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# NH-1,2,3-triazoles as versatile building blocks in denitrogenative transformations

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The utilization of NH-1,2,3-triazoles as easily accessible building blocks in denitrogenative ring cleavage transformations with electrophiles to provide multifunctionalized nitrogen heterocycles and *N*-alkenyl compounds is reviewed. Leveraging the ready availability of NH-1,2,3-triazoles, these processes provide a convenient route to a range of pharmaceutically relevant heterocyclic cores and *N*-alkenyl compounds. The synthetic usefulness of *in situ* acylated NH-1,2,3-triazoles as viable alternatives to widely explored *N*-sulfonyl-1,2,3-triazoles in ring cleavage processes is highlighted.

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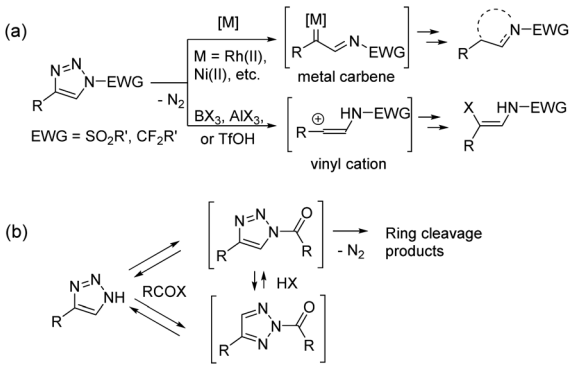
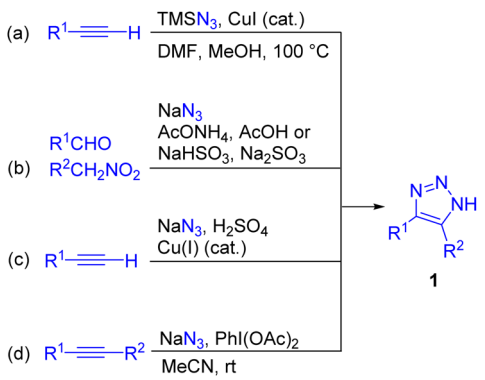
1,2,3-Triazoles are nitrogen heterocycles with versatile reactivity[1] and great medicinal importance.[2] Since the discovery of azide–alkyne click chemistry in 2002,[3] triazole derivatives have gained enormous attention in organic, medicinal, biomolecular, and material sciences. Among them, 1,2,3-triazoles bearing an electron-withdrawing group at position N1 are of special importance because of their propensity to undergo N1–N2 bond cleavage in denitrogenative triazole ring opening transformations (Scheme 1a).[1] *N*-sulfonyl-1,2,3-triazoles[4] and *N*-fluoroalkyl-1,2,3-triazoles[5] are the most explored building blocks, which undergo ring cleavage under metal catalysis or by the action of Lewis or Brønsted acids. Very recently, a new strategy based on the use of NH-1,2,3-triazoles involving the installation of an electron-withdrawing group with *in situ* ring cleavage was described and used with success (Scheme 1b). The present review features the use of free NH-1,2,3-triazoles **1** in denitrogenative transformations, proceeding *via* *N*-acyl-1,2,3-triazoles or their analogues as key intermediates.

*N*-unsubstituted NH-1,2,3-triazoles **1**, considered in the present review, are the simplest and most readily available triazoles.[6] They can be prepared by azide–alkyne cycloaddition[6] or alternative methods such as cycloaddition/elimination with activated ketones[7] or nitroalkenes.[7] In the last five years, there has been a notable surge of innovative methods for the synthesis of NH-1,2,3-triazoles and several one-pot protocols from inexpensive and commercially available reagents have been developed.[6] To underline the most efficient and practical routes, NH-1,2,3-triazoles were synthesized from $TMSN_3$ and alkynes *via* CuI-catalysed cycloaddition (Scheme 2a),[8] or sodium azide, aldehydes and nitroalkanes *via* a tandem Henry reaction/[3 + 2] cycloaddition (Scheme 2b),[9a–c] including recently developed green chemistry approaches.[9d–h] In 2022, NH-1,2,3-triazoles became available from $NaN_3/H_2SO_4$ and alkynes, which is so far the simplest and the most straightforward route, although the generation of $HN_3$ raises safety concerns (Scheme




Scheme 1 Schematic representation of the utilization of (a) *N*–sulfonyl- or *N*-fluoroalkyl-substituted 1,2,3-triazoles and (b) NH-1,2,3-triazoles in ring cleavage denitrogenative transformations.




Scheme 2 Overview of efficient routes for the synthesis of NH-1,2,3-triazoles (a–d).

