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Adding a Structural Context to the Deprotometalation and Trans-Metal-Trapping Chemistry of Phenyl-substituted Benzotriazole


Abstract: Organometallic bases are becoming increasingly complex since mixing components can lead to bases superior to single component bases. To better understand this superiority, it is useful to study metalated intermediate structures prior to quenching. This study is on 1-phenyl-1H-benzotriazole, which was previously deprotonated by an in situ ZnCl₂·TMEDA/LiTMP mixture and then iodinated. Here, reaction with LiTMP exposes the deficiency of the single component base as the crystalline product obtained was [(4-R-1-(2-lithiophenyl)-1H-benzotriazole·3THF)], [R=2-C₆H₄(Ph)N], in which ring opening of benzotriazole and N₂ extrusion had occurred. Supporting lithiation by adding Bu₂Al(TMP) induces trans-metal-trapping where C-Li bonds transform into C-AI bonds to stabilise the metalated intermediate. X-ray diffraction studies revealed homodimeric [(4-R-1-phenyl-1H-benzotriazole)], [R=2-C₆H₄(Ph)N·TMP·Li], and its heterodimeric isomer [(4-R-1-phenyl-1H-benzotriazole)(1-(2-R'-phenyl)-1H-benzotriazole)], whose structure and slow conformational dynamics (activation enthalpy ΔH° = 71 ± 5 kJ mol⁻¹, Gibbs free activation energy at 298 K ΔG°‡ = 67 ± 5 kJ mol⁻¹ and activation energy E_a = 75.4 ± 0.4 kJ mol⁻¹) were probed by solution NMR spectroscopy.

Introduction

The benefit of studying the structures of organometallic bases (deprotonating agents) and the metalated intermediates of their deprotonation (C-H to C-metal exchange) reactions is all too apparent. Conveying this idea in a complete context, Schlosser[4] remarked eloquently that “no difference in organometallic reactivity patterns can be rationalised unless the metal and its specific interactions with the accompanying carbon backbone, the surrounding solvent, and the substrate of the reaction, are explicitly taken into account”. In other words, reactivity and structures are inseparable. At the time this quotation was written, most organometallic bases were single-metal species; whereas today mixed-metal bases are often used in synthetic campaigns. This presence of two or occasionally more distinct metals within the base composition can add greatly to the complexity involved, making structural information even more valuable towards attaining a full understanding of the chemistry. Without such structural characterisation, our recent report of rare ortho-meta’ and meta-meta’ deprotonations of several aromatic substrates, the latter, a seeming heterodoxy to the doctrine of directed ortho-metalation (DOM) may have lacked a plausible explanation. However, the rationale proposed was of a special template effect whereby the ring structure of the “pre-inverse-crown” base[5] controls the number and regioselectivity of deprotonations within the “inverse crown” product to a greater extent than the directing group attached to the aromatic substrate. This small but significant step towards realising designer bases has stimulated us to explore areas of deprotonation chemistry where structural information is scant or missing completely. Mongin’s recent report of the deprotonation of 1-aryl-H-benzotriazoles by a lithium-zinc mixture (Scheme 1 shows examples where aryl = phenyl) caught our eye in this context for its interesting regioselectivities determined indirectly through iodination of the active metallo intermediates. This paper also summarised the few lithiation studies carried out on these heterocycles, none of which, significantly, reported any structurally characterised lithio intermediates. Surprisingly few crystal structures are known even for N-lithiated benzotriazoles. A notable exception is cationic [(PMDETA·Li)(C₆H₄N₂)] reported by Wright[7] which was not synthesised from benzotriazole (C₆H₄N₂N₂) but via a remarkable oxidative dehydrocoupling of o-phenylenediamine triggered by a lithium-tin(II) alkyl-amido base. While seminal studies by Katritzky[8] have established benzotriazole as a versatile synthetic auxiliary, its importance extends to numerous material and biochemical applications. Here in this study we set about gaining reactivity and structural insights into the deprotonetalation chemistry of 1-phenyl-1H-benzotriazole 1.

Scheme 1. Stepwise deprotonization/iodination reactions of 1-phenyl-1H-benzotriazole as reported by Mongin et al.

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Results and Discussion

Deprotolitiation without trans-metal-trapping: With only modest success in trying to fill the structural gaps in knowledge between reactants and iodinated products in the mixed lithium-zinc base system (see later in article), we turned to a related lithium-aluminium system that has previously provided valuable insights into related deprotonmetalations of anisole and N,N-dimethylphenylamine (DMPEA). An added incentive for including aluminium is the paucity of crystallographically characterised aluminium benzotriazoles with a rare exception being a series of phenoxbenzotriazole derivatives studied for their photoluminescent properties.

