

Cite this: *Chem. Sci.*, 2021, 12, 15318

All publication charges for this article have been paid for by the Royal Society of Chemistry

Intermolecular CDC amination of remote and proximal unactivated C_{sp}³–H bonds through intrinsic substrate reactivity – expanding towards a traceless directing group†‡

Suresh Rajamanickam,^a Mayank Saraswat,^b Sugumar Venkataramani^{*b} and Bhisma K. Patel^{†a}

An intermolecular radical based distal selectivity in appended alkyl chains has been developed. The selectivity is maximum when the distal carbon is γ to the appended group and decreases by moving from $\gamma \rightarrow \delta \rightarrow \epsilon$ positions. In –COO– linked alkyl chains, the same distal γ -selectivity is observed irrespective of its origin, either from the alkyl carboxy acid or alkyl alcohol. The appended groups include esters, N–H protected amines, phthaloyl, sulfone, sulfinamide, nitrile, phosphite, phosphate and borate esters. In borate esters, boron serves as a traceless directing group, which is hitherto unprecedented for any remote C_{sp}³–H functionalization. The selectivity order follows the trend: 3° benzylic > 2° benzylic > 3° tertiary > α to keto > distal methylene ($\gamma > \delta > \epsilon$). Computations predicted the radical stability (thermodynamic factors) and the kinetic barriers as the factors responsible for such trends. Remarkably, this strategy eludes any designer catalysts, and the selectivity is due to the intrinsic substrate reactivity.

Received 9th August 2021
Accepted 26th October 2021

DOI: 10.1039/d1sc04365j

rsc.li/chemical-science

Introduction

Functionalization of inert C_{sp}³–H bonds with a high degree of selectivity is one of the most challenging yet desirable avenues in organic synthesis. In living systems, the enzyme cytochrome P₄₅₀ uses an intricate binding pocket to achieve this transformation in appended alkyl chains with precise selectivity onto a particular substrate.¹ Chemists have successfully functionalized C_{sp}³–H bonds adjacent to π -systems,² heteroatoms^{2b,3} or using directing groups.⁴ Lately, chemists have developed designer metal catalysts or molecular recognition units to functionalize C_{sp}³–H bonds of the same type without the assistance of directing groups.⁵ The catalysts/oxidants achieve selectivity through electronic, steric and stereo-electronic factors inherited in the substrates; though it is quite often that the examined substrates are electronically biased.²

Several strategies have emerged for the non-directed remote C_{sp}³–H functionalization of aliphatic compounds. For instance,

the methine and methylene C–H bonds have been selectively oxidized using Fe(PDP)/H₂O^{6a–c} and NO₂[Fe–bTAML]/m-CPBA^{6d} in complex substrates. An electrochemical method demonstrates the oxyfunctionalization of electron-rich methylene carbon centers at remote positions.^{7a} Intermolecular remote C_{sp}³–H bromination,^{7b} chlorination^{7c} and xanthylation^{7d} have been accomplished utilizing *N*-halo and *N*-xanthylamides under irradiation of visible light Zhdankin's azidoiodine method. Indeed, it has been used in association with an Fe(II)/i-Pr-PyBox-catalyst for the selective azidation of electron-rich, remote methine over methylene and methyl centers.⁸ Subsequently, azidation is achieved using a Mn-catalyst,^{9a} and radical trifluoromethylthiolation (SCF₃) is mediated by AgSCF₃ (ref. 9b and c). The site selectivity of substrates can be altered by fine-tuning the catalytic systems.^{9d} Synthetic chemists have made significant strides in the field of remote C_{sp}³–H functionalization; however, this area remains as exciting as ever, and several reactions are yet to be discovered and generalized.

Compounds containing C–N bonds find extensive utility in diverse areas of chemistry, and several reaction methods have emerged. Historically, the Hofmann–Löffler–Freitag (HLF) reaction is employed for intramolecular amination, involving the C_{sp}³–H bond.^{10a,b} The modern HLF reaction is achieved *via* electrochemical^{11a} or a combination of photo and electrochemical processes.^{11b} Currently, the direct conversion of the C–H bond to the C–N bond is demonstrated by nitrene chemistry,¹² directing groups,^{4a,b,13a,b} and cross dehydrogenative coupling (CDC) strategies.^{2a,b,g–i,14} Performing intermolecular aminations *via* nitrene

^aDepartment of Chemistry, Indian Institute of Technology Guwahati, North Guwahati Address, Assam-781039, India. E-mail: patel@iitg.ac.in

^bDepartment of Chemical Sciences, Indian Institute of Science Education and Research (IISER) Mohali, Sector 81, Knowledge City, Manauli, SAS Nagar, 140306, India. E-mail: sugumarv@iisermohali.ac.in

† Dedicated to Professor Srinivasan Chandrasekaran on the occasion of his 70th birthday.

‡ Electronic supplementary information (ESI) available. CCDC 2077948 and 2070229. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d1sc04365j

chemistry is a tedious task as the metallo-nitrene intermediates are unstable, and the scope is limited mainly to electronically biased benzylic or tertiary sites.^{12d,g-i} Strategies based on directing groups require two additional steps during the synthesis—pre-functionalization of the starting materials followed by deprotection. Moreover, the reductive elimination step in the catalytic cycle is hampered by the high binding affinity of the newly introduced amino moieties.¹⁵ The CDC predominantly works for activated C–H bonds that are adjacent to heteroatoms,^{3f,g} or π -systems.^{2b,c,g-i} Intermolecular amination *via* CDC remains hitherto undiscovered in substrates that can engage remote, unactivated methylene sites in the coupling process. Moreover, the question arises, whether the selectivity in a structurally diverse substrate is a result of steric, or stereo-electronic factors or the outcome of the catalyst and oxidant.

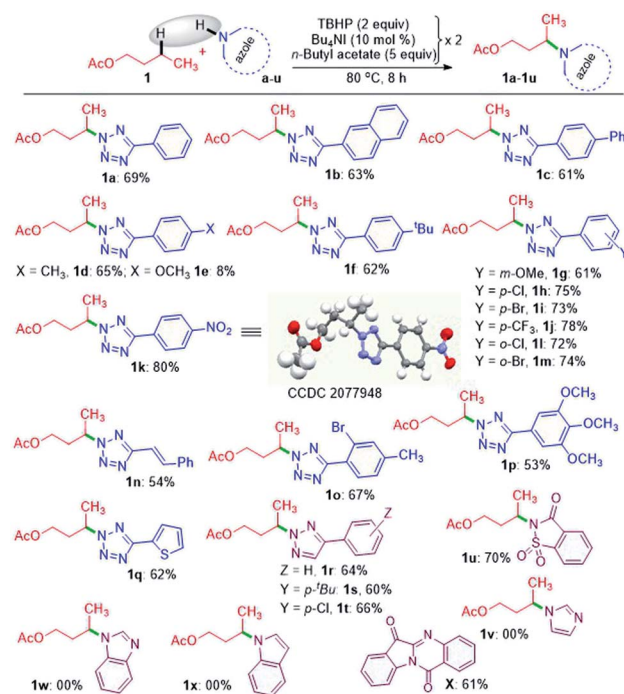
The tetrazole moiety is highly admired among N-heterocyclic compounds: it acts as a bioisostere of carboxylic acid surrogates due to similar pK_a 's and renders high metabolic stability.¹⁶ For instance, biphenyl tetrazoles are key intermediates in the production of multibillion-dollar angiotensin II receptor antagonists, a class of drugs for treating high blood pressure known as sartan drugs; pemirolast—an antiallergic drug; and azosemide—a diuretic. Also, many of the cephalosporine derivatives possess tetrazole units (Fig. S1, see ESI†).¹⁶ The $[Mn^{III}(ClPC)]^{12d}$ and $Rh_2(esp)_2$ (ref. 12g–i) catalysts can efficiently differentiate large substrate variation which is applied to complex molecules, such as terpenes, steroids, alkaloids, peptides, and lactams.^{12d,g-i} In most cases, the functionalization is limited to a position in electronically biased substrates, such as the tertiary or benzylic.^{12d,g-i,n} Thus, it is challenging to achieve amination at undirected C–H bonds with predictable site-selectivity without a *de novo* approach. Can a remote and rational C_{sp^3} –H amination strategy be developed based on the intrinsic substrate reactivity that can be expanded to a traceless directing group? If so, what are the possible determining factors?