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2c).[10] Finally, the approach utilizing azidyl radical–alkyne cycloaddition with the use of the $NaN_3/PhI(OAc)_2$ system in mild conditions is highly efficient for complex disubstituted triazoles such as 4,5-diaryltriazoles (Scheme 2d),[11a] and it was also possible to efficiently synthesize these compounds without an oxidant, albeit only under harsh conditions (MW heating at 200 °C).[11b] More examples of novel synthetic methods to access NH-1,2,3-triazoles appeared in recent reviews.[6]

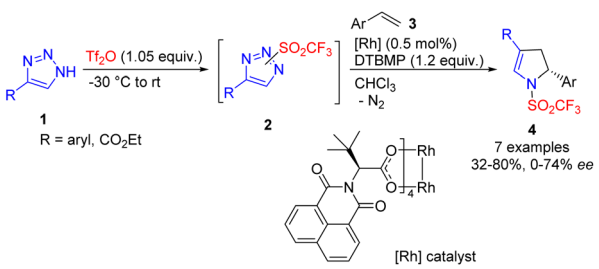
Due to better atom economy, the utilization of readily available NH-triazoles is advantageous compared to the use of *N*-sulfonyl- or *N*-fluoroalkyl-triazoles. Additionally, access to NH-1,2,3-triazoles was possible using "alkyne-free" methods. Both, primary nitro compounds and aromatic aldehydes are easily accessible industrial scale products.[12]

One of the first denitrogenative transformations of 1,2,3-NH-triazoles **1** was reported in 2014. In this process, *in situ* sulfonylation with triflic anhydride and 2,6-di(*tert*-butyl)-4-methylpyridine (DTBMP) as a base was used to generate reactive *N*-triflyl triazoles **2**.[13] Their ring cleavage by a chiral Rh(II) catalyst in the presence of an excess of alkene **3** afforded 2,3-dihydropyrroles **4** with low to good enantiocontrol (Scheme 3).

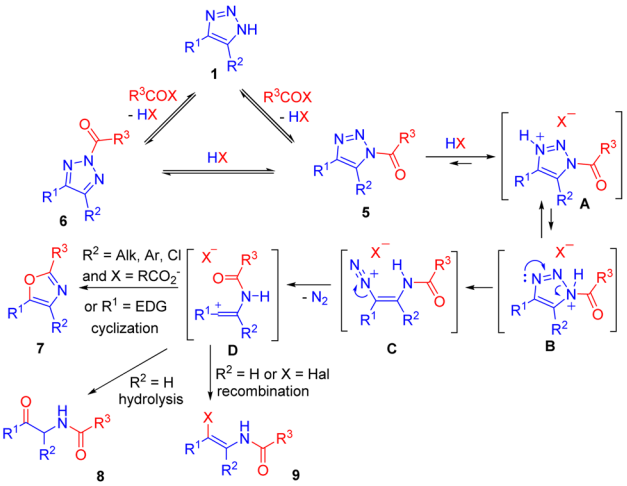
However, besides triflation, there are no other examples of *in situ* sulfonylation of NH-1,2,3-triazoles followed by denitrogenative transformations. Therefore, this reaction is limited to the extremely electron-accepting triflyl group and analogous ring cleavage did not proceed with other *N*-sulfonyl triazoles.

In contrast to sulfonylation, acylation of NH-1,2,3-triazoles is more versatile and has developed into a highly active area of research in recent years.[14] Tandem acylation followed by ring cleavage without isolation of *N*-acyltriazoles was performed using acyl halides or acid anhydrides. The mechanism of this transformation, recently confirmed by us,[14] involved the formation of N1 (**5**) and N2-acylated (**6**) triazoles in equilibrium, followed by acid-mediated cleavage of the former. Denitrogenation and formation of a vinyl cation in an irreversible step was the driving force of N2–N1-acyltriazole interconversion, which ensured the complete transformation of triazoles into ring cleavage products **7–9** (Scheme 4).

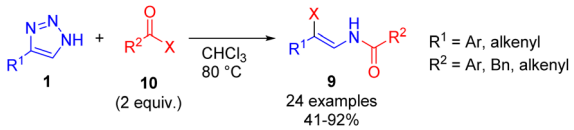
Cleavage of NH-1,2,3-triazoles **1** with an excess of acyl halides **10** (X = Cl, Br) under elevated temperature led to the formation of β-enamido halides **9** in moderate to good yields (Scheme 5).[15] β-Enamido halides are difficult to access by other synthetic routes and are present in natural products, which underlines the synthetic value of the method.




Scheme 3 Synthesis of 2,3-dihydropyrroles by cleavage of NH-triazole in the presence of triflic anhydride. DTBMP – 2,6-di(*tert*-butyl)-4-methylpyridine.




Scheme 4 Mechanism of NH-triazole cleavage with electrophiles.

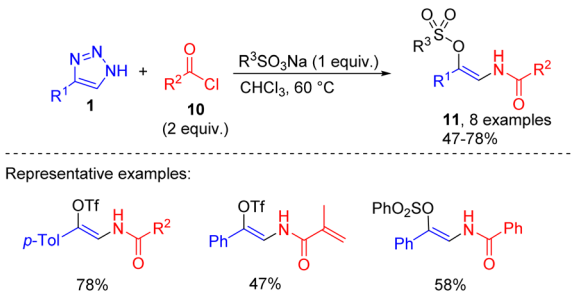



Scheme 5 Cleavage of NH-1,2,3-triazoles with acyl halides.