In the lithium-aluminium question in hand, LiTMP performs the metalation of the C-H substrate while the diacylaluminium amide \( \text{Bu}_2\text{Al}(\text{TMP}) \) performs the trans-metal-trapping of the developing C- carbonan (TMP = 2,2,6,6-tetramethylpiperidide). As explained elsewhere, we do not use the more common term transmetalation as it usually signifies the exchange of metals in separate condensates, whereas here the aluminium compound inserts into the C-Li bond to trap and simultaneously stabilise the carbane within the one mixed lithium-aluminium compound, hence trans-metal-trapping seems more appropriate. We first tested the lihtiation of \( \text{1} \) with LiTMP on its own. Though lithiations of benzotriazoles have normally been performed at low temperatures, for a direct comparison with the aluminium system we carried out the reaction at room temperature for two hours in \( \text{n} \)-hexane and then THF was added to the mixture. As expected this unsupported (without aluminium trapping) lithiation reaction was unclean producing an unpromising dark solution over several days produced one crop of red crystals identified by X-ray crystallography as the dilithio benzotriazole derivative \([\{4-R-1-(2-lithiophenyl)-1H-benzotriazole-3THF\}_2]\), \( \text{2} \), where \( R = 2\)-C\( \text{H}_6\)(Ph)NLi. The crystalline yield obtained was only 10%.

As revealed in Figure 1a, crystalline \( \text{2} \) exists as a discrete, centrosymmetric dimer. The structure is easily comprehensible from consideration of a monomeric unit. A phenyl-benzenetriazole scaffold is substituted in two distinct positions. The 4-position of the benzo ring is occupied by a diphenylamido-lithium fragment (containing Li1-N4) connected by a new C-C (C2-C19) bond involving ortho-C(19), (see Figure 2a) while the 2-position of the 1-phenyl substituent has been lithiated (Li2-C8, see Figure 2b).

Dimerisation occurs through four bonds, Li1-N2′ (the central N atom of the triazole unit), Li2-N4′ (the N atom of the amide) and their symmetrical equivalents Li1′-N2 and Li2′-N4 respectively. Bonding to two triazole N atoms (N3 and N2′), one amido N atom (N4) and one THF O atom (O1), distorted tetrahedral Li1 is distinct from Li2 whose distorted tetrahedral environment is made up of an ortho-C atom (C8), an amido N atom (N4) and two THF O atoms (O2 and O3). A different perspective of the structure (Figure 1b) shows the internal Li1/Li1′ atoms belong to an essentially planar 6-5-6-5-6 fused ring system (plane of central 6 membered ring lies at 13.64(5)° to benzotriazole plane) with the Li1/Li1′ atoms connecting the two symmetrically equivalent benzotriazole frameworks through their N2N3/N3′N2′ junctions. External Li2/Li2′ atoms, more sterically accessible enabling greater THF solvation, lie either side of the fused ring plane as do the amido atoms (N4/N4′). Both the 1-phenyl and 4-phenyl ring planes twist out of the fused ring plane by 39.0(5)° and 46.17(7)° respectively to which they are attached (via N1 and C2 respectively), while the second phenyl ring of the amido unit lies more remote but overlapping slightly with the fused core. Key bond dimensions are given in the legend to Figure 1. The Li-N bond lengths span the narrow range, 2.087(5) – 2.140(5)Å, as the anionic (N4) or dative (N2, N3) nature of the bonding is offset by the different hybridisations involved (N2: N3: sp², N4, sp³). Similar Li-N (azole) bond lengths [1.992(5)Å; 2.044(5)Å] are encountered in the dimeric N-lithiated benzotriazole iron(II) amide co-complex ([Li(bta)(HMDS)]\( _2 \)) (HMDS = 1,1,3,3,3-hexamethyldisilazide) a close analogy of 2 as evidenced by its same 6-5-6-5-6 fused ring centrepiece (Figure 3). Interestingly, this bimetallic structure was interpreted in the context of the ring-laddering principle conceived by Snith and Wade on the basis of their seminal studies of lithium amide structures. Thus it was interpreted as an iron amide intermolecularly intercepted dimeric unit of polymeric \([\text{DMSO-Li}(\text{bta})]_{\infty}\) (DMSO = dimethylsulfoxide) an infinite ladder structure characterised crystallographically by Schleyer (Figure 3). The 6-5-6-5-6 fused ring motifs also present in
Significantly, switching the alkali metal from lithium to potassium in [(HMPA·K(bta)]_2 fused (KN)_2 maintains the polymeric ladder constitution though the single 6-membered (LiNNLiNN) repeating unit is replaced by alternating, the composition and structure of benzotriazole scaffold, namely the C2-C19 bond (a) and the Li2-C8 bond (b). Views of Figure 2.