Results and discussion

A nitrogen centered radical (NCR) is generated by reacting tetrazole with a combination of iodide and peroxide or *tert*-butyl hydroperoxide (TBHP)/tetrabutylammonium iodide (TBAI).^{17a,b} In the absence of any other radical coupling partners, the generated NCR abstracts a methyl (–CH₃) group from the TBHP giving a N²-methylated tetrazole exclusively.^{17a} Taking cues from this and the non-directed remote C_{sp^3} –H bond functionalization strategies,^{6–8,9a} particularly Wu's and Tang's trifluoromethylation,^{9b,c} we envisaged the incorporation of N-heterocycles (tetrazoles) at an unactivated remote methylene C_{sp^3} –H position. With this idea in mind, we set up a reaction at 70 °C between 5-phenyl-2H-tetrazole **a** and *n*-butyl acetate **1** in the presence of Bu₄NI and the oxidant, TBHP (5–6 M in decane). As anticipated, the site-selective γ -aminated product **1a** was achieved in 21% yield. Since the oxidant, TBHP, is dissolved in *n*-decane, the *in situ* generated tetrazole radical reacted with it affording an indiscriminate mixture of aminated *n*-decane products, which is quite unsolicited. This preliminary result was quite encouraging, as the remote CDC of **1** with **a** evades

any directing group or designer catalyst. Next, the optimization parameters were scrutinized by varying the reaction temperature, oxidant, catalyst, and its loading to improve the reaction efficacy. After extensive exploration (Table S1, see ESI†), it was revealed that subjecting **a** (0.5 mmol) to two iterative additions of 10 mol% of Bu₄NI, two equivalents of oxidant (aq TBHP), and 330 μ l of **1** at the beginning and after an interval of 2.5 h at 80 °C afforded product **1a** in 69% yield after 8 h (Table S1, see ESI†).

A range of substituted tetrazoles **a–q** were subjected to the optimized conditions, and the results are summarized in Scheme 1. Tetrazole **a** and π -conjugated tetrazoles **b** and **c** coupled with **1** giving products (**1a–1c**) in modest yields. The key feature here is the occurrence of exclusive amination at the γ -position of **a**, leading to a single regioisomer. The regioselectivity can be attributable to a slight kinetic preference in the ease of formation of the radical center in **1** at γ over its α position. {For further details, see the Computational studies section (Fig. 2)}. The phenyl ring of aryl tetrazole possesses either electron-donating **d–g** or electron-withdrawing **h–m** substituents smoothly coupled with **1** to afford their γ -tetrazolyl products **1d–1m** in moderate to good yields (Scheme 1). When a π -conjugated tetrazole **n** reacted with **1**, it gave γ -aminated product **1n**. Furthermore, aryl tetrazoles **o** and **p** having multiple substituents afforded their respective products **1o**, and **1p**. A thiophenyl tetrazole **q** provided its product **1q** in modest yield. The regioselectivity of the product was ascertained by X-ray crystallographic analysis, as shown in Scheme 1. Recently the Patel group achieved similar N² alkylation using TBAI/TBHP



Scheme 1 Substrate scope for the intermolecular amination of *n*-butyl acetate. ^a Reaction conditions: azole (0.5 mmol), *n*-butyl acetate (5 equiv. \times 2), Bu₄NI (10 mol% \times 2) and aq TBHP (2 equiv. \times 2) at 80 °C for 8 h in an inert atmosphere. ^b Isolated yields.



combination in the presence of various organic peroxides.^{17a} Computations revealed the determining factors for exclusive N² regioselectivity such as the high spin density at the N² center, low transition state barriers, and better thermodynamic stability of the products.^{17a}

Next, we sought the possibilities of remote amination of **1** with other azoles and N-heterocycles, such as triazoles **r-t**, saccharin **u**, imidazole **v**, benzimidazole **w** and indole **x** (Scheme 1), while aryltriazoles **r-t** and saccharin **u** reacted successfully giving γ -functionalized products **1r-t** and **1u**; imidazole **v** and benzimidazole **w** failed to react (Scheme 1). However, indole **x** provided tryptanthrine **X** in 61% yield, which is well documented.¹⁸ The failure to undergo similar amination for **v** and **w** is any one or a combination of the factors, namely (i) delocalization and stabilization of the radical with no or minimal radical character at the nitrogen center, (ii) significant activation energy, and (iii) unfavourable thermodynamic stability of the product.^{17a}

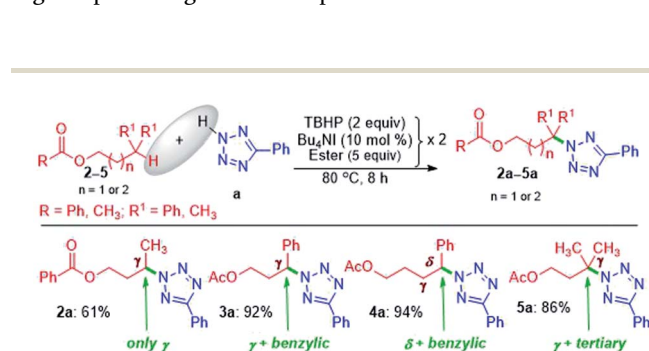
The successful γ -amination of ester **a** encouraged us to test a similar strategy with an aromatic ester **2** which provided a γ -tetrazolyl product **2a** in 61% yield (Scheme 2). This result suggests that both methyl and phenyl groups have identical or no influence when present on the carbonyl side of the ester, and the fate is dictated by the ester –COO– moiety. However, after mutually swapping the position of the phenyl and the methyl group as in **3**, the γ -tetrazolyl product **3a** was isolated in a very high yield (92%). The phenyl group, which is now present at the γ -position, is electronically biased as it is also benzylic; thus, there is a positive influence on the selectivity and the yield. Further, an exclusive δ -functionalization without a trace of γ -product in ester **4a** confirmed the complete electronic biasness.^{12d} Further, an ester having a γ -methine hydrogen (3° C–H), as in **5**, provided an exclusive γ -tetrazolyl product **5a**. Thus, the current strategy evades designer catalysts, and the selectivity is due to the inherent substrate reactivity.

Now, a further query arose: other than the benzylic position, does this protocol always provide γ -selectivity, or is this amination feasible beyond this position, particularly for substrates having longer alkyl chains? With this objective in mind, ester **6** possessing γ and δ positions was tested, which provided an inseparable mixture of γ and δ aminated products **6'a** and **6a** in a combined yield of 73% in the ratio of 1 : 4.8 (Scheme 3). The higher percentage of the δ -product **6a** is due to the better

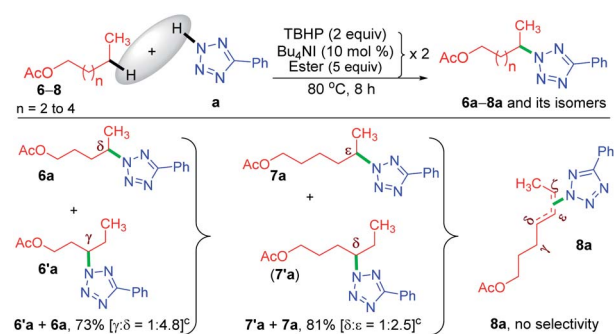
stability of the distal δ -carbon center compared to the γ -centered radical. In product **1a**, the selectivity is at the γ -center and in **6a** it is at the δ -center, and both happen to be at the distal carbon (secondary). This raises further inquisitiveness: whether the selectivity is dictated by the distance from the ester moiety (γ or δ position) or only by the distal methylene carbon. When hexyl acetate **7** was subjected to identical reaction conditions, it provided an inseparable mixture of δ and ϵ aminated products, **7'a** and **7a** in the ratio of (1 : 2.5) in a combined yield of 81% (Scheme 3), suggesting a preferential distal selectivity. Undoubtedly, the ϵ -selective product **7a** was the major one, but the degree of distal selectivity decreased considerably. No preferential selectivity was observed for *n*-heptyl acetate **8**, as it provided a mixture of δ , ϵ , and ζ aminated products whose exact ratio could not be ascertained. The distal selectivity is governed by the electronic influence imparted by the –COO– group along the alkyl chain.^{7a-d,9b,c}

In this regard, we computationally estimated the C–H bond dissociation energies (BDEs) for the compounds having more than four carbons in the alkyl chain (C-5 ester **6** and C-6 ester **7**).¹⁹ Surprisingly, the values are nearly the same for the distal positions. The observed lower C–H BDEs and their difference (0.1–0.2 kcal mol^{–1}) are not only consistent with the isolated mixture of products but also the difference in the stability [inferred from the comparison of C–H BDEs at all the centers] that can be accountable for the distal selectivity.²⁰

Now, the query arises about the selectivity in short-chain esters, say C-2 esters **9** and **10** possessing no γ -carbon, and a C-3 ester having a terminal but a primary γ -carbon. With this objective in mind, we treated **9** and **10** with **a**. The amination took place only at the α -position to yield products **9a** and **10a** in lower yields with no traces of β products (Scheme 4). Despite the superior stability of the α -C radical, the reduced yields of products **9a** and **10a** are due to the competitive *N*-methylation of tetrazole.^{17a} For tetrazoles **h** and **i**, the products **9h** and **9i** were isolated in their pure form. However, tetrazole **d** provided an inseparable mixture of α -aminated product **9d** and 2-methyl-5-(*p*-tolyl)-2H-tetrazole **dm**^{17a} in 3.4 : 1 ratio. The C-3 ester **11** provided two isomeric products, namely α -aminated (**11a**, 35%)

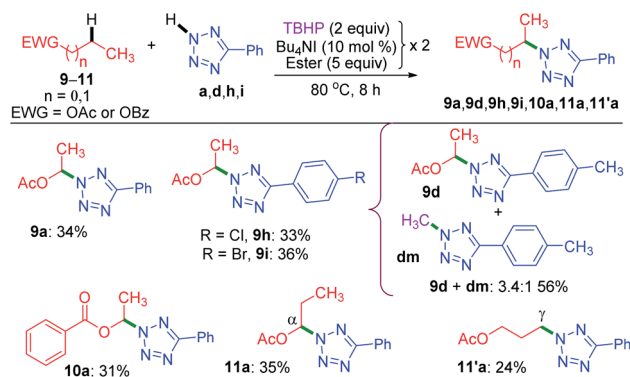


Scheme 2 Substrate scope of amination at remote methylene, benzylic and tertiary-methine sites. ^a Reaction conditions: azole (0.5 mmol), ester (5 equiv. \times 2), Bu₄NI (10 mol% \times 2) and aq TBHP (2 equiv. \times 2) at 80 °C for 8 h. ^b Isolated yields.