This transformation in the presence of sodium sulfonates was employed in the synthesis of enamido triflates or sulfonates **11**. Mainly compounds with the phenacyl group at the nitrogen were accessed by the mentioned route (Scheme 6).[15]

An alternative method, applicable to the synthesis of β-fluoroacylenamido triflates is based on the formation of N2-acyltriazoles **6** *via* the *in situ* acylation of NH-1,2,3-triazoles with fluorinated acid anhydrides followed by their treatment with triflic acid, which proceeds through N2–N1 acyltriazole interconversion and ring cleavage (Scheme 7).[14]

The products are useful building blocks, that can get involved in Pd-catalysed cross-coupling substitution reactions of the triflate group to access multifunctionalized enamide derivatives – attractive drug candidates and synthetic intermediates.[16] None of the methods mentioned was applicable with the less reactive alkyl-substituted acylating agents ($Ac_2O$, AcCl),




Scheme 6 Synthesis of β-enamido triflates and sulfonates from NH-1,2,3-triazoles, acyl halides, and sodium sulfonates.

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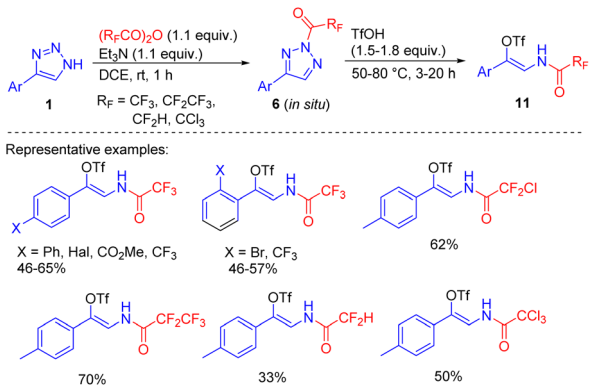
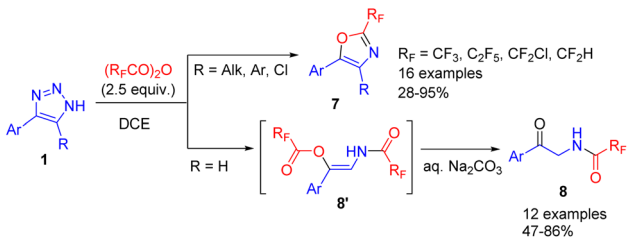



Scheme 7 One-pot two step synthesis of β-fluoroacylenamido triflates. DCE = 1,2-dichloroethane.

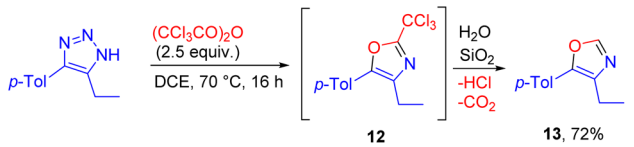



Scheme 8 Synthesis of fluoroalkylated oxazoles and 2-acylaminoketones from NH-1,2,3-triazoles with fluoroalkylated acid anhydrides.

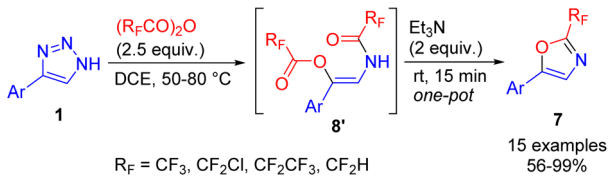
because the resulting acyltriazoles were resistant to ring cleavage even at elevated temperatures.[14]

The cleavage of NH-1,2,3-triazoles with an excess of fluoroalkylated acid anhydrides led to highly pharmaceutically relevant 2-fluoroalkyl oxazoles 7 (in the cases of 4,5-disubstituted triazoles) or 2-acylaminoketones (for 4-substituted triazoles, R = H) (Scheme 8).[17] In the first case intramolecular cyclization took place, whereas in the second, unstable β-acyloxyenamide **8′** formed, which underwent ester hydrolysis to 2-acylaminoketone **8** upon treatment with an aqueous base. The difference in chemoselectivity was attributed to the increased vinyl cation stability of disubstituted examples, which made them more prone to intramolecular cyclization.