Mechanistic considerations: The most interesting aspect of the composition and structure of 2 is what they infer about its possible mechanism of formation. Two features stand out. First, the new C2-C19 bond that creates the biphenyl unit, and second the diphenylamido function which, originating from a 1-phenyl-1H-benzotriazole molecule implies a dinitrogen (N_2) extrusion has taken place. Scheme 2 shows a plausible mechanism that can account for these events. Starting from a mono-deprotonated derivative of 1 with the carbanionic centre located at the 4-position of the benzo ring (consistent with Mongin’s in situ zinc study) if the formation of this monolithium complex is slow then it in turn can react with non-metalated molecules of starting material 1. Nucleophilic attack of the ipso-sp^2C bonded to N1 by this carbanion will induce extrusion of N_2 to create an attached diphenylamido function. Since the reaction solution contained two equivalents of LiTMP per benzotriazole molecule, excess base is available to lithiate the 1-phenyl substituent under DoM control to give the isolated product 2. Alternatively the second TMP lithiation could have taken place at the onset of the reaction prior to the nucleophilic substitution/dinitrogen extrusion process. The conclusion from this part of the study is that LiTMP lithiation under the ambient temperature conditions studied is as anticipated ineffectual. The unsupported (in the sense of not having bimetallic synergetic stabilisation, see below) highly polar lithio intermediates are unstable and can undergo heterocycle-destroying side reactions. One such side reaction has been described here but given that the isolated yield of the product was only about 10% and the NMR spectra of reaction filtrates were unclean showing starting material, 2 and other unidentified products, it can be assumed that other side reactions are taking place.

Deprotolithiation with trans-metal-trapping via an aluminium protocol: We repeated the LiTMP lithiation of 1 but this time in the presence of the dialkylaluminium amide iBu_2Al(TMP). Due to the considerable steric bulk of both metal compounds they do not form an aluminate [Li’(iBu_2Al(TMP))_2] so LiTMP is free to deprotonate the triazole (note TMP-aluminates have been found to be weak bases compared to lithium bases, generally incapable of deprotonating C-H bonds). The reaction was carried out in two different stoichiometries with one or two molar equivalents of the base mixture to one equivalent of the triazole (Scheme 3). Mono- and di-deprotonation respectively were expected. In practice, only monodeprotonation was observed under either stoichiometry, but interestingly this was manifested in two distinct crystalline solids.

Established by X-ray crystallography, the formula of the minor product is [4-R’-1-phenyl-1H-benzotriazole]_2, 3 where R’=iBu_2Al(µ-TMP)Li. This C_2 symmetric dimeric structure was produced from a 1:1, benzotriazole:base stoichiometric reaction.
Its classification as the minor (isolable) product reflects that it was isolated only from one reaction, whereas several repeated attempts under seemingly identical conditions afforded the major product, the heterodimeric isomer of 3, [(4-R’-1-phenyl-1H-benzotriazole)·(1-(2-R’-phenyl)-1H-benzotriazole)]. 4. With respect to the starting material 1, a single deprotonation has taken place at the 4-position of the benzo ring (C5) and two identical molecules have joined up to form the homodimeric structure of 3 (Figure 4). One of these benzo-deprotonated molecules forms one half of 4 with the other half a molecule of 1 selectively deprotonated at the ortho position of the phenyl ring (C12) to complete the tetracnucle (2xAI; 2xLi) structure (Figure 5). Formally these deprotonations are aluminations (C-H to C-Al exchanges) as in both 3 and 4 the deprotonated C atoms bond to Al through C5-Al1 and C12-Al1/C49-Al2 respectively, with the coordination sphere of each Al atom completed by two iBu ligands and a N atom from TMP which bridges to Li. Within the monomeric moiety of 3 the Li atom bonds to TMP (Li1-N4) and the outer unsubstituted triazole N (Li1-N3), with the dimerization connection being to the central triazole N (Li1-N2)’ to complete a distorted trigonal planar, (N3)x lithium coordination. Though the two distinct Li atoms in 4 also coordinate to 3 N atoms they do it differently from that in 3 and from each other. Thus, Li1 binds to two central triazole N atoms (Li1-N2, Li1-N6) and a TMP N (Li1-N4); whereas Li2 binds to two unsubstituted outer triazole N atoms (Li2-N3, Li2-N7) and a TMP N (Li2-N6 and Li2-N3) connecting the two distinct monomeric, dinuclear moieties. Though all three structures 2, 3 and 4 share the same 6-5-6-5-6 fused ring core, conformationally these cores are distinct (Figure 6 shows a superimposition of the three cores). The fused ring core of 2 has an essentially planar, linear-ladder shape in contrast to those of 3 and 4 which are severely puckered. In 4 it is butterfly shaped with a Li...Li hinge and benzotriazolyl wings whose planes lie at 50.9(4)° to one another, whereas in 3 one of the benzotriazole rings is almost perpendicular relative to the other at 86.9(4)°. These conformational contrasts are a consequence of the different coordinations at the Li junctions which connect the two benzotriazolyl frameworks, coupled with the different substitution patterns within these frameworks. In 3 the 1-phenyl substituents project mainly above the 6-5-6-5-6 fused ring core while the iBuAl(TM) units are situated below it. In 4 the deprotonated phenyl ring lies twisted with respect to the triazole ring to which it is attached [49.61(6)°], while the inclination of the non-deprotonated phenyl ring is marginally less at 44.2(6)°. As in 3, the iBuAl(TM) units are situated for the most part underneath the fused ring core which as a consequence of steric restraint must bend upwards in contrast to its near planar disposition in 2. To the best of our knowledge there are no previous examples of C-H aluminated benzotriazolyl-based crystal structures for direct comparison with 3 and 4. Benzotriazole derivatives substituted by phenol appendages at the 2H-N position have been structurally determined but these do not contain Al-C bonds as their Al centres interact only with N and O atoms. All the Al-C bonds within 3 and 4 fall in the narrow range 2.003(3) – 2.101(2) Å. The mean Al-C(i-Bu) bond length (2.0127 Å) is in the range of
other 4-coordinate Al centres bound to iso-butyl ligands in lithium aluminate structures (for example in ((Bu2Al(NPh)Li)[THF])[4]; mean 2.0085 Å; in ((PhC=O)Ni)2Li(TMP)(Bu2AlBu2)[4]; mean terminal bond length, 2.034 Å). Little variation is found in the Al-N(TMP) bonds within 3 and 4 with a mean length of 1.8716 Å. The Li-N(TMP) bonds span 2.028 – 2.094(4) Å, while the Li-N(triazolyl) bonds span 2.035(4) – 2.195(4) Å with longest bond [2.195(4) Å] belonging to the monomer connecting Li(2) - N(3) junction in complex 4.

