (f) Torssell, K. B. G. *Natural Product Chemistry*, Taylor and Francis, New York, **1997**. [↑](#endnote-ref-6)
7. (a) Zask, A.; Murphy, J.; Ellestad, G. A. *Chirality* **2013**, *25*, 265-274. (b) LaPlante, S. R.; Fader, L. D.; Fandrick, K. R.; Fandrick, D. R.; Hucke, O.; Kemper, R.; Miller, S. P. F.; Edwards, P. J. *J. Med. Chem*. **2011**, *54*, 7005-7022. (c) Laplante, S. R,; Edwards, P. J.; Fader, L. D.; Jakalian, A.; Hucke, O. *Chem. Med. Chem*. **2011**, *6*, 505-513. (d) Clayden, J.; Moran, W. J.; Edwards, P. J.; Laplante, S. R. *Angew. Chem. Int. Ed*. **2009**, *48*, 6398-6401. [↑](#endnote-ref-7)
8. (a) Tietze, L. F.; Schuster, H. J.; von Hof, J. M.; Hampel, S. M.; Colunga, J. F.; John, M. *Chem. Eur. J.* **2010**, *16*, 12678-12682. (b) Clayden, J. *Chem. Commun*. **2004**, 127-135. (c) Bowles, P.; Clayden, J.; Helliwell, M.; McCarthy, C.; Tomkinson, M.; Westlund, N. *J. Chem. Soc., Perkin Trans. 1* **1997**, 2607-2616. (d) Clayden, J. *Angew. Chem. Int. Ed.* **1997**, *36*, 949-951 and references cited therein. (e) Cuyegkeng, M. A.; Mannschreck, A. *Chem. Ber.* **1987**, *120*, 803-809. (f) Ackerman, J. H.; Laidlaw, G. M.; Snyder, G. A. *Tetrahedron Lett*. **1969**, 3879-3882. [↑](#endnote-ref-8)
9. (a) Berthelot-Bréhier, A.; Panossian, A.; Colobert, F.; Leroux, F. R. *Org. Chem. Front*. **2015**, 2, 634-644. (b) Clayden, J.; Worrall, C. P.; Moran, W. J.; Helliwell, M. *Angew. Chem. Int. Ed.* **2008**, *47*, 3234-3237. (c) Koide, H.; Hata, T.; Uemura, M. *J. Org. Chem*. **2002**, *67*, 1929-1935. (d) Thayumanavan, S.; Beak, P.; Curran, D. P. *Tetrahedron Lett*. **1996**, *37*, 2899-2902. [↑](#endnote-ref-9)
10. (a) Clayden, J.; Lai, L. W.; Helliwell, M. *Tetrahedron* **2004**, *60*, 4399-4412 and references cited therein. (b) Rios, R.; Kimeno, C.; Carroll, P. J.; Walsh, P. J. *J. Am. Chem. Soc.* **2002**, *124*, 10272-10273. (c) Dai, W. M.; Lau, C. W. *Tetrahedron Lett.* **2001**, *42*, 2541-2544. [↑](#endnote-ref-10)
11. (a) Dai, W. M.; Yeung, K. K. Y.; Chow, C. W.; Williams, I. D. *Tetrahedron: Asym.* **2001**, *12*, 1603-1613. (b) Clayden, J.; Johnson, P.; Pink, J. H.; Helliwell, M. *J. Org. Chem*. **2000**, *65*, 7033-7040. [↑](#endnote-ref-11)
12. (a) Meyers, A. I.; Nelson, T. D.; Moorlag, H.; Rawson, D. J.; Meier, A. *Tetrahedron* **2004**, *60*, 4459-4473. (b) Clayden, J.; McCarthy, C.; Cumming, J. G. *Tetrahedron Lett.* **2000**, *41*, 3279-3283. [↑](#endnote-ref-12)
13. (a) Wiberg, K. B. In Greenberg, A.; Breneman, C. M.; Liebman, J. F. (eds.), *The Amide Linkage: Structural Significance in Chemistry, Biochemistry, and Materials Science*, Wiley, New York, NY, **2003**; pp 33-46. (b) Scherer, G.; Kramer, M. L.; Schutkowski, M.; Reimer, U.; Fischer, G. *J. Am. Chem. Soc*. **1998**, *120*, 5568-5574. (c) Stewart, W. E.; Siddall, T. H. *Chem. Rev*. **1970**, *70*, 517-551. [↑](#endnote-ref-13)
14. (a) Bragg, R. A.; Clayden, J.; Morris, G. A.; Pink, J. H. *Chem. Eur. J.* **2002**, *8*, 1279-1289. (b) Clayden, J.; Frampton, C. S.; McCarthy, C.; Westlund, N. *Tetrahedron* **1999**, *55*, 14161-14184. (c) Clayden, J.; McCarthy, C.; Helliwell, M. *Chem. Commun.* **1999**, 2059-2060. (d) Clayden, J. *Synlett*, **1998**, 810-816. (e) Ahmed, A.; Bragg, R. A.; Clayden, J.; Lai, L. W.; McCarthy, C.; Pink, J. H.; Westlund, N.; Yasin, S. A. *Tetrahedron* **1998**, *54*, 13277-13294. [↑](#endnote-ref-14)