4,5-disubstituted NH-1,2,3-triazole reacted with trichloroacetic anhydride to give 2-unsubstituted oxazole **13**, due to the low stability of the trichloromethyl-substituted product **12** during silica gel column chromatography. The whole transformation is a rare and unique case of a reaction involving trichloroacetic anhydride as a one-carbon building block (Scheme 9).[17]



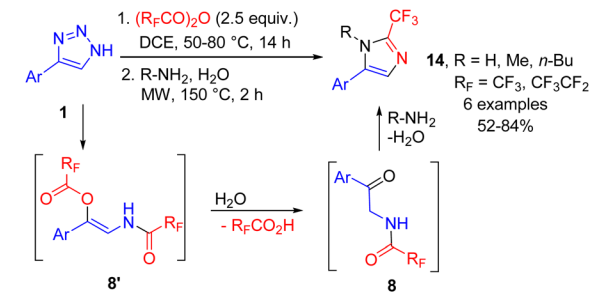

Scheme 9 Formation of 2-unsubstituted oxazole from NH-1,2,3-triazole and trichloroacetic anhydride.



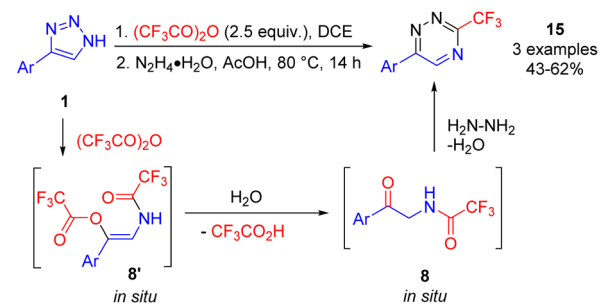

Scheme 10 One-pot synthesis of fluoroalkylated oxazoles from NH-1,2,3-triazoles, acid anhydrides and $Et_3N$.

The limitation of oxazole synthesis to only disubstituted triazoles was overcome by the cyclization of *in situ* formed β-acyloxyenamide **8′** to oxazoles 7 using $Et_3N$ and proceeded quickly and nearly quantitatively under ambient conditions.[18] This one-pot triazole cleavage procedure provided an efficient access to 2-fluoroalkylated oxazoles from monosubstituted triazoles in good to excellent yields (Scheme 10).

The easy access to fluorinated 2-acylaminoketones **8** was utilized in a number of one-pot syntheses of fluoroalkylated heterocycles directly from NH-1,2,3-triazoles **1**. First, 2-fluoroalkyl imidazoles **14** were prepared by cleavage with trifluoroacetic or perfluoropropanoic anhydrides, followed by the treatment of the ketamide intermediate with an aqueous solution of the primary amine (or ammonium acetate for R = H) under microwave conditions. The acid formed after hydrolysis of the enamide to yield 2-acylaminoketone promoted the Robinson–Gabriel cyclization of the latter. This procedure afforded imidazoles **14** in moderate to good yields in a one-pot manner starting from triazoles (Scheme 11).[17]




Scheme 11 One-pot synthesis of 2-fluoroalkyl-imidazoles from NH-1,2,3-triazoles.




Scheme 12 Synthesis of 3-fluoroalkyl-1,2,4-triazines from NH-1,2,3-triazoles.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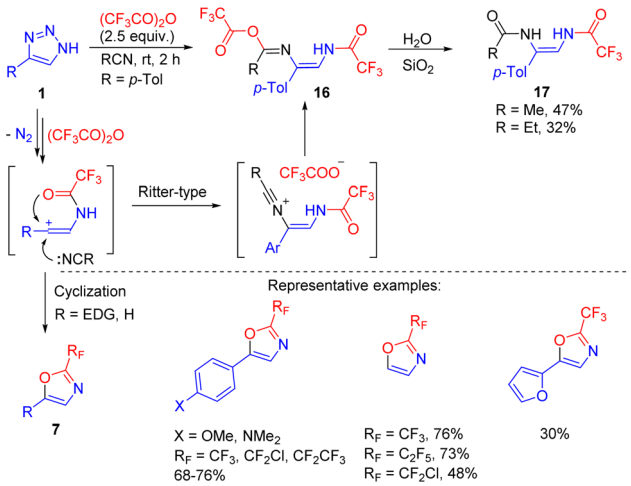
Alternatively, condensation of the formed acyloxyenamide **8′** with hydrazine hydrate after switching the solvent to acetic acid provided fluoroalkylated 1,2,4-triazines **15** (Scheme 12).[17]

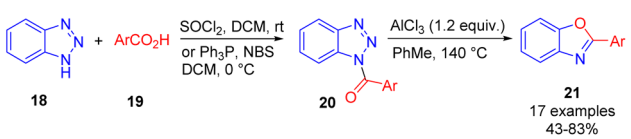
The formation of the vinyl cation intermediate in *N*-acyltriazole cleavage was confirmed by changing the solvent from a chlorinated one to acetonitrile or propionitrile.[18] In one special case (Scheme 13, R = *p*-Tol) adducts **16** of the Ritter reaction were formed and hydrolysed to bis(enamides) **17**. However, this reaction was not general and in the cases of electron-richer triazoles, cyclization of the vinyl cation to oxazoles **7** took place (Scheme 13, R = H, EDG). This route is an alternative to one mentioned above (Scheme 10), and is applicable to electron-rich substrates. The straightforward formation of oxazoles **7** rather than enamides **8′** in polar MeCN was explained by the decreased stability of the vinyl cation–trifluoroacetate anion contact ion pair, which prevented recombination and favoured cyclization.[18]