**NMR spectroscopic studies of the lithium aluminate systems:** Full details and copies of the NMR spectra of the new compounds 3-4 are provided in the Supporting Information. It was not possible nor indeed considered worthwhile to assign the resonances for the LiTMP induced benzotriazolyl ring-opening product 2 because it was clearly formed as a mixture of crystallographically characterised 2, starting material and at least one unknown species. The homodimer 3 was characterised by both 1H and 12C NMR spectra recorded in C6D6 solution. Though the spectra are complicated and several resonances overlap, with the help of 2D NMR experiments (COSY, HSQC and HMBC experiments) we were able to assign all of the expected resonances. An interesting feature is that, although the structure in the solid state contains only one type of TMP ligand, the 1H resonances associated with it in solution consist of two χCHHN (at 1.57 ppm and 0.87-0.70 ppm), two χCHHN (at 1.12-0.98 ppm and 0.29 ppm) and two CH2 resonances (at 1.57 ppm and 1.12-0.98 ppm), a pattern which fits with the five TMP resonances observed in the 12C NMR spectrum. This suggests the TMP ligand is held rigidly in an asymmetrical arrangement, which is consistent with its bridging placement between the Li and Al centres in the crystal structure of 3. The highest frequency resonance in the 1H spectrum at 8.30 ppm can be assigned to the C-H on the benzo ring adjacent to the C-Al bond (its associated C-H resonance comes at 135.8 ppm). While we could not observe the C-Al resonance directly in the 12C NMR spectrum, we found it at 162.1 ppm through a 1H, 12C-HMBC experiment. Assigning and interpreting the 1H and 12C NMR spectra for the heterodimeric lithium aluminate 4 proved more of a challenge due to the inherently greater complexity of the data. The causes of this were a combination of the structural asymmetry of 4 and the existence of a slow chemical exchange process. We elected to interrogate the NMR signal assignments and the structure and dynamics for 4 using a battery of 1- and 2-dimensional NMR techniques. The slow-exchange (on the chemical shift NMR timescale) dynamics were notably evident through the occurrence of strong exchange cross-peaks in 2D [1H, 1H] EXSY NMR data and detected by 1H resonance lineshape variation over a range of temperatures either side of ambient. Critically, no exchange cross-peaks were detected in the aromatic resonance region of the 2D [1H, 1H] EXSY NMR data. This ruled out a see-saw chemical bond-making and breaking process within the heterodimer complex and led us to consider other alternative explanations. Exchange cross-peaks occurring solely within the aliphatic proton resonance range led us to surmise

