

REVIEW

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Recent developments in dehydration of primary amides to nitriles

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Dehydration of amides is an efficient, clean and fundamental route for the syntheses of nitriles in organic chemistry. The two imperative functional groups viz., amide and nitrile groups have been extensively discussed in the literature. However the recent development in the century-old dehydration method for the conversion of amides to nitriles has hardly been reported in one place, except a lone review article which dealt with only metal catalysed conversions. The present review provides broad and rapid information on the different methods available for the nitrile synthesis through dehydration of amides. The review article has major focus on (i) non-catalyzed dehydrations using chemical reagents, and (ii) catalyzed dehydrations of amides using transition metal, non-transition metal, organo- and photo-catalysts to form the corresponding nitriles. Also, catalyzed dehydrations in the presence of acetonitrile and silyl compounds as dehydrating agents are highlighted.

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1. Introduction

Nitriles are naturally found in various bacteria, fungi, plants and animals and are extensively applied as raw materials for the production of a variety of pharmaceuticals, agrochemicals,

polymers, materials, *etc.*¹ Examples of pharmaceuticals containing nitrile groups include *vildagliptin*, an anti-diabetic drug² and *anastrozole*, a drug used for treating breast cancer (Fig. 1).³ Moreover, nitrile functionalities have been utilized as versatile intermediates in organic synthesis and they can be readily converted to various other important functional groups viz., aldehydes,⁴ carboxylic acids,⁵ esters,⁶ primary amines,⁷ imines,⁸ substituted oximes,⁹ heterocycles,¹⁰ amides,¹¹ *etc.*¹² Synthesis of such a highly valuable nitrile functionality has been one of the promising research fields in organic chemistry.¹³

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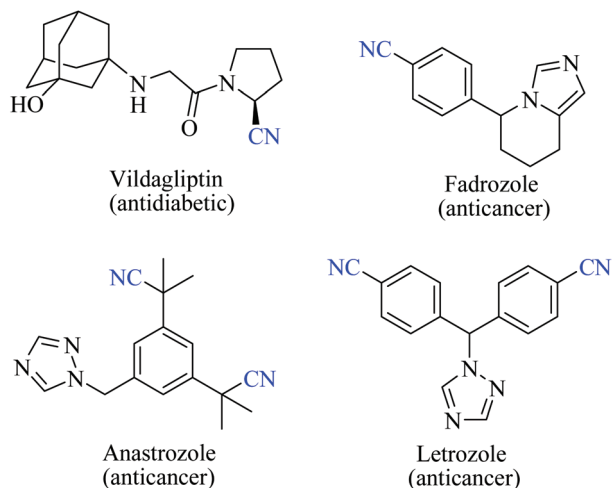
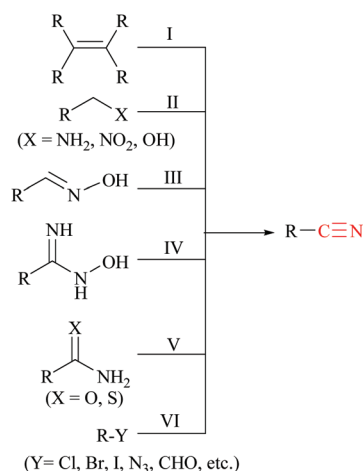


Fig. 1 Examples of $-C\equiv N$ containing drug molecules.

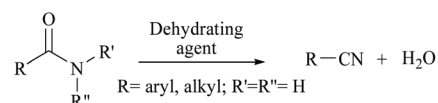
Various methods^{13,14} have been developed for the syntheses of nitriles using different synthetic strategies including the addition reaction of multiple bonds (I),¹⁵ oxidation of amines (II),¹⁶ reduction of nitroalkanes (II),¹⁷ reactions of alcohols (II),¹⁸ dehydration or rearrangement of oximes (III) and amidoximes (IV),¹⁹ dehydration of amides or water shuffling with amides (V),²⁰ dehydrosulfurization of thioamides (V),²¹ substitution of aryl and alkyl-halides/azides (VI),²² and reaction of aldehydes (VI)²³ (Scheme 1). The classical methods *viz.*, nucleophilic substitution of alkyl halides, the Sandmeyer reaction of diazotized aniline, the Rosenmund-von Braun reaction of aryl halides, *etc.*, have limited applications due to the usage of highly toxic cyanide sources.²⁴ On the other side, synthetic procedures using non-metallic cyanide sources²⁵ have also been available for the large-scale production of nitriles like acrylonitrile by ammoxidation of *n*-propene, adiponitrile by hydrocyanation of 1,3-butadiene, and so on.²⁶ However, these methods are also limited by their poor functional group toler-