Several efficient NH-1,2,3-triazole ring cleavage protocols were also developed for NH-benzotriazole **18**, which can be easily and regioselectively acylated on N1. The treatment of the formed *N*-acylbenzotriazole **20** with $AlCl_3$ as a Lewis acid promotor facilitated ring cleavage leading to benzoxazoles under relatively harsh conditions (Scheme 14).[19]

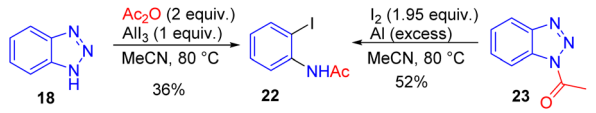
Rare examples of *ortho*-iodoacetanilide **22** formation in moderate yields from NH- and related *N*-acylbenzotriazole were reported in which the $All_3/Ac_2O$ system or aluminium and iodine in acetonitrile were used.[20] These are the only cases of *N*-acetylbenzotriazole **23** ring cleavage known. Importantly, the reaction of *N*-acetylbenzotriazole **23** with $AlCl_3$ was not efficient



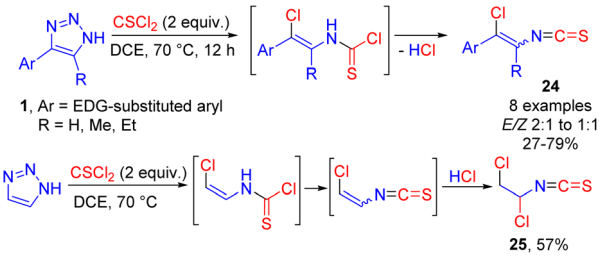

Scheme 13 Formation of bis(enamides) **17** and oxazoles **7** by the cleavage of NH-1,2,3-triazoles with trifluoroacetic anhydride in nitrile solvent.




Scheme 14 Synthesis of benzoxazoles from NH-benzotriazoles.




Scheme 15 Formation of *o*-iodoacetanilide by $All_3$-mediated cleavage of N1-acetylbenzotriazole.



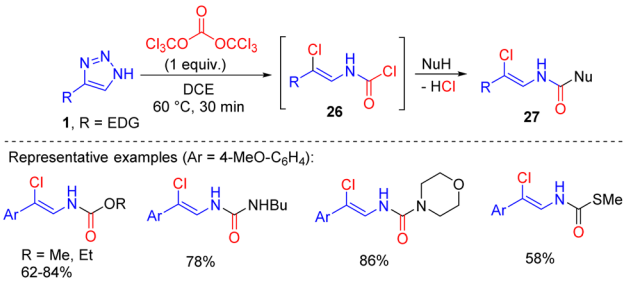

Scheme 16 Cleavage of NH-1,2,3-triazoles with thiophosgene.

and led only to deacylation, and not to the desired ring cleavage product (Scheme 15).[19]

Cleavage of electron-rich 4-aryl-NH-1,2,3-triazoles **1** was successfully achieved with thiophosgene leading to the formation of vinyl isothiocyanates **24** by HCl elimination from the vinyl chloride intermediate (Scheme 16).[21] The vinyl isothiocyanate moiety is present in natural products with antifungal and antibacterial activity and is difficult to access by traditional methods. Switching from electron-rich aromatic NH-triazoles to unsubstituted NH-1,2,3-triazole afforded product **25** of HCl addition across the double bond in moderate yield.

A similar transformation with triazoles **1** bearing an electron-rich aryl or alkenyl substituent in position 4 proceeded with triphosgene.[21] The *in situ* formed carbamoyl chlorides **26** were treated with nucleophiles to gain access to multifunctional compounds **27**, such as *N*-alkenyl carbamates, ureas and thiocarbamates (Scheme 17).

The denitrogenative transformation of NH-1,2,3-triazoles was studied also on more complex substrates such as 4-(1-hydroxycyclobutyl)-1,2,3-triazoles **28**. Their cleavage with acyl chlorides **10** catalysed by triflic acid provided efficient access to cyclic enaminones **29** (Scheme 18).[22] The reaction proceeded *via* the cleavage of *N*-acyltriazole and semipinacol rearrangement




Scheme 17 Synthesis of multifunctional *N*-alkenyl compounds by the cleavage of NH-1,2,3-triazoles with triphosgene.