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**Figure 5.** Molecular structure of [[4-R’-1-phenyl-1H-benzotriazole][1-(2-R’-phenyl)-1H-benzotriazole]], [R’=iBuAl(μ-TMP)Li]. 4. Thermal ellipsoids are displayed at 35% probability, hydrogen atoms, one disordered component of iBu and one n-hexane molecule of crystallization have been omitted for clarity. The dashed lines illustrate Li···C contact: Li1···C12, 2.695(4). Selected bond lengths (Å) and bond angles (°): L1-N2, 2.064(4); L1-N4, 2.048(4); L1-N6, 2.053(4); L2-N3, 2.195(4); L2-N7, 2.083(4); L2-N8, 2.094(4); A1-C15, 2.101(2); A1-N4, 1.9659(19); A1-C22, 2.015(2); A1-C26, 2.020(2); A2-C44, 2.053(4); A1-C12, 2.195(4); A1-C40, 2.083(4); A1-C49, 2.094(4); Al-C12, 2.034 Å with longest bond [2.195(4) Å] belonging to the monomer connecting Li(2) - N(3) junction in complex 4.

**Figure 6.** Superimposition of the 6-5-6-5-6 fused ring cores of (a) complex 2, (b) complex 3 and (c) complex 4. Blue ellipsoids represent nitrogen atoms and yellow ellipsoids represent lithium atoms.
that slow conformational flexibility was the most likely source of these observations. To quantify the process, a series of temperature-dependent 2D $^1$H, $^1$H EXSY NMR data sets were acquired for 4 in the temperature range 263 ≤ T ≤ 323 K at mixing times varying in the range 5 ≤ $\tau_m$ ≤ 200 ms (see Supporting Information). The data gave activation enthalpy $\Delta H^\ddagger = 71 \pm 5$ kJ mol$^{-1}$, Gibbs free energy of activation at 298 K, $\Delta G^\ddagger_{298} = 67 \pm 5$ kJ mol$^{-1}$ and activation energy $E_a = 75.4 \pm 0.4$ kJ mol$^{-1}$, values that are consistent with conformational rearrangement dynamics rather than bond breaking and making. To explore more closely the conformations and to firmly establish the nature of the solution-phase species representing 4, we made strides towards assigning the data to allow key through-space information, acquired via 2D $^1$H, $^1$H NOESY NMR data, to be interpreted. For this we acquired NMR data on a sample of 4 in toluene-$d_8$ cooled to 213 K, at which temperature the exchange process was sufficiently arrested to ensure that the NOESY data were dominated by genuine NOEs, exchange cross-peaks being suppressed below the level of detection. With the assignment of the aromatic proton resonances confirmed at this temperature, we used these and the representative crystal structure of 4 to determine the most likely NOE cross-peak assignments.

The complexity of the data meant that specific, unambiguous assignments were not easily obtained from the NOE data for 4. However, it was possible to identify NOEs that were associated specifically with one of the four methyl groups (methyl-1) of the TMP moiety designated here as TMP(b) whose nitrogen centre is identified as N4 in Figure 5. The NOEs, shown in Figure 7 as white dotted lines, occurred between the methyl-1 protons and those protons of the aromatic rings lying in close proximity as indicated in the Figure and provided in Table S2 of the Supporting Information.

As shown (Supporting Information Fig S7), the proton resonance arising from the NMR-assigned TMP(b) methyl group has two identities at low temperature related by chemical exchange and whose chemical shift separation $\Delta \delta = 1.0$ ppm. The conformation shown in Fig. 7, allows the NOEs arising from the proton resonance of the methyl group at $\delta^H = 0.6$ ppm to be assigned. The relatively low chemical shift is consistent with magnetic shielding; here the TMP(b) methyl group is located above an aromatic ring plane as shown in Fig. 7. By extension, it follows that the same methyl group can also be located in a less shielded environment ($\delta^H = 1.6$ ppm) through conformational exchange. By implication, this suggests that the conformational flexibility revealed by the NMR data of 4 in solution is associated with movement within the structure that relocates TMP(b) methyl-1. This could be readily achieved through a conformational change in the 7-membered ring containing the TMP(b) nitrogen centre (N4 in Figure 5) which would be consistent with the thermodynamic quantities obtained by exchange data analysis. This is more likely than the equivalent conformational rearrangement of the 6-membered ring containing the TMP(a) nitrogen centre (N8 in Figure 5). Such a process would require the climbing of higher energy than the one determined by our exchange data analysis. Other large proton chemical shift changes were identified that arise by the same process, but unambiguous assignment of these signals was not possible. NMR resonances from groups closely related to TMP(b)-methyl-1 were also affected in a similar manner on the basis of which it is proposed that the dynamic event observed for 4 in solution is concerted motion of the 7-membered ring N4-Li1-N2-N1-C7-C12-Al1 (Figure 5). Consideration of these same NOE data confirms that the molecule exists as the intact heterodimer when in solution. The simultaneous appearance of NOEs to TMP(b)-methyl-1 protons from the aromatic proton resonances of both of the 6-membered aromatic rings attached to each benzotriazole shows that the two halves of the complex are in close proximity to the identified TMP(b) methyl group, allowing NOEs to evolve from both sides of the molecule simultaneously.