15. (a) Wang, X.; Fu, J.-m.; Snieckus, V. *Helv. Chim. Acta* **2012**, *95*, 2680. (b) Snieckus, V.; Macklin, T. in *Handbook of C-H Transformations*; Dyker, G., Ed.; Wiley-VCH: Weinheim, Germany, **2005**; Vol. 1, p 106. (c) Whisler, M. C.; MacNeil, S.; Snieckus, V.; Beak, P. *Angew. Chem., Int. Ed.* **2004**, 43, 2206-2225. (d) Fu, J. M.; Snieckus, V. *Can. J. Chem.* **2000**, *78*, 905. [↑](#endnote-ref-15)
16. Jørgensen, K. B.; Rantanen, T.; Dörfler, T.; Snieckus, V. *J. Org. Chem.* **2015**, *80*, 9410-9424. [↑](#endnote-ref-16)
17. Beak has shown that directed *ortho* metalation in amides is dependent on the dihedral angle between the hydrogen abstracted and the proximate amide carbonyl. Our biaryl cases deal with very different geometry considerations in remote metalation of aryl C-H and C-CH3 deprotonations: Beak, P.; Kerrick, S. T.; Gallagher, D. J. *J. Am. Chem. Soc.* **1993**, *115*, 10628-10636. [↑](#endnote-ref-17)
18. Mohri, S.-I.; Stefinovic, M.; Snieckus, V. *J. Org. Chem*. **1997**, *62*, 7072-7073. [↑](#endnote-ref-18)
19. (a) Friebolin, H., *Basic One- and Two-Dimensional NMR spectroscopy, 3rd Ed.*, Wiley: Weinheim, Germany, **1998**, chapter 11. (b) Günther, H., *NMR spectroscopy – Basic principles, concepts, and applications in chemistry*, 2nd Ed., Wiley: New York, **1995**, chapter 9. (c) Allerhand, A.; Gutowsky, H. S.; Jonas, J.; Meinzer, R. A. *J. Am. Chem. Soc.*, **1966**, *88*, 3185-3194. [↑](#endnote-ref-19)
20. Kincaid, J. F.; Eyring, H.; Stearn, A. E. *Chem. Rev.*, **1941**, *28*, 301. [↑](#endnote-ref-20)
21. The available NMR probe could not safely achieve a higher temperature. [↑](#endnote-ref-21)
22. Meier, B. H.; Ernst, R. R. *J. Am. Chem. Soc*. **1979**, *101,* 6441. [↑](#endnote-ref-22)
23. Förster, H.; Vögtle, F. *Angew. Chem., Int. Ed.* **1977**, *16*, 429-441. [↑](#endnote-ref-23)
24. Baker, R. W.; Brkic, Z.; Sargent, M. V.; Skelton, B. W.; White, A. H. *Aust. J. Chem*. **2000**, *53*, 925-938. [↑](#endnote-ref-24)
25. Berg, U.; Bladh, H. *Acta Chem. Scand.* **1998**, *52*, 1380-1385. [↑](#endnote-ref-25)
26. Wolf, C.; Hochmuth, D. H.; König, W. A.; Roussel, C*. Liebigs Ann*. **1996**, 357-363. [↑](#endnote-ref-26)
27. Bott, G.; Field, L. D.; Sternhell, S. *J. Am. Chem. Soc*. **1980**, *102*, 5618-5626. [↑](#endnote-ref-27)
28. (a) Gaussian 09, Revision A.02, Frisch, M. J.; *et al*., Gaussian, Inc., Wallingford CT, **2009**. [See SI for full reference]. (b) While M06-2X functional has been successfully employed in the computational prediction of rotation barriers (see ref 28c) we observed that in our case it yielded ~4 kJ/mol higher barriers for the Ar-Ar and Ar-CO rotations and ~4 kJ/mol lower barriers for the C-N and combined rotations than M06L functional. (c) Jackson, K. E.; Mortimer, C. L.; Odell, B.; McKenna, J. M.; Claridge, T. D. W.; Paton, R. S.; Hodgson, D. M. *J. Org. Chem.* **2015**, *80*, 9838-9846. [↑](#endnote-ref-28)
29. Two possible transition states were considered where either the carbonyl or the NR2 group of the amide is towards the *o*-aryl substituent. (See SI for details.) [↑](#endnote-ref-29)
30. (a) Clayden, J.; Lund, A.; Vallverdú, L.; Helliwell, M. *Nature* **2004**, 431, 966-971. (b) Betson, M. S.; Bracegirdle, A.; Clayden, J.; Helliwell, M.; Lund, A.; Pickworth, M.; Snape, T. J.; Worrall, C. P. *Chem. Commun*. **2007**, 754-756. [↑](#endnote-ref-30)