Scheme 3 Substrate scope for the amination of alkyl acetates. ^a Reaction conditions: 5-phenyl-2H-tetrazole (0.5 mmol), ester (5 equiv. \times 2), Bu₄NI (10 mol% \times 2) and aq TBHP (2 equiv. \times 2) at 80 °C for 8 h. ^b Isolated yields. ^c Products obtained as an inseparable regioisomeric mixture and the ratio determined by ¹HMR analysis.



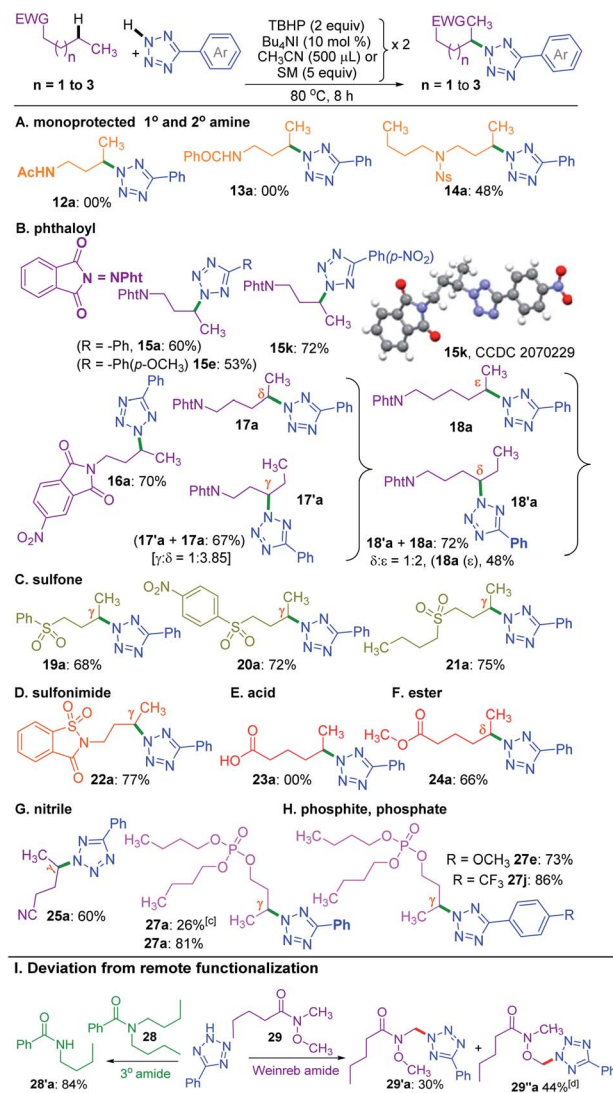


Scheme 4 Substrate scope for intermolecular amination of esters.^a Reaction conditions: aryl tetrazole (0.5 mmol), ester (5 equiv. \times 2), Bu₄NI (10 mol% \times 2) and aq TBHP (2 equiv. \times 2) at 80 °C for 8 h. ^a Isolated yield.

and γ -aminated (**11'a**, 24%) products without giving any β -isomer. Both the α - and γ -functionalized products **11a** and **11'a** might be forming *via* a radical pathway due to the inductive (–I) influence of the ester group. An alternative pathway for the α -functionalized product **11a** originating *via* the formation of an oxocarbenium is less likely.^{21a}

So far, the substrates examined are ester appended alkyl moieties that dictate the site-selective amination by intrinsic reactivity. Other electron-withdrawing groups, such as amide, phthaloyl, ketone, sulfone, cyano, phosphate, electron-deficient heterocycles, and sulfonimide, were tested to establish similar influences (Scheme 5). Primary alkyl amides **12** and **13** both failed to provide any product. Interestingly, a nosyl protected dibutyl amine **14** provided mono γ -tetrazolyl product **14a**, which suggests the detrimental influence of the free N–H group. This fact was further confirmed when a phthaloyl protected *n*-butyl amine **15** coupled with **a** and provided the product **15a** in 60% yield. Additionally, substrate **15** reacted with **e** and **k**, and delivered exclusive γ -tetrazolylated products **15e** and **15k**. The regio- and site-selectivity of the product was confirmed by X-ray crystallography (Scheme 5). In a similar vein, 5-nitrophthalic anhydride protected *n*-butyl amine **16** delivered product **16a**.

To determine the supremacy of the distal methylene selectivity, the chain length was increased from C-4 to C-5. A phthaloyl protected *n*-pentyl amine **17** gave an inseparable mixture of γ : δ isomeric products **17'a** and **17a** in the ratio of 1 : 3.8. On the other hand, for an *n*-hexyl amine derivative **18**, the δ : ϵ selectivity was found to be 1 : 2. These results reconfirmed the dominance of distal methylene selectivity. Thus, the presence of electron-withdrawing moieties such as esters and protected secondary amines is all capable of dictating the regioselective amination in an alkyl chain at the distal methylene carbon. Interestingly, aryl butyl sulfones **19** and **20** also provided exclusive γ -aminated products **19a** and **20a**, whereas a dibutyl sulfone **21** possessing two butyl chains yielded only a mono γ -aminated product **21a** (Scheme 5). An *n*-butyl saccharin **22** possessing a strong electron-withdrawing sulfonimide group provided product **22a**. Unprotected alkyl carboxylic acid, such as hexanoic acid **23**, failed to deliver any product,



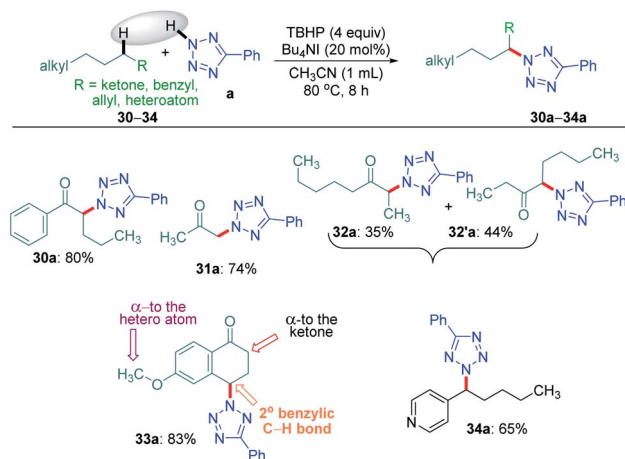
Scheme 5 Substrate scope for the amination of various electron withdrawing groups. ^a Reaction conditions: (A–D), aryl tetrazole (0.5 mmol), substrates **12**–**22** (0.5 mmol), Bu₄NI (10 mol% \times 2), aq TBHP (2 equiv. \times 2) and CH₃CN (500 μ L \times 2) at 80 °C for 8 h. (E–H), azoles (0.5 mmol), substrates **23**–**27** (5 equiv. \times 2), Bu₄NI (10 mol% \times 2) and aq TBHP (2 equiv. \times 2) at 80 °C for 8 h. ^b Isolated yield. ^c Tributyl phosphite (**26**) used as the starting material. ^d Aryl tetrazole (0.5 mmol), substrates **28**, **29** (0.5 mmol), Bu₄NI (10 mol% \times 2), aq TBHP (2 equiv. \times 2) and CH₃CN (500 μ L \times 2) at 80 °C for 8 h. ^e Isolated along with an inseparable mixture of uncharacterized impurities.

suggesting the detrimental influence of the free carboxy group. However, when the acid was converted to methyl hexanoate **24**, it afforded the δ -aminated product **24a**. Thus, there is a distinct distal selectivity irrespective of the attached appended group in an alkyl chain (either from the alkyl carboxylic acid or alkyl alcohol). Similar distal methylene tetrazolylated was observed for valeronitrile **25** affording product **25a**. As further testimony to the distal selectivity, a reaction between a tributyl phosphite **26** and **a** provided a mono γ -aminated product with concurrent oxidation of the phosphite to a phosphate derivative **27a** in 26% yield. The lower yield obtained is due to the consumption of the

oxidant in converting the phosphite to a phosphate. However, a fully oxidized species **27** provided mono γ -aminated product **27a** in 81% yield. Again, high yields were obtained when phosphate **27** reacted with aryl tetrazoles **e** and **j**, giving products **27e** and **27j**, respectively (Scheme 5).