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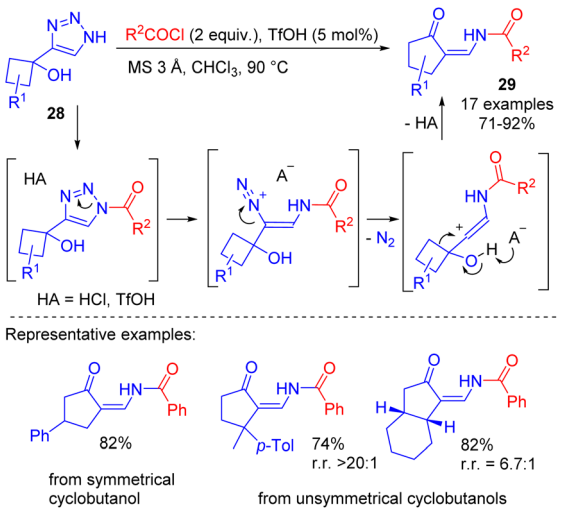


Scheme 18 Synthesis of cyclic enaminones by TfOH-catalyzed cleavage of 4-(1-hydroxycyclobutyl)-1,2,3-triazoles with acyl chlorides.

cascade. The procedure was found to be easily scalable to give multifunctional substrates in good yields.

## Conclusions

In conclusion, NH-1,2,3-triazoles are commercially available or easily synthesized starting materials that exhibit a remarkable versatility in transformations to diverse nitrogen-containing heterocycles and functionalized *N*-alkenyl compounds *via* denitrogenative cleavage. *In situ* prepared *N*-acylated 1,2,3-triazoles are key intermediates in these transformations. Acid-mediated triazole ring opening of *N*-acylated 1,2,3-triazoles, followed by nitrogen elimination affords vinyl cation intermediates, which undergo a variety of reactions such as cyclization or heteroatom capture. Further development of denitrogenation of NH-1,2,3-triazoles accompanied by C–C bond forming reactions, C–H insertion or rearrangement of the vinyl cation can be expected, providing access to a structural diversity of products with potential applications in drug development. Moreover, due to easy availability of NH-1,2,3-triazoles they are excellent starting materials for the development of new industrial synthetic processes.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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## Notes and references

1 (*a*) Chemistry of 1,2,3-triazoles, in *Topics in Heterocyclic Chemistry*, ed. W. Dehaen and V. A. Bakulev, Springer, 2015, vol. 40; (*b*) H. M. Davies and J. S. Alford, Reactions of metallocarbenes derived from *N*-sulfonyl-1,2,3-triazoles, *Chem. Soc. Rev.*, 2014, **43**, 5151; (*c*) W. Li and J. Zhang, Synthesis of Heterocycles through Denitrogenative Cyclization of Triazoles and Benzotriazoles, *Chem. – Eur. J.*, 2020, **26**, 11931.

2 S. Kumar, B. Sharma, V. Mehra and V. Kumar, Recent accomplishments on the synthetic/biological facets of pharmacologically active 1*H*-1,2,3-triazoles, *Eur. J. Med. Chem.*, 2021, **212**, 113069.

3 (*a*) C. W. Tornøe, C. Christensen and M. Meldal, Peptidotriazoles on Solid Phase: [1,2,3]-Triazoles by Regiospecific Copper(I)-Catalyzed 1,3-Dipolar Cycloadditions of Terminal Alkynes to Azides, *J. Org. Chem.*, 2002, **67**, 3057; (*b*) V. V. Rostovtsev, L. G. Green, V. V. Fokin and K. B. Sharpless, A Stepwise Huisgen Cycloaddition Process: Copper(I)-Catalyzed Regioselective "Ligation" of Azides and Terminal Alkynes, *Angew. Chem., Int. Ed.*, 2002, **41**, 2596.

4 (*a*) M. Akter, K. Rupa and P. Anbarasan, 1,2,3-Triazole and Its Analogues: New Surrogates for Diazo Compounds, *Chem. Rev.*, 2022, **122**, 13108; (*b*) B. Chattopadhyay and V. Gevorgyan, Transition-Metal-Catalyzed Denitrogenative Transannulation: Converting Triazoles into Other Heterocyclic Systems, *Angew. Chem., Int. Ed.*, 2012, **51**, 862; (*c*) P. Anbarasan, D. Yadagiri and S. Rajasekar, Recent Advances in Transition-Metal-Catalyzed Denitrogenative Transformations- of 1,2,3-Triazoles and Related Compounds, *Synthesis*, 2014, **46**, 3004; (*d*) Z.-F. Xu, H. C. Dai, L. H. Shan and C.-Y. Li, Metal-Free Synthesis of (*E*)-Monofluoroenamine from 1-Sulfonyl-1,2,3-triazole and $Et_2O{\cdot}BF_3$ *via* Stereospecific Fluorination of α-Diazoimine, *Org. Lett.*, 2018, **20**, 1054; (*e*) D. Yang, L. Shan, Z.-F. Xu and C.-Y. Li, Metal-free synthesis of imidazole by $BF_3{\cdot}Et_2O$ promoted denitrogenative transannulation of *N*-sulfonyl-1,2,3-triazole, *Org. Biomol. Chem.*, 2018, **16**, 1461.