Deprotonization via a lithium-zinc base mixture: The logical starting point for this part of our investigation was to follow Mongin’s original literature preparation (see Scheme 1 in the introduction). A quantitative (99%) yield of the 4,2'-diido derivative of 1 was reported on mixing 1 with three equivalents of TMP(H)/nBuLi (i.e., in situ LiTMP) and one equivalent of ZnCl$_2$-TMEDA in THF solution at room temperature for two hours followed by quenching with three equivalents of the
electrophile iodine. From other literature on this base mixture it is certain that a lot of chemistry is concealed in the reaction coordinate between reactant 1 and the final 4,2'-diiodo product. Intuitively one might expect that the base mixture would undergo a complete metathesis to generate the ate "LiZn(TMP)₂" especially as ate reagents have been fundamental to many of the recent advances made in metalation chemistry. However, DOSY NMR data in the reaction solvent (THF) point to an alternative partial metathesis terminating at neutral, weakly basic Zn(TMP)₂, with "Li₂Zn(TMP)₂," forbidden on sterics, leaving LiTMP (or a co-complexed/solvated version) as the only starting base in the mixture strong enough to C-H deprotonate a benzotriazole. Following deprotonation of presumably the benzo-4 position (using less base Mongin found that this regioselectivity was greatly favoured), the lithiated benzotriazole could undergo transmetalation with Zn(TMP)₂ to make a monoazincated benzotriazole and LiTMP (Scheme 4). In turn, this second LiTMP can deprotonate the ortho-2' position of the phenyl substituent to form the dideprotonated, mixed lithium-zinc benzotriazole. Two possible scenarios can be envisaged from here to reach the diiodo product. Either this mixed lithium-zinc species could be iodinated directly or alternatively its Li centre could undergo a fast second transmetalation to form a zinc dianionic benzotriazolate and LiTMP prior to iodination. Endeavours to gather experimental evidence that would shed light on the above possibilities have proved frustratingly challenging. We have made several attempts to obtain crystalline or solid forms of possible metalo intermediates by following this literature procedure but omitting the final iodination step. The excellent solubility of the metal intermediate/s in THF solution has hampered this quest. Even cooling the reaction solution to -33°C failed to induce the deposition of any solid. This is noteworthy since the mixture contains lithium chloride, at least formally, though the aforementioned DOSY study proposed the possibility that this metathetical coproduct might not be free but is incorporated within the active lithiating base of formula LiTMP·2LiCl·TMEDA. With few exceptions it is notoriously difficult to crystallise metalo complexes from THF solutions of reaction mixtures containing salt-supported organometallic bases. In this case we enjoyed a modicum of success as setting the reaction solution aside at room temperature for about a month produced a tiny crop of very small crystals which, when subjected to an X-ray crystallographic study, proved to be only weakly diffracting. The poorly diffracting nature of the crystals is undoubtedly correlated to large amounts of disorder in two separate major structural features. Disordered, THF solvated Li⁺(C₆H₅O)⁻ centres (that is metallated THF anions) are present at approximately half of the possible sites at the core of an ordered metallocycle while disordered THF present as "solvent of crystallisation" could only be treated by using the SQUEEZE routine implemented within PLATON to remove it from the structural model. Despite the poor quality of the data, the connectivity of the main structural feature is well-defined, comprising a hexameric ring of alternating Zn atoms and dideprotonated benzotriazole ligands with zinc bridging between the 4-position of one ligand and the 2'-position of another (Figure 8), leading us to presume an approximate formula of [(C₆H₅N₂)(C₆H₅Zn)₃(Li)₃(C₆H₅O)₃(THF)₃]·nTHF, 5. Though there is no evidence to say this species represents the bulk product of the reaction its 4,2'-dideprotonation matches up well with the 4,2'-diiodination observed by Mongin following the quenching step. Also it appears to concur with the second scenario mentioned above whereby zinc stabilises the two dideprotonated benzotriazole centres to release an equivalent of LiTMP, which, in turn, can deprotonate THF solvent molecules to generate C₆H₅O⁻ anions. Deprotonative metalation of THF usually forces its ring opening and decomposition but significantly α-
deprotonated THF anions (C₆H₄O⁻) can exist intact under bimetallic (Na-Zn or Li-Al) synergetic stabilisation so a similar Li-Zn effect could be in operation here. It is also pertinent to note that zinccyclic structures are known. For example, our earlier work probing the alkali-metal-mediated zirconation of N,N-diethyl-thiophene-2-carboxamide by the sodium zircono reagent [(TMEDA)Na(TMP)(Bu)Zn(Bu)] produced a nice example of a 16-membered (ZnCCC) ring. This smaller zinccycle is related to that in 5 as the ring arises from double deprotonation of the thiophene unit (compare that of the benzotriazole unit in 5), but in contrast it was produced via direct zirconation (Zn-H exchange) unlike the indirect lithiation/zinc-trapping mechanism at work in the synthesis of 5 (or the lithiation/aluminium-trapping in 3 and 4).