On the other hand, an *n*-butyl chain bearing a tertiary amide such as *N,N*-dibutylbenzamide **28** in the presence of 5-phenyl-2*H*-tetrazole **a** under the standard reaction conditions did not provide the γ -tetrazolylated product rather underwent dealkylation producing *N*-butylbenzamide **28'a** in 84% yield (Scheme 5). Similar dealkylation using 5,10,15,20-tetraphenylporphyrinato iron(III) chloride–Bu^oOOH is reported.^{21b} The reaction between *N*-methoxy-*N*-methylpentanamide (Weinreb amide) (**29**) and tetrazole under the present conditions provided a multitude of products. The major CDC product occurs adjacent to oxygen and nitrogen heteroatoms. The CDC product adjacent to the nitrogen atom **29'a** could be isolated in its pure form. But the product adjacent to oxygen **29''a** ended up along with other inseparable uncharacterized products.

Intriguingly, another electron-withdrawing group, a ketone, possessing a γ -methylene site, as in valerophenone **30** led to α -functionalized product **30a** in 80% yield (Scheme 6). Acetone **31** having two symmetrical methyl groups α - to carbonyl reacted with **a** to yield product **31a**. Unsymmetrical dialkyl ketone having two sets of alpha hydrogen as in **32** afforded a regioisomeric mixture of products **32a** and **32'a** (1 : 1.25) in 35% and 44% yields, respectively. There is a marginal preference towards the α -side of the longer alkyl chain. This observation is consistent with Zhang's oxidative imidation of ketone with saccharin.²² Computationally BDEs and the spin densities are estimated to be quite similar for both the regioisomeric radicals. However, there is a slight kinetic bias (0.2–0.3 kcal mol^{−1}) for the formation of radicals at the alpha position to the longer alkyl chain, which accounts for a marginal preference for forming **32'a** over **32a** (see Fig. 4 in the Computational studies section).

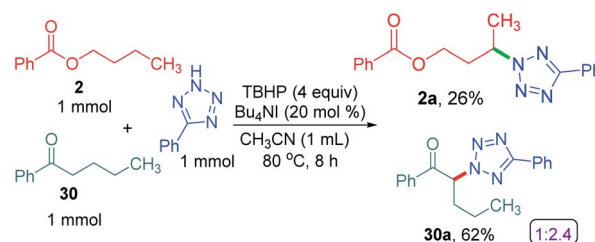


Scheme 6 Substrate scope for alpha-site-selective amination. ^a Reaction conditions: 5-phenyl-2*H*-tetrazole (0.5 mmol), substrates **30–34** (0.5 mmol), Bu₄NI (20 mol%), aq TBHP (4 equiv.) and CH₃CN (1 mL) at 80 °C for 8 h. ^b Isolated yield.

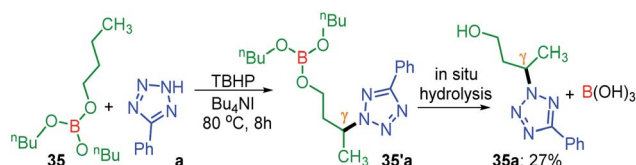
We examined another keto containing substrate, 6-methoxy tetralone **33**, having three potential amination sites namely a benzylic, a methoxy and an alpha C–H to the ketone to access the most preferential site. Substrate **33** provided an exclusive mono-aminated product **33a** at its benzylic position, without affecting the other two sites (Scheme 6). This preferential amination at the benzylic position is further evident when an alkyl pyridine **34** gave its exclusive benzylic product **34a** in 65%.

To determine the preferential selectivity order between an α -carbon to ketone and a distal methylene carbon in an ester, an intermolecular competitive reaction between **2** and **30** was performed (Scheme 7). Interestingly preferential amination took place at the α position of ketone in **30** over the distal methylene carbon in **2** in the ratio of 2.4 : 1 (Scheme 7).

Unlike protected alcohol, amine, amide and carboxylic acid, this method is completely unsuccessful in its free forms. Boron is known to be electron deficient, so for an alkoxy borane, the following question arises: (i) whether the attached alkyl alcohol in the form of alkoxy borane can undergo similar substrate-induced remote amination; (ii) Can the borylated amino alcohols be *in situ* hydrolyzed to generate their amino alcohols and serve as a traceless directing group? With this objective, tributyl borate **35** was subjected to the reaction conditions where mono-amination took place at one of the distal methylene carbons giving **35a** in 27% yield (Scheme 8). No doubt the reaction is induced by the central boron atom and proceeds *via* intermediate **35'a**, but due to the usage of aqueous TBHP the B–O bond got hydrolyzed to free alcohol before completion of amination. Thus, neither the use of aqueous TBHP nor the decane solution of TBHP is suitable for alkyl borate. The peroxy reagent, *tert*-hexyl hydroperoxide (THHP), available in its pure form, was used instead of TBHP. Using the TBAI/THHP combination, the

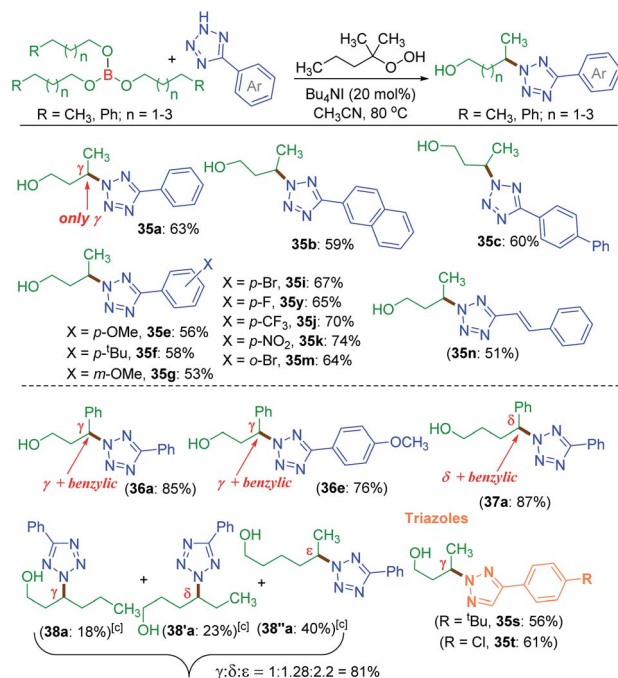


Scheme 7 Intermolecular selectivity between α -carbon to ketone and distal carbon. ^a Reaction conditions: 5-phenyl-2*H*-tetrazole (1 mmol), substrates **2** and **30** (1 mmol), Bu₄NI (20 mol%), aq TBHP (4 equiv.) and CH₃CN (1 mL) at 80 °C for 8 h. ^b Isolated yields.



Scheme 8 Site-selective amination of tributyl borate. ^a Reaction conditions: 5-phenyl-2*H*-tetrazole (0.5 mmol), tributyl borate (5 equiv.), Bu₄NI (20 mol%), aq TBHP (5 equiv.) and CH₃CN (1 mL) at 80 °C for 8 h. ^b Isolated yield. Intermediate **35'a** was not isolated.





Scheme 9 Traceless directing group strategy for amination using borate esters. ^a Reaction conditions: 5-phenyl-2H-tetrazole (0.5 mmol), borate ester **35**–**38** (0.5 mmol), Bu_4NI (20 mol%), *tert*-hexyl hydroperoxide (5 equiv.) and CH_3CN (1 mL) at 80 °C for 8 h. ^b Isolated yield. ^c Trihexyl borate (**38**) was used.

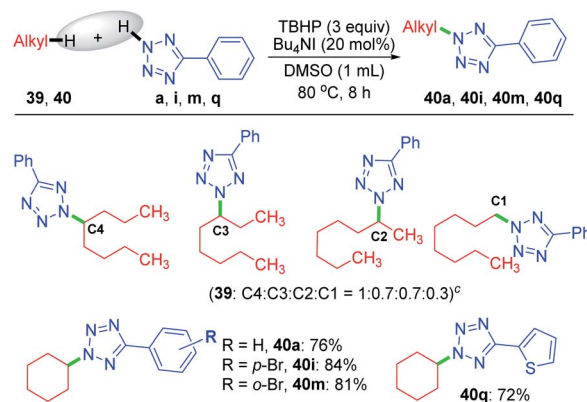
yield of **35a** improved up to 63% (Scheme 9). Thus this is a unique illustration of boron serving as a traceless directing group in any remote functionalization and possesses great synthetic potential.