5 (*a*) V. Motornov, A. Markos and P. Beier, A rhodium-catalyzed transannulation of *N*-(per)fluoroalkyl-1,2,3-triazoles under microwave conditions – a general route to *N*-(per) fluoroalkyl-substituted five-membered heterocycles, *Chem. Commun.*, 2018, **54**, 3258; (*b*) V. Motornov and P. Beier, Chemoselective Aza-[4+3]-annulation of *N*-Perfluoroalkyl-1,2,3-triazoles with 1,3-Dienes: Access to *N*-Perfluoroalkyl-Substituted Azepines, *J. Org. Chem.*, 2018, **83**, 15195; (*c*) V. Motornov, V. Košťál, A. Markos, D. Taffner and P. Beier, General approach to 2-fluoroalkyl 1,3-azoles *via* the tandem ring opening and defluorinative annulation of *N*-fluoroalkyl-1,2,3-triazoles, *Org. Chem. Front.*, 2019, **6**, 3776; (*d*) O. Bakhanovich, V. Khutorianskyi, V. Motornov and P. Beier, Synthesis of *N*-perfluoroalkyl-3,4-disubstituted pyrroles by rhodium-catalyzed transannulation of *N*-fluoroalkyl-1,2,3-triazoles with terminal alkynes, *Beilstein J. Org. Chem.*, 2021, **17**, 504; (*e*) A. Markos, S. Voltrová, V. Motornov, D. Tichý, B. Klepetářová and P. Beier, Stereoselective Synthesis of (*Z*)-β-Enamido Triflates and Fluorosulfonates from *N*-Fluoroalkylated Triazoles, *Chem. – Eur. J.*, 2019, **25**, 7640; (*f*) A. Markos, L. Janecký,


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T. Chvojka, T. Martinek, H. Martinez-Seara, B. Klepetářová and P. Beier, Haloalkenyl Imidoyl Halides as Multifacial Substrates in the Stereoselective Synthesis of *N*-Alkenyl Compounds, *Adv. Synth. Catal.*, 2021, **363**, 3258.

6 (*a*) T. Opsomer and W. Dehaen, Metal-free syntheses of *N*-functionalized and NH-1,2,3-triazoles: an update on recent developments, *Chem. Commun.*, 2021, **57**, 1568; (*b*) C. P. Parambil, S. P. Veethil and W. Dehaen, The Synthesis of Five-Membered N-Heterocycles by Cycloaddition of Nitroalkenes with (In)Organic Azides and Other 1,3-Dipoles, *Synthesis*, 2022, **54**, 910.

7 Use of activated ketones: (*a*) R. Prakash, T. Opsomer and W. Dehaen, Triazolization of Enolizable Ketones with Primary Amines: A General Strategy toward Multifunctional 1,2,3-Triazoles, *Chem. Rec.*, 2021, **21**, 376; (*b*) J. John, J. Thomas and W. Dehaen, Organocatalytic routes toward substituted 1,2,3-triazoles, *Chem. Commun.*, 2015, **51**, 10797, Use of nitroalkenes. (*c*) V. A. Motornov, S. L. Ioffe and A. A. Tabolin, [3+ 2]-Annulation reactions with nitroalkenes in the synthesis of aromatic five-membered nitrogen heterocycles, *Targets Heterocycl. Syst.*, 2019, **23**, 237.

8 T. Jin, S. Kamijo and Y. Yamamoto, Copper-Catalyzed Synthesis of *N*-Unsubstituted 1,2,3-Triazoles from Nonactivated Terminal Alkynes, *Eur. J. Org Chem.*, 2004, 3789.

9 (*a*) R. Hui, M. Zhao, M. Chen, Z. Ren and Z. Guan, One-Pot Synthesis of 4-Aryl-NH-1,2,3-Triazoles through Three-Component Reaction of Aldehydes, Nitroalkanes and $NaN_3$, *Chin. J. Chem.*, 2017, **35**, 1808; (*b*) L. Wu, X. Wang, Y. Chen, Q. Huang, Q. Lin and M. Wu, 4-Aryl-NH-1,2,3-Triazoles *via* Multicomponent Reaction of Aldehydes, Nitroalkanes, and Sodium Azide, *Synlett*, 2016, **27**, 437. (*c*) Q. Hu, Y. Liu, X. Deng, Y. Li and Y. Chen, Aluminium(III) Chloride-Catalyzed Three-Component Condensation of Aromatic Aldehydes, Nitroalkanes and Sodium Azide for the Synthesis of 4-Aryl-NH-1,2,3-triazoles, *Adv. Synth. Catal.*, 2016, **358**, 1689. For "green" protocols to NH-1,2,3-triazoles, see; (*d*) A. Garg, R. Hazarika, N. Dutta, B. Dutta and D. Sarma, Bio-waste Derived Catalytic Approach Towards NH-1,2,3-Triazole Synthesis, *ChemistrySelect*, 2021, **6**, 7266; (*e*) A. Garg, D. Sarma and A. A. Ali, *Curr. Res. Green Sustainable Chem.*, 2020, **3**, 100013; (*f*) J. Sultana, A. Garg, A. Kulshrestha, S. S. Rohman, B. Dutta, K. Singh, A. Kumar, A. K. Guha and D. Sharma, *Catal. Lett.*, 2023, **153**, 3516; (*g*) P. Phukan, R. Chetia, R. Boruah, S. Konwer and D. Sharma, *Adv. Mater.*, 2021, **2**, 6996; (*h*) S. Bagheri, M. J. Nejad, F. Pazoki, M. K. Miraki and A. Heydari, *ChemistrySelect*, 2019, **4**, 11930.