Conclusions

The aim of this study was to gain some structural comprehension of the metalation chemistry of heterocyclic substrates by complicated multicomponent organometallic bases. Studies of this type are essential to unravel the complexity of these base mixtures, which can often outperform single component bases such as LiTMP. Though excellent work by Mongin et al had quantified the yields and selectivities of the metalation of 1-phenyl-1H-benzotriazole with sequential in situ ZnCl₂/TMEDA/LiTMP and iodine protocols, no information was gleaned on the reactive metalo intermediates, which mediate the iodine-hydrogen exchange reaction. As commonly found in mixed-metal base systems, extracting quality crystalline samples from bulk THF solutions is no straightforward task and that synthetic routes were prepared according to literature procedures. LiTMP[3] and Bu₃Al(TMP) were prepared according to literature procedures. NMR spectra were recorded on a Bruker AVANCE 400 NMR spectrometer, operating at 400.1 MHz for 1H, 155.5 MHz for 13Li, 102.4 MHz for 27Al and 100.6 MHz for 13C. 1H and 13C NMR spectra were referenced to the appropriate solvent signal, 1Li NMR spectra were referenced against LiCl in D₂O at 0.00 ppm and 27Al NMR spectra were referenced against AlCl₃ in D₂O at 0.00 ppm. Elemental analyses of the compounds 3-4 were carried out using a Perkin Elmer 2400 elemental analyser.

Experimental Section

General procedures: All reactions and manipulations were performed under a protective atmosphere of dry pure argon gas using standard Schlenk tube or glovebox techniques. Solvents were dried by heating to reflux over sodium benzophenone ketyl and distilled under nitrogen prior to use. Deuterated NMR solvents were degassed and stored over 4 Å molecular sieves to use. 2,2,6,6-tetramethylpiperidine THF (H) was purchased from Merck KGaA, and stored over activated 4 Å molecular sieves. nBuLi (1.6 M in n-hexane solution), AlCl₃ and Bu₃AlCl were purchased from Aldrich and used as received. 1-Phenylbenzotriazole was purchased from Aldrich and used as received. THF and methylcyclohexane were purchased from Aldrich and used as received. THF and methylcyclohexane were degasified and stored over 4 Å molecular sieves.

Synthesis of [(4-R-1-(2-lithiophenyl)-1H-benzotriazole-3THF)][R = 2-C₆H₄(Pn)N][Li], 2. In a Schlenk tube, 1-phenyl-1H-benzotriazole (195 mg, 1 mmol) was suspended in n-hexane (15 mL) and LiTMP (294 mg, 2 mmol) was then added. The resulting yellow suspension was stirred for 2 h at room temperature. THF (2 mL) was then added and the reaction mixture was heated until a dark solution was achieved. The solution was placed in a freezer operating at -15 °C. After a few days, crystals of 2 were obtained (132 mg, 0.10 mmol, 10%). The 1H NMR spectrum of the reaction shows that the metallation is not clean and that there is starting material. 1H NMR (400.13 MHz, δ in THF, 300 K): 7.90-7.81 (m), 7.68-7.47 (m), 7.32 (m), 7.08 (m, 6.89 (d, 2H), 6.70 (t, 1H), 3.60 (m, THF), 1.77 ppm (m, THF). We observe by NMR spectroscopy a small amount of 1-phenyl-1H-benzotriazol-3THF in the reaction. 1H-NMR (400.13 MHz, δ in THF, 300 K) of C₆H₄N⁺: δ 8.08 (‘d’, 1H, 4 = 8.4 Hz, J = 0.9 Hz, Hz), 7.86 (m, 3H, Hz, Hz), 7.83 (m, 2H, Hz, Hz), 7.55 (dd, 1H, J = 8.0 Hz, J = 6.9 Hz, J = 1.1 Hz, Hz), 7.40 (tt, 1H, J = 6.9 Hz, J = 1.1 Hz, Hz), 7.41 ppm (dd, 1H, J = 8.2 Hz, J = 7.0 Hz, J = 1.0 Hz, Hz).