Employing the modified reaction conditions *i.e.* using *tert*-hexyl hydroperoxide (THHP), γ -amination of **35** was accomplished using an array of electronically diverse aryl tetrazoles (Scheme 9). The use of π -conjugated aromatic hydrocarbons bearing tetrazoles, such as naphthyl **b** and biphenyl **c**, provided good yields of their corresponding γ -aminated butanols **35b** and **35c**. The phenyl ring of tetrazoles substituted with electron-donating groups such as **e**–**g** and electron-withdrawing groups **i**, **k** and **m** all coupled efficiently with **35** to give their respective amino-alcohols **35i**, **35k** and **35m** in yields ranging from 53–74% (Scheme 9). The efficacy of the amination was demonstrated with a cinnamyl tetrazolyl moiety **n**, which afforded product **35n**. The synthetic utility of this transformation was extended to borate **36** and **37** possessing γ and δ phenyl groups respectively {(**36a**, 85%), (**36e**, 76%) and (**37a**, 87%)}. The high product yields obtained for substrates **36** and **37** at their γ and δ position are due to the benzylic nature of this traceless directing group assisted strategy. This strategy was then employed to a longer alkyl chain bearing borate **38** with **a** as the aminating partner. The reaction provided three isolable positional isomers namely γ (**38a**, 18%), δ (**38'a**, 23%) and ϵ (**38''a**, 40%). This result clearly shows distal as well as distance selectivity in the order ($\epsilon > \delta > \gamma$). Unlike in the ester appended alkyl chain (**8a**, Scheme 3) here, the selectivity is much more distinct and the regioisomeric aminoalcohol could be isolated in their

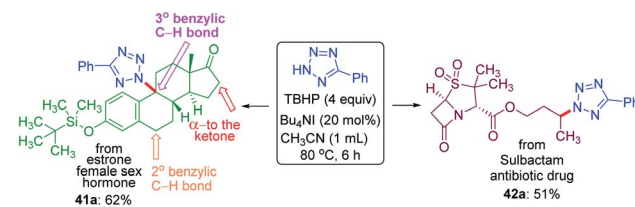
pure form. Finally, this traceless amination strategy was successfully applied to two triazoles **s** and **t** with tributylborate **35**, and both provided triazolyl alcohols **35s** and **35t** demonstrating the power of this traceless strategy (Scheme 9).

Despite the extreme inertness of linear alkane, amination takes place in the decane present in the decane solution of TBHP. This compelled us to use an aqueous solution of TBHP for subsequent investigations. To see if any selectivity can be achieved in a shorter linear chain hydrocarbon having no electronic bias such as *n*-octane **39**, *n*-octane **39** was treated with tetrazole **a** in the presence of TBHP– Bu_4NI combination in DMSO, resulting in a mixture of inseparable aminated products **C4** : **C3** : **C2** : **C1** with a 1 : 0.7 : 0.7 : 0.3 ratio as determined by ^1H NMR. In contrast, a cyclic hydrocarbon such as cyclohexane **40** reacts with a diverse range of electronically substituted tetrazoles **a**, **i**, **m** and **q** to afford good yields of tetrazole-*N*-cycloalkylated products (Scheme 10).

To demonstrate the site-selective intermolecular CDC amination strategy towards late-stage synthetic applications, we subjected estrone **41**, a female sex hormone and sulbactam **42** an antibiotic to our present protocol (Scheme 11). For the estrone **41** having both 2° and 3° benzylic carbon and α -carbon to the ketone, the amination took place at the sterically hindered 3° benzylic position (**41a**, 62% yield) without affecting the other two sites suggesting the dominance of the electronic



Scheme 10 Substrate scope of *N*-cycloalkylation of aryl tetrazoles. ^a Reaction conditions: aryl tetrazole (0.5 mmol), hydrocarbon **39**, **40** (1.0 mmol), Bu_4NI (20 mol%), aq TBHP (4 equiv.) and DMSO (1 mL) at 80 °C for 8 h. ^b Isolated yield. ^c Isomer ratio determined by ^1H NMR.



Scheme 11 Late stage amination of biologically active molecules. ^a Reaction conditions: 5-phenyl-2H-tetrazole (0.25 mmol), substrates **41** (0.25 mmol), **42** (1 mmol), Bu_4NI (20 mol%), aq TBHP (4 equiv.) and CH_3CN (1 mL) at 80 °C for 6 h.

over the steric factor. The sulbactam **42** is a β -lactamase inhibitor used in combination with the antibiotic ampicillin for the treatment of bacterial infections resistant to β -lactam antibiotics. The butyl esters moiety of sulbactam **42** underwent site-selective intermolecular CDC amination giving product **42a** in 51% isolated yield at the electron-rich remote methylene C(sp³)-H bond of the ester group.

Thus, the preferential order for site-selective amination obtained so far in an appended alkyl chain from the overall investigation follows 3° benzylic > 2° benzylic > α to keto > distal methylene ($\gamma > \delta > \epsilon$).

Mechanistic studies

To shed light on the mechanistic pathway of this site-selective remote intermolecular amination, a series of control experiments were performed (Scheme S1, see ESI†) from which the following conclusion was drawn: (i) the reaction is non-ionic; (ii) non-involvement of hypoiodite [Bu₄N]⁺[IO][−]; (iii) radical nature of the reaction and formation of a radical center on the ester moiety and involvement of the ^tBuOO radical; (iv) formation of the N-centered tetrazolyl radical. Based on our findings and following the previous literature,^{7–9,17,23} a plausible mechanistic pathway is depicted in Scheme 12. Initially, oxidation of iodide by TBHP generates the *tert*-butoxyl radical, hydroxyl anion, and iodine. The *in situ* generated iodine reacts with another molecule of TBHP and a hydroxyl ion to afford a *tert*-butylperoxy radical and a molecule of water. Consequently, these radicals abstract a hydrogen atom from the ester and aryl tetrazole resulting in the formation of both carbon C and nitrogen-centered N radical intermediates. Finally, the coupling of carbon C and nitrogen centered radical N affords the desired intermolecular aminated product **1a** (Scheme 12).

Computational studies

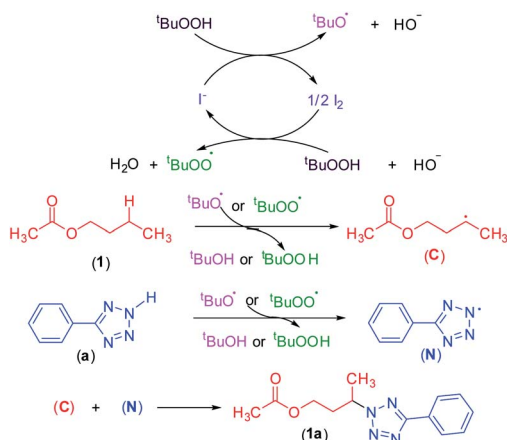
To understand the reasons and also to gain additional insights on the selectivity, we have performed DFT computations at (U) M06-2X²⁴/6-311G(d,p)²⁵ and (U)wB97XD²⁶/6-311G(d,p) levels of theory using Gaussian 09.²⁷ For the majority of the substrates

and their radical isomers, geometries have been optimized at both levels of theory. Furthermore, their relative stability has been compared using bond dissociation energies (BDEs) for deducing the most stable radical among the isomers. Bond dissociation energies and spin densities have been estimated to enumerate the kinetic and thermodynamic stability of the radicals. All these results have been compared, which showed excellent consistency with the experimentally observed selectivity and the yields. In the preceding section, the decisive factors in the selectivity for each type of substrate are discussed based on the computational studies:

(a) Unactivated alkanes. In the case of *n*-octane **39**, four isomeric radicals are possible, among them, the 2° radicals, namely (2-, 3- and 4-dehydro *n*-octyl radicals) are more stable compared to the 1° radical. Among these, the 2° radicals (having radical centers at C2, C3 and C4) have nearly the same BDEs and spin densities, which makes them equally probable to react; in other words, the reaction did not lead to any selectivity (Fig. 1, and Table S4, see ESI†).

(b) Esters. In the case of *n*-butyl acetate, four carbon-centered radicals are possible in the butyl chain, however, the α -radical with the unpaired electron positioned alpha to the ester oxygen is found to be thermodynamically stable on account of the lower spin density and BDE (Fig. 2a), whereas the δ -radical is the least stable due to less hyperconjugation. Despite occupying the second in the series among those four isomeric radicals, the γ -radical formation is a kinetically preferred channel based on the results at both the levels of theory (Fig. 2b). This has been illustrated from the reaction between the ^tBuO radical and *n*-butyl acetate. Interestingly, a higher positive entropy of the product is also favored by its formation over the α -radical. The distal selectivity has also been observed upon increasing the chain length of the ester. In the case of **6** and **7**, product selectivity has been restricted to γ -/ δ - and δ -/ ϵ -, respectively (Fig. 2c and d). Once again, the BDEs and spin densities showed excellent consistency for the distal selectivity and, more importantly, the isolated yields of regioisomers.