10 D. Jankovic, M. Virant and M. Gazvoda, Copper-Catalyzed Azide–Alkyne Cycloaddition of Hydrazoic Acid Formed *In Situ* from Sodium Azide Affords 4-Monosubstituted-1,2,3-Triazoles, *J. Org. Chem.*, 2022, **87**, 4018.

11 (*a*) L. Hu, C. Muck-Lichtenfield, T. Wang, G. He, M. Gao and J. Zhao, Reaction between Azidyl Radicals and Alkynes: A Straightforward Approach to NH-1,2,3-Triazoles, *Chem. – Eur. J.*, 2016, **22**, 911; (*b*) S. Roshandel, S. C. Suri, J. C. Marcischak, G. Rasul and G. K. S. Prakash, Catalyst and solvent free microwave-assisted synthesis of substituted 1,2,3-triazoles, *Green Chem.*, 2018, **20**, 3700.

12 Benzaldehydes are by the order of magnitude less expensive than corresponding aryl acetylenes according to price lists of common chemical suppliers. Also, nitroalkanes are very inexpensive large scale industrial products. See: N. Ono, *The Nitro Group in Organic Synthesis*, John Wiley & Sons, New York, 2002.

13 S. W. Kwok, L. Zhang, N. P. Grimster and V. V. Fokin, Catalytic Asymmetric Transannulation of NH-1,2,3-Triazoles with Olefins, *Angew. Chem., Int. Ed.*, 2014, **53**, 3452.

14 V. Motornov, R. Pohl, B. Klepetářová and P. Beier, *N*-Acyl-1,2,3-triazoles – key intermediates in denitrogenative transformations, *Chem. Commun.*, 2023, **59**, 9364.

15 T. Wang, Z. Tang, H. Luo, Y. Tian, M. Xu, Q. Lu and B. Li, Access to (*Z*)-β-Substituted Enamides from *N*1-H-1,2,3-Triazoles, *Org. Lett.*, 2021, **23**, 6293.

16 (*a*) Q. Lu, T. Wang, Q. Wu, L. Cheng, H. Luo, L. Liu, G. Chu, L. Wang and B. Li, C–H heteroarylation of aromatics *via* catalyst free $S_N2'$ coupling cycloaromatization, *Green Chem.*, 2022, **24**, 4399; (*b*) T. Chvojka, A. Markos, S. Voltrová, R. Pohl and P. Beier, Ligand-dependent stereoselective Suzuki–Miyaura cross-coupling reactions of β-enamido triflates, *Beilstein J. Org. Chem.*, 2021, **17**, 2657.

17 V. Motornov and P. Beier, Access to Fluoroalkylated Azoles and 2-Acylaminoketones *via* Fluorinated Anhydride-Mediated Cleavage of NH-1,2,3-Triazoles, *Org. Lett.*, 2022, **24**, 1958.

18 V. Motornov and P. Beier, One-pot synthesis of 4-substituted 2-fluoroalkyloxazoles from NH-1,2,3-triazoles and fluoroalkylated acid anhydrides, *New J. Chem.*, 2022, **46**, 14318.

19 A. S. Singh, N. Mishra, D. Kumar and V. K. Tiwari, Lewis-Acid-Mediated Benzotriazole Ring Cleavage (BtRC) Strategy for the Synthesis of 2-Aryl Benzoxazoles from *N*-Acylbenzotriazoles, *ACS Omega*, 2017, **2**, 5044.

20 B. Dong, Y. Liu, P. Yang, D. Sang, J. Tian, L. Li and S. Long, Denitrogenative cleavage of benzotriazoles and benzotriazinones, and selective *N*-desulfonylation of benzotriazoles by aluminum halides, *Tetrahedron Lett.*, 2022, **103**, 153965.

21 V. Motornov and P. Beier, Synthesis of *N*-vinyl isothiocyanates and carbamates by the cleavage of NH-1,2,3-triazoles with one-carbon electrophiles, *Org. Biomol. Chem.*, 2023, **21**, 1147.

22 M. Xu, L. Liu, T. Wang, H. Luo, M. Hou, L. Du, X. Xin, Q. Lu and B. Li, Acid-catalyzed ring-expansion of 4-(1-hydroxycyclobutyl)-1,2,3-triazoles, *Org. Chem. Front.*, 2022, **9**, 1065.

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