Synthesis of [(4-R'-1-phenyl-1H-benzotriazole-3THF)][R' = (Bu)₃Al(µ- TMP)Li], 3. In a Schlenk tube, 1-phenyl-1H-benzotriazole (195 mg, 1 mmol) and LiTMP (147 mg, 1 mmol) were suspended in n-hexane (15 mL) to give a pale yellow suspension. A solution of Bu₃Al(TMP) (281 mg, 1 mmol) in n-hexane (3 mL) was added immediately to give a beige suspension, which was stirred for 2 h at room temperature. The
suspension was filtered and the solid was washed with n-hexane (3 x 4 mL). Crystals of 3 suitable for an X-ray diffraction study were obtained by heating a suspension in n-hexane/methylcyclohexane to get a solution, and crystals grew when the Schlenk tube was placed in a refrigerator operating at 4 °C (319 mg, 0.30 mol%, 30%).

$^{13}C$ NMR (400.13 MHz, CD$_2$Cl$_2$, 300 K): $\delta$ 87.3 (q, C$_{20}$), 86.5 (q, C$_{22}$), 2.25 (v br s, 2H, CH$_2$Bu), 1.57 (s, 7H, CH$_2$TMP + γCH$_2$TMP), 1.46-1.29 (overlapped br s + br d, 12H, CH$_2$Bu), 1.12-0.98 (br m, 8H, CH$_2$TMP + βCH$_2$TMP), 0.87-0.70 (overlapped br s, 5H, γCH$_2$TMP + CH$_2$Bu), 0.29 ppm (v br s, 2H, CH$_2$TMP). $^1$H NMR (155.5 MHz, CD$_2$Cl$_2$, 300 K): $\delta$ 2.50 ppm. $^{13}$C$^{[1]}$H NMR (100.6 MHz, CD$_2$Cl$_2$, 300 K): $\delta$ 153.8 (C$_{2}$-benzotriazole), 135.9 (C$_{2}$-Ph), 135.8 (C$_{1}$), 131.6 (C$_{3}$-benzotriazole), 130.9 (C$_{5}$), 130.2 (C-Ph), 124.6 (C-Ph), 106.0 (C$_{4}$), 53.2 (C$_{13}$-TMP), 43.5 (TMP), 37.7 (Bu), 30.3 (CH$_2$Bu), 30.0 (br s, TMP), 29.2 (v br s, CH$_2$Bu), 26.1 (TMP), 27.3 (Bu), 18.6 ppm (TMP). We only observe by 2D NMR spectroscopy the Al-C$_{2}$H$_3$. $^{11}$C-HMBCC NMR: $\delta$ 162.1 ppm (Al-C$_{2}$H$_3$). We can observe by NMR spectroscopy a small amount of methylcyclohexane from the crystallization. $^{13}$C NMR (400.13 MHz, CD$_2$Cl$_2$, 300 K) of C$_{2}$H$_4$: $\delta$ 1.70-0.61 (m, 5H), 1.30-1.15 (m, 4H), 0.86 ppm (d, 5H), $^{13}$C NMR (400.13 MHz, CD$_2$Cl$_2$, 300 K) of C$_{2}$H$_6$: $\delta$ 35.6, 32.9, 26.7, 26.6, 23.0 ppm.

Elemental analysis calc. (%) for CuH$_{13}$O$_{2}$AlN$_{2}$Li: C, 73.11; H, 9.78; N, 10.66; found: C, 73.15; H, 9.72; N, 11.21.

**Synthesis of [[4-R'-1-phenyl-1H-benzotriazole][1-(2-R''-phenyl)-1H-benzotriazole]].** [R' = (Bu)$_2$Al/μI]TLi, 4. In a Schlenk tube, THF (0.50 mL, 3 mmol) was added in THF (7 mL) and then reacted with BuAlI (562 mg, 2 mmol) in n-hexane (3 mL) added to give a beige suspension, which was stirred for 2 h at room temperature. The suspension was filtered and the solid was washed with n-hexane (3 x 4 mL). Crystals of 4 suitable for an X-ray diffraction study were obtained by heating a suspension in n-hexane/methylcyclohexane to get a solution, and crystals grew when the Schlenk tube was placed in a freezer operating at -15 °C (357 mg, 0.34 mol%, 34%).

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**Keywords:** aluminium • lithium • metallaletion • structure elucidation • triazoles • zinc

**References**


Sedate lithiation: Conventional lithiation of 1-phenyl-1H-benzotriazole destroys the heterocycle by ring opening, N₂ extrusion and C-C bond formation, but adding the trapping agent iBu₂AlTMP transforms the initial unstable C-Li to stable C-Al bonds sedating the sensitive carbanion to keep the triazole scaffold intact. This trans-metal-trapping was established through isolation and structural characterization of bimetallic “intermediates”.