In the case of *n*-propyl acetate **11** (Scheme 4), computations predict the α -radical as the thermodynamically most stable radical as well as kinetically more favored, whereas the γ -radical is the least stable among the isomers (Fig. 3a). Besides that, the γ radical has the least preference. However, the inspection of the optimized geometries of radicals indicates that both α and γ radicals attain a zigzag geometry with a planar radical center, whereas the β radical attained a twisted geometry upon



Scheme 12 The putative mechanism for an oxidative C-N bond formation.

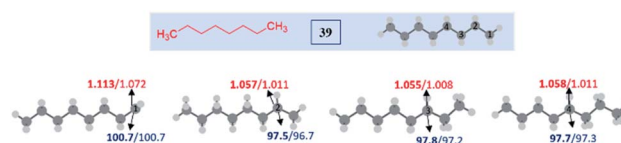


Fig. 1 Computed data on the distal selectivity in unactivated alkane (*n*-octane) **39**. For each case, spin densities are represented in red and the BDEs (in kcal mol^{−1}) are mentioned in blue; bold – (U)M06-2X/6-311G(d,p) and normal font – (U)wB97XD/6-311G(d,p) levels of theory.



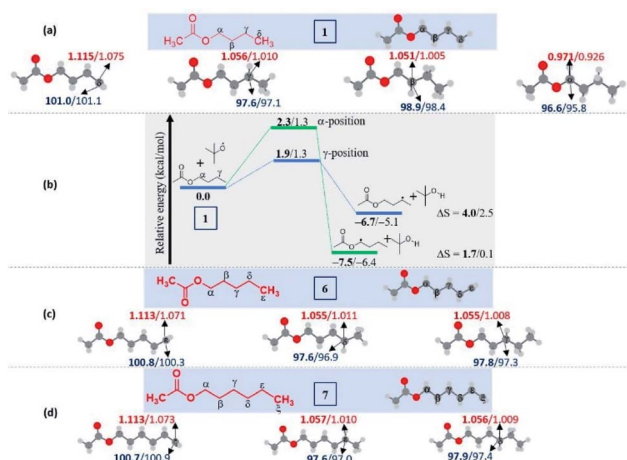


Fig. 2 Computed data on the distal selectivity in esters: (a) *n*-butyl acetate **1**; (b) energy profile depicting the kinetic favourability of the γ -radical over α -radical formation in **1** through hydrogen abstraction by the ^tBuO radical (the corresponding transition states are indicated in Fig. S7 in the ESI[†]); (c) *n*-pentyl acetate **6** and, (d) *n*-hexyl acetate **7**; (for each case, spin densities are represented in red and the BDEs (in kcal mol⁻¹) are mentioned in blue; the energies (in kcal mol⁻¹) relative to the reactants, and the thermodynamic entropy changes (in cal K⁻¹ mol⁻¹) accompanying the reactions are indicated; bold – (U) M06-2X/6-311G(d,p) and normal font – (U)wB97XD/6-311G(d,p) levels of theory).

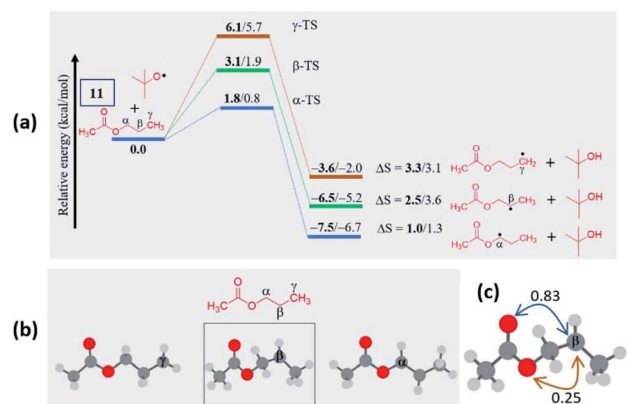


Fig. 3 (a) Computed energy profiles depicting the kinetic and thermodynamic favorability of radical formation in **11**; the energies relative to the **11** + ^tBuO radical are indicated (in kcal mol⁻¹), the thermodynamic entropy changes accompanying the reactions are indicated (in cal K⁻¹ mol⁻¹); (the corresponding transition states are indicated in Fig. S8 in the ESI[†]); (bold – (U)M06-2X/6-311G(d,p) and normal font – (U)wB97XD/6-311G(d,p) levels of theory). (b) Optimized geometries of possible isomeric radical intermediates in *n*-propyl acetate (**11**) at the (U)M06-2X/6-311G(d,p) level of theory; (c) The second-order perturbation energies (in kcal mol⁻¹) from the natural bond orbital (NBO) analysis of the β -radical of *n*-propyl acetate (**11**) at the (U)M06-2X/6-311G(d,p) level of theory.

optimization (Fig. 3b). The attempts to optimize the zigzag β radical structure led to a saddle point. To enumerate the reason, we have also performed NBO analysis,²⁸ which showed a weak interaction between the radical center and the lone pairs of oxygen (Fig. 3c). Presumably, the twist in the alkyl chain

containing the radical center at the β -position, along with the lone pairs of oxygen (at the anomeric position) prevents the approach of the tetrazolyl radical to form the N–C bond. On the other hand, if the chain length of the ester increases (for *e.g.* *n*-butyl acetate), the geometries did not show any twists. So, the distal product selectivity dominates (Fig. 2a). In the case of short-chain esters (ethyl acetate), the α -position is favored over the β -position, which can be accounted for their thermodynamic stability and the ease of formation of the corresponding radicals (Fig. S5–S7, see ESI[†]).

(c) **Ketones.** Unlike esters, the ketones prefer the α -selectivity in almost all cases. The preference mainly arises due to the delocalization of the radical character with the carbonyl group, which can be reflected in the lower spin density values at those centers. Apart from the thermodynamic stabilization of the α -radical, the lower BDE suggests that the ease of radical formation at the α -position over distal positions indicates a kinetic preference. Notably, this can be observed in the case of aryl alkyl ketone **30**, where the major product is **30a**, an alpha selective product (Fig. 4a). The situation is quite interesting in unsymmetrical ketones with two different alkyl chains as both the α -positions are equally favorable. However, the experimental results showed a mixture of products with a marginal preference for the C–N bond formation at the long alkyl chain over the short alkyl chain (Fig. 4b). Despite the spin densities and the BDEs being comparable, a slight kinetic preference (0.2–0.3 kcal mol⁻¹) to form a radical at the alpha position to the longer alkyl chain has been predicted. Presumably, lower kinetic favorability led to a mixture of products and a marginal

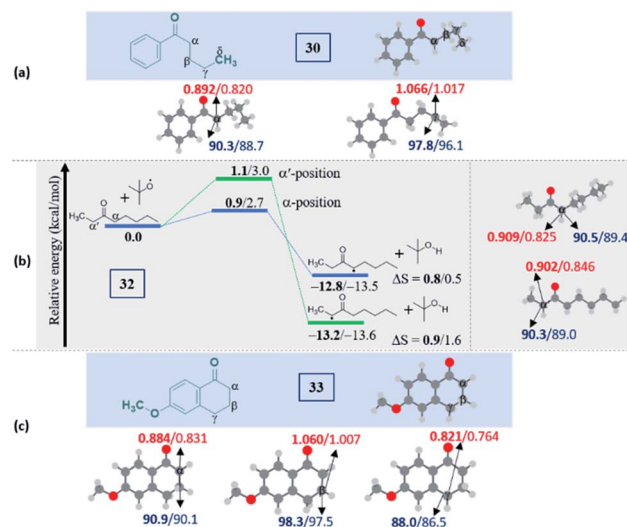


Fig. 4 Computed data on the distal selectivity in ketones: (a) valerophenone **30**; (b) energy profile depicting the formation of the radicals **32a** and **32'a** by hydrogen abstraction by the ^tBuO radical from **32** (the corresponding transition states are indicated in Fig. S9 in the ESI[†]); (c) **33** (for each case, spin densities are represented in red and the BDEs (in kcal mol⁻¹) are mentioned in blue; the energies (in kcal mol⁻¹) relative to the reactants, and the thermodynamic entropy changes (in cal K⁻¹ mol⁻¹) accompanying the reactions are indicated; bold – (U) M06-2X/6-311G(d,p) and normal font – (U)wB97XD/6-311G(d,p) levels of theory).

preference for forming **32/a** over **32a** (Fig. S9, see ESI†). The importance of radical stability in the selectivity can be envisaged in **33**, where the selectivity is not at the α -position to the carbonyl. Instead, the major product is found to be γ -selective (Fig. 4c). The domination of the resonance stabilization at the γ -position over the α -position is responsible for such a remarkably high selectivity. Indeed, this situation can be rationalized from the lower spin density and BDE at the γ -position than at the α -position.