31. Clayden, J.; Pink, J. H. *Angew. Chem. Int. Ed.* **1998**, *37*, 1937-1939. [↑](#endnote-ref-31)
32. Tilly, D.; Magolan, J.; Mortier, J. *Chem. Eur. J.* **2012**, *18*, 3804-3820. [↑](#endnote-ref-32)
33. (a) Castanet, A.-S.; Tilly, D.; Véron, J.-B.; Samanta, S. S.; De, A.; Ganguly, T.; Mortier, J. *Tetrahedron* **2008**, *64*, 3331-3336. (b) Tilly, D.; Samanta, S. S.; Castanet, A.-S.; De, A.; Mortier, J. *Eur. J. Org. Chem.* **2006**, 174-182. (c) Tilly, D.; Samanta, S. S.; De, A.; Castanet, A.-S.; Mortier, J. *Org. Lett*. **2005**, *7*(5), 827-830. [↑](#endnote-ref-33)
34. Tilly, D.; Fu, J.-M.; Zhao, B.-P.; Alessi, M.; Castanet, A.-S.; Snieckus, V.; Mortier, J. *Org. Lett.* **2010**, *12*(1), 68-71. [↑](#endnote-ref-34)
35. Kli´s, T.; Luli´nski, S.; Serwatowski, J. *Curr. Org. Chem*. **2008**, *12*, 1479-1501. [↑](#endnote-ref-35)
36. McMurry, J.; «Organic Chemistry», 9th Ed., Cengage Learning, USA, **2016**, p. 710-711. [↑](#endnote-ref-36)
37. Bürgi, H. B.; Dunitz, J. D.; Lehn, J. M.; Wipf, G. *Tetrahedron* **1974**, *30*, 1563-1572. [↑](#endnote-ref-37)
38. Cieplak, A. S., *Organic addition and elimination reactions; transformation paths of carbonyl derivatives* in *Structure Correlation vol. 1*; Bürgi, H.-B.; Dunitz, J. D. (Eds), VCH, Weinheim, Germany, **1994**, p. 205-302 (page 210-218). [↑](#endnote-ref-38)
39. Clayden, J.; Davies, R. P.; Hendy, M. A.; Snaith, R.; Wheatley, A. E. H. *Angew. Chem., Int. Ed.* **2001**, *40*, 1238-1240. [↑](#endnote-ref-39)
40. Armstrong, D. R.; Clayden, J.; Haigh, R.; Linton, D. J.; Schooler, P.; Wheatley, A. E. H. *Chem. Commun*. **2003**, 1694-1695. [↑](#endnote-ref-40)
41. (a) Vincent, M. A.; Smith, A. C.; Donnard, M.; Harford, P. J.; Haywood, J.; Hillier, I. H.; Claydon, J.; Wheatley, A. E. H. *Chem. Eur. J.* **2012**, *18*, 11036-11045. (b) Wheatley, A. E. H.; Clayden, J.; Hillier, I. H.; Smith, A. C.; Vincent, M. A.; Taylor, L. J.; Haywood, J. *Beilstein J. Org. Chem.* **2012**, *8*, 50-60. [↑](#endnote-ref-41)
42. Calculations were performed at the CPCM (THF) M06L/6-311++G(d) // CPCM (THF) B97XD/6-31+G(d) level of theory. [↑](#endnote-ref-42)
43. Various ligation states of Li were considered: only *i*Pr2NH base; two THF molecules; and a combination of one *i*Pr2NH and one THF molecule. The latter was found to be the most favored in all considered stages of the reaction. [↑](#endnote-ref-43)
44. (a) Modarresi-Alam, A.R.; Amirazizi, A.; Bagheri, H.; Bijanzadeh, H.R.; Kleinpeter, E. *J. Org. Chem,* **2009**, *74*, 4740-4746. (b) Günther, H., *NMR spectroscopy – Basic principles, concepts, and applications in chemistry*, 2nd edition, Wiley: New York, 1995, chapter 9. (c) Friebolin, H., *Basic One- and Two-Dimensional NMR spectroscopy*, 3rd edition, Wiley: Weinheim, Germany, 1998, chapter 11. (d) Yamagami, C.; Takao, N.; Takeuchi, Y. *Aust. J. Chem.* **1986**, *39*, 457-463. [↑](#endnote-ref-44)
45. Kincaid, J.F.; Eyring, H.; Stearn, A.E. *Chem. Rev.* **1941**, *28*, 301. [↑](#endnote-ref-45)
46. Meier, B. H.; Ernst, R.R. *J. Am. Chem. Soc.* **1979**, *101*, 6441. [↑](#endnote-ref-46)
47. EXSYCalc Version 1.0, Juan Carlos Cobas, M Martin Pastorm Copyright © MestReC 2004 [↑](#endnote-ref-47)
48. (a) Zolnai, Z.; Juranic,N.; Vikic-Topic, D.; Macura, S. *J. Chem. Inf. Comput. Sci.* **2000**, *40*, 611-621. (b) Lu, J.; Ma, D.; Hu, J.; Tang, W.; Zhu, D. *J. Chem. Soc. Dalton Trans.* **1998**, 2267-2273. [↑](#endnote-ref-48)