It is noteworthy to mention that the delocalization of spin density and changes in the bond distances between the radical center and the carbonyl carbon (shortening) indicate the stabilization of the alpha radicals in both valerophenone (**30**) and methylcaproate (**24**) (Fig. S10, see ESI†). However, valerophenone and methylcaproate react in a contrasting fashion exhibiting different product selectivities. We presume that the lone pairs of the ester oxygen disfavor the approach of the tetrazole radical in forming the C–N bond. Overall, the kinetic factors associated with the reactions control the selectivity in esters, whereas the thermodynamic stability of the radicals dictates the selectivity in ketones. On the other hand, the unactivated systems did not show any selectivity at all.

Conclusions

We have developed and rationalized site-selective intermolecular amination *via* CDC at unactivated, non-acidic, remote methylene positions without the aid of any directing group or designer catalysts under metal-free conditions. This site-selective amination takes place with a remarkable level of selectivity with a range of electron-withdrawing groups possessing alkyl chains of various lengths. Unprotected functional groups such as alcohol, amines, amides, and carboxylic acid-containing ionizable hydrogens are unsuccessful substrates for this strategy. Alkyl borated substrates gave identical selectivity giving free amino alcohol where the appended boron atom serves as a traceless directing group which is unprecedented in any remote C_{sp³}–H functionalization. In a di-alkyl ester, the selectivity is dictated by the –COO– group giving the same γ selectivity irrespective of its origin either from the alcohol or carboxy acid. In an appended alkyl chain, the preference is always for the distal carbon. It is best when the position is at the γ carbon to the directing group. The site-selectivity is favored at the methine C–H over methylene and methyl of the same length alkyl chain. The degree of distal selectivity decreases as the chain length increases, which is noticeable up to the ϵ position beyond which it is unnoticeable. The ketone and benzylic functionalities bearing longer alkyl chains provided α -aminated (proximal) products rather than remote-functionalization, and when the remote site is also benzylic, high yields of products are obtained. Substrates possessing multiple amenable sites undergo selective mono-amination, which allows late-stage functionalization. Linear hydrocarbons having no electronic bias are completely unselective, whereas cyclohexane provided mono aminated products. In this site-selective amination strategy, N-heterocyclic azoles (tetrazole and triazole) and saccharine served as the nitrogen centered radicals. From the

control experiments carried out, a radical–radical cross-coupling between carbon and nitrogen center radicals has been proposed for this oxidative C–N bond formation. Based on the DFT computations, such high selectivity has been attributed to the thermodynamic stability (in ketones) or kinetic factors (in esters) associated with the radicals. Furthermore, the unactivated alkanes led to a mixture of products without any selectivity, which corroborates nearly the same stability of various distal radical isomers. Thus, we have rationalized site selectivity amination without the *de novo* approach and found it is solely dependent on the intrinsic reactivity of the substrate.

Data availability

Optimization of reaction parameters, mechanistic studies, crystallographic description, all experimental procedures, characterisation data, computational details, and copies of ¹H, ¹³C{¹H}, ¹⁹F and, ³¹P NMR spectra for all compounds featured in this manuscript are provided in the ESI.†

Author contributions

S. R. and B. K. P. conceived, designed and executed the project. S. R. conducted all the experiments and with B. K. P. analysed the data. M. S. and S. V. performed the computational calculations and interpreted the data. S. R., B. K. P., M. S. and S. V. all prepared the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

B. K. P. acknowledges the support of this research by SERB (EMR/2016/007042) and CSIR 02(0365)/19-EMR-II. S. R. acknowledges IIT Guwahati for the fellowship. S. V. acknowledges IISER Mohali for the financial support from SERB (CRG/2019/005744) and DST Indo-Czech Republic bilateral scientific research cooperation, Ministry of Science and Technology (DST/INT/CZ/P-17/2019). M. S. thanks the CSIR and SERB for the fellowship. We are also thankful to IITG central instrument facilities for spectroscopic analysis. We thank Dr P. Rajamalli, MRC, IISc Bangalore, for providing lab facilities and Anjali Mahadevan, IISER Mohali for help in the computations.

Notes and references

- (a) F. P. Guengerich and T. L. Macdonald, *Acc. Chem. Res.*, 1984, **17**, 9; (b) M. Sono, M. P. Roach, E. D. Coulter and J. H. Dawson, *Chem. Rev.*, 1996, **96**, 2841; (c) F. P. Guengerich, *J. Biochem. Mol. Toxicol.*, 2007, **21**, 163; (d) P. R. Ortiz de Montellano, *Chem. Rev.*, 2010, **110**, 932.
- (a) R. T. Gephart III, D. L. Huang, M. J. B. Aguila, G. Schmidt, A. Shahu and T. H. Warren, *Angew. Chem., Int. Ed.*, 2012, **51**, 6488; (b) Y. Lv, K. Sun, T. Wang, Y. Wu, G. Li, W. Pu and S. Mao, *Asian J. Org. Chem.*, 2016, **5**, 325; (c) G.-J. Deng,



- F. Xiao and L. Yang, CHAPTER 5 Cross dehydrogenative coupling reactions involving allyl, benzyl and alkyl C-H bonds, in *From C-H to C-C bonds: Cross-dehydrogenative-coupling*, The Royal Society of Chemistry, 2015, pp. 93–113; (d) R. Vanjari and K. N. Singh, *Chem. Soc. Rev.*, 2015, **44**, 8062; (e) T. A. Ramirez, B. Zhao and Y. Shi, *Chem. Soc. Rev.*, 2012, **41**, 931; (f) I. Bosque, R. Chinchilla, J. C. Gonzalez-Gomez, D. Guijarro and F. Alonso, *Org. Chem. Front.*, 2020, **7**, 1717; (g) Z.-W. Hou, D.-J. Liu, P. Xiong, X.-L. Lai, J. Song and H.-C. Xu, *Angew. Chem., Int. Ed.*, 2021, **60**, 2943; (h) Z. Ruan, Z. Huang, Z. Xu, S. Zeng, P. Feng and P.-H. Sun, *Sci. China: Chem.*, 2021, **64**, 800; (i) Z.-W. Hou, L. Li and L. Wang, *Org. Chem. Front.*, 2021, **8**, 4700.
- 3 (a) Z. Li and C.-J. Li, *J. Am. Chem. Soc.*, 2004, **126**, 11810; (b) C.-Y. Huang, H. Kang, J. Li and C.-J. Li, *J. Org. Chem.*, 2019, **84**, 12705; (c) G. Majji, S. K. Rout, S. Rajamanickam, S. Guin and B. K. Patel, *Org. Biomol. Chem.*, 2016, **14**, 8178; (d) C.-J. Li, *Acc. Chem. Res.*, 2009, **42**, 335; (e) Z. Li, D. S. Bohle and C.-J. Li, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**, 8928; (f) M. K. Lakshman and P. K. Vuram, *Chem. Sci.*, 2017, **8**, 5845; (g) S. Rajamanickam, G. Majji, S. K. Santra and B. K. Patel, *Org. Lett.*, 2015, **17**, 5586.
- 4 (a) J. He, T. Shigenari and J.-Q. Yu, *Angew. Chem., Int. Ed.*, 2015, **54**, 6545; (b) Q. Gou, G. Liu, Z.-N. Liu and J. Qin, *Chem.-Eur. J.*, 2015, **21**, 15491; (c) G. He and G. A. Chen, *Angew. Chem., Int. Ed.*, 2011, **50**, 5192; (d) J. J. Topczewski, P. J. Cabrera, N. I. Saper and M. S. Sanford, *Nature*, 2016, **531**, 220; (e) C. Sambiasi, D. Schönbauer, R. Blicke, T. Dao-Huy, G. Pototschnig, P. Schaaf, T. Wiesinger, M. F. Zia, J. Wencel-Delord, T. Besset, B. U. W. Maes and M. Schnürch, *Chem. Soc. Rev.*, 2018, **47**, 6603; (f) S. Rej, Y. Ano and N. Chatani, *Chem. Rev.*, 2020, **120**, 1788; (g) Y. Xu and G. Dong, *Chem. Sci.*, 2018, **9**, 1424; (h) L. Cuesta and E. P. Urriolabeitia, CHAPTER 8 Coordination-directed metallation strategy for C-H functionalization, in *C-H and C-X Bond functionalization: Transition metal mediation*, The Royal Society of Chemistry, 2013, pp. 262–309; (i) Z. Chen, B. Wang, J. Zhang, W. Yu, Z. Liu and Y. Zhang, *Org. Chem. Front.*, 2015, **2**, 1107.
- 5 (a) K. Liao, T. C. Pickel, V. Boyarskikh, J. Bacsá, D. G. Musaev and H. M. L. Davies, *Nature*, 2017, **551**, 609; (b) K. Liao, S. Negretti, D. G. Musaev, J. Bacsá and H. M. L. Davies, *Nature*, 2016, **533**, 230; (c) K. Liao, Y.-F. Yang, Y. Li, J. N. Sanders, K. N. Houk, D. G. Musaev and H. M. L. Davies, *Nat. Chem.*, 2018, **10**, 1048; (d) W. Liu, Z. Ren, A. T. Bosse, K. Liao, E. L. Goldstein, J. Bacsá, D. G. Musaev, B. M. Stoltz and H. M. L. Davies, *J. Am. Chem. Soc.*, 2018, **140**, 12247.
- 6 (a) M. S. Chen and M. C. White, *Science*, 2007, **318**, 783; (b) M. S. Chen and M. C. White, *Science*, 2010, **327**, 566; (c) M. C. White and J. Zhao, *J. Am. Chem. Soc.*, 2018, **140**, 13988; (d) S. Jana, M. Ghosh, M. Ambule and S. S. Gupta, *Org. Lett.*, 2017, **19**, 746.
- 7 (a) Y. Kawamata, M. Yan, Z. Liu, D.-H. Bao, J. Chen, J. T. Starr and P. S. Baran, *J. Am. Chem. Soc.*, 2017, **139**, 7448; (b) V. A. Schmidt, R. K. Quinn, A. T. Brusoe and E. J. Alexanian, *J. Am. Chem. Soc.*, 2014, **136**, 14389; (c) R. K. Quinn, Z. A. Könst, S. E. Michalak, Y. Schmidt, A. R. Szklarski, A. R. Flores, S. Nam, D. A. Horne, C. D. Vanderwal and E. J. Alexanian, *J. Am. Chem. Soc.*, 2016, **138**, 696; (d) W. L. Czaplyski, C. G. Na and E. J. Alexanian, *J. Am. Chem. Soc.*, 2016, **138**, 13854.
- 8 (a) A. Sharma and J. F. Hartwig, *Nature*, 2015, **517**, 600; (b) R. R. Karimov, A. Sharma and J. F. Hartwig, *ACS Cent. Sci.*, 2016, **2**, 715.
- 9 (a) X. Huang, T. M. Bergsten and J. T. Groves, *J. Am. Chem. Soc.*, 2015, **137**, 5300; (b) H. Wu, Z. Xiao, J. Wu, Y. Guo, J.-C. Xiao, C. Liu and Q.-Y. Chen, *Angew. Chem., Int. Ed.*, 2015, **54**, 4070; (c) S. Guo, X. Zhang and P. Tang, *Angew. Chem., Int. Ed.*, 2015, **54**, 4065; (d) P. E. Gormisky and M. C. White, *J. Am. Chem. Soc.*, 2013, **135**, 14052.
- 10 (a) A. W. Hofmann, *Ber. Dtsch. Chem. Ges.*, 1879, **12**, 984; (b) K. Löffler and C. Freytag, *Ber. Dtsch. Chem. Ges.*, 1909, **42**, 3427.
- 11 (a) X. Hu, G. Zhang, F. Bu, L. Nie and A. Lei, *ACS Catal.*, 2018, **8**, 9370; (b) F. Wang and S. S. Stahl, *Angew. Chem., Int. Ed.*, 2019, **58**, 6385.
- 12 (a) S. Blakey and N. M. Weldy, C-N Bond formation by C-H functionalization via metal-catalyzed nitrene insertion, in *C-N Bond formation by C-H functionalization via metal-catalyzed nitrene insertion; in catalytic transformations via C-H activation*, ed. J.-Q. Yu, Georg Thieme Verlag, Stuttgart, 2015, vol. 2, pp. 249–286; (b) D. N. Zalatan and J. D. Bois, Metal-catalyzed oxidations of C-H to C-N bonds, in *Metal-catalyzed oxidations of C-H to C-N bonds in C-H activation*, ed. J.-Q. Yu and Z. Shi, Springer, Berlin, Heidelberg, 2009, vol. 292, pp. 347–378; (c) S. M. Paradine, J. R. Griffin, J. Zhao, A. L. Petronico, S. M. Miller and M. C. White, *Nat. Chem.*, 2015, **7**, 987; (d) J. R. Clark, K. Feng, A. Sookezian and M. C. White, *Nat. Chem.*, 2018, **10**, 583; (e) F. Collet, C. Lescot, C. Liang and P. Dauban, *Dalton Trans.*, 2010, **39**, 10401; (f) S. Wiese, Y. M. Badiei, R. T. Gephart, S. Mossin, M. S. Varonka, M. M. Melzer, K. Meyer, T. R. Cundari and T. H. Warren, *Angew. Chem., Int. Ed.*, 2010, **49**, 8850; (g) K. W. Fiori and J. Du Bois, *J. Am. Chem. Soc.*, 2007, **129**, 562; (h) E. N. Bess, R. J. DeLuca, D. J. Tindall, M. S. Oderinde, J. L. Roizen, J. Du Bois and M. S. Sigman, *J. Am. Chem. Soc.*, 2014, **136**, 5783; (i) J. L. Roizen, D. N. Zalatan and J. Du Bois, *Angew. Chem., Int. Ed.*, 2013, **52**, 11343; (j) C. Liang, F. Robert-Peillard, C. Fruit, P. Müller, R. H. Dodd and P. Dauban, *Angew. Chem., Int. Ed.*, 2006, **45**, 4641; (k) C. Lescot, B. Darses, F. Collet, P. Retailleau and P. Daban, *J. Org. Chem.*, 2012, **77**, 7232; (l) J. L. Roizen, D. N. Zalatan and J. Du Bois, *Angew. Chem., Int. Ed.*, 2013, **52**, 11343; (m) J. J. Fleming and J. D. Bois, *J. Am. Chem. Soc.*, 2006, **128**, 3926; (n) H. Lu, V. Subbarayan, J. Tao and X. P. Zhang, *Organometallics*, 2010, **29**, 389.
- 13 (a) X. Wu, K. Yang, Y. Zhao, H. Sun, G. Li and H. Ge, *Nat. Commun.*, 2015, **6**, 6462; (b) Q. Gou, Y.-W. Yang, Z.-N. Liu and J. Qin, *Chem.-Eur. J.*, 2016, **22**, 16057.
- 14 J. Pan, M. Su and S. L. Buchwald, *Angew. Chem., Int. Ed.*, 2011, **50**, 8647.
- 15 J. F. Hartwig, *Acc. Chem. Res.*, 1998, **31**, 852.



- 16 E. Vitaku, D. T. Smith and J. T. Njardarson, *J. Med. Chem.*, 2014, **57**, 10257.
- 17 (a) S. Rajamanickam, C. Sah, B. A. Mir, S. Ghosh, G. Sethi, V. Yadav, S. Venkataramani and B. K. Patel, *J. Org. Chem.*, 2020, **85**, 2118; (b) H. Aruri, U. Singh, M. Kumar, S. Sharma, S. K. Aithagani, V. K. Gupta, S. Mignani, R. A. Vishwakarma and P. P. Singh, *J. Org. Chem.*, 2017, **82**, 1000.
- 18 C. Wang, L. Zhang, A. Ren, P. Lu and Y. Wang, *Org. Lett.*, 2013, **15**, 2982.
- 19 (a) J. M. Hudzik, J. W. Bozzelli and J. M. Simmie, *J. Phys. Chem. A*, 2014, **118**, 9364; (b) L. Bu, P. N. Ciesielski, D. J. Robichaud, S. Kim, R. L. McCormick, T. D. Foust and M. R. Nimlos, *J. Phys. Chem. A*, 2017, **121**, 5475.
- 20 (a) D. Šakić and H. Zipse, *Adv. Synth. Catal.*, 2016, **358**, 3983; (b) G. D. P. Gomes, Y. Loginova, S. Z. Vatsadze and I. V. Alabugin, *J. Am. Chem. Soc.*, 2018, **140**, 14272.
- 21 (a) G. Majji, S. Guin, S. K. Rout, A. Behera and B. K. Patel, *Chem. Commun.*, 2014, **50**, 12193; (b) L. Constantino and J. Iley, *Org. Biomol. Chem.*, 2004, **2**, 1894.
- 22 Y. Lv, Y. Li, T. Xiong, Y. Lu, Q. Liu and Q. Zhang, *Chem. Commun.*, 2014, **50**, 2367.
- 23 E. Shi, Y. Shao, S. Chen, H. Hu, Z. Liu, J. Zhang and X. Wan, *Org. Lett.*, 2012, **14**, 3384.
- 24 Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215.
- 25 R. Krishnan, J. S. Binkley and R. Seeger, *J. Chem. Phys.*, 1980, **72**, 650.
- 26 J.-D. Chai and M. Head-Gordon, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6615.
- 27 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, A. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, J. J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, T. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 09 Revision A.02*, Gaussian, Inc., Wallingford CT, 2016.
- 28 (a) E. D. Glendening, A. E. Reed, J. E. Carpenter and F. Weinhold, *NBO version 3.1*, 2003; (b) J. P. Foster and F. Weinhold, *J. Am. Chem. Soc.*, 1980, **102**, 7211.