Scheme 1 Various reported routes for nitrile synthesis.

ance due to harsh reaction conditions.²⁷ Alternatively, alkyl/aryl halides have also been catalytically converted to the corresponding amides using naturally abundant small molecules *viz.*, CO_2 , NH_3 and urea.²⁸ In fact, syntheses of nitriles from nitrogen containing compounds such as amine, amides, *etc.*, will be better alternatives than their non-nitrogen counterparts, as additional steps for the incorporation of the nitrogen functionality, involving the usage of stoichiometric reagents and generation of metal waste, can be avoided. Amides, apart from their easy availability, with a general formula of $RC(=O)NR'R''$, are commonly found in numerous synthetic and natural products, besides their crucial role as basic building blocks of proteins in every form of life.²⁹ Direct synthesis of nitriles by dehydration of amides with benign and low-cost catalysts and dehydrating agents under simple reaction conditions is particularly attractive from the green chemistry point of view. Furthermore, in the conversion of amides to nitriles, water molecules are the only value added side product which theoretically adds merit to the sustainable synthesis (Scheme 2). Thus, the dehydration of amides would be one of the fundamental and clean synthetic routes for the syntheses of nitriles.¹³ Simple heating at very high temperatures could lead to the dehydration of the amide groups to form the corresponding nitriles, whether catalysts and/or dehydrating agents are present or not. However, in order to overcome the problems associated with such harsh temperature, the dehydration methods were often carried out using stoichiometric amounts of dehydrating agents in the presence of catalysts, since the conversion of primary amides to nitriles by eliminating water is typically endergonic.¹³ Another interesting method for the preparation of nitriles from amides is "transfer hydration" which is based on the "water" acceptor-donor capacity of the substrate.¹⁶ Under this method, powerful water donors such as amides and aldoximes are dehydrated into nitriles in the presence of dehydrating agents such as acetonitrile. However, selectivity and excess use of the water donor are the major drawbacks of this method. Hence, it is a challenging goal to develop eco-friendly ways for the conversion of amides to the corresponding nitriles.

It is worth noting that recently, Al-Huniti and Croatt have reviewed the metal catalyzed conversion of amides to nitriles.¹³ However, our aim is to present broad and rapid information on the various available methods for the dehydration of amides to nitriles under non-catalyzed and catalyzed conditions. The catalytic dehydrations included for the discussion are based on transition metal, non-transition metal, organo- and photo-catalysts in the presence of acetonitrile and silyl compounds as dehydrating agents. It is hoped that by compiling the widely scattered information on the dehydration of



Scheme 2 Dehydration of amides leading to clean products.

amides, the present review will grasp the attention of a broad readership due to the potential applications of nitriles.

The present review shall describe the merits and demerits of the present methods and pertaining challenges for dehydration of amides by categorizing them into two parts: part I shall deal with the non-catalyzed dehydration methods using chemical reagents and in part II applications of catalysts in dehydration shall be discussed.

2. Non-catalyzed dehydration of amides using chemical reagents

The first dehydration of amides was performed by Wöhler and Liebig in 1832 who reported that pyrogenetic decomposition of benzamide over barium oxide yielded phenyl cyanide and water.³⁰ The first use of phosphorus pentoxide to dehydrate ammonium salts or amides was introduced by Dumas in 1847.³¹ However, chemical reagents either as dehydrating agents or as catalysts are usually required for the mild conversion of amides to nitriles. Thus, the direct preparation of nitriles was generally achieved by the dehydration of the corresponding amides using classical dehydrating reagents such as Lewis acids, basic reducing agents and other efficient chemical reagents that are discussed in the following sections.

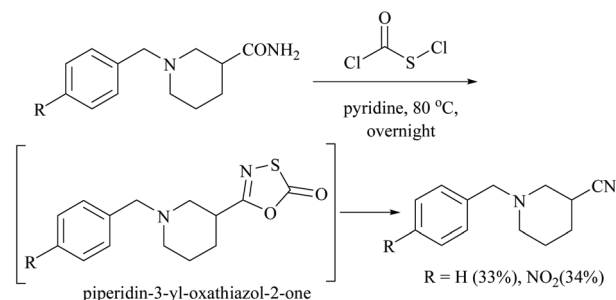
Rickborn and Jensen reported the dehydration of 4-*tert*-butylcyclohexanecarboxamide to the corresponding nitrile using phosphorus pentoxide (P_2O_5), thionyl chloride and also phosphorus oxychloride as dehydrating agents.³² Kenyon and Ross used P_2O_5 for the dehydration of 2-methyl-3-phenylpropanamide to obtain the corresponding nitrile.³³ Mahajan *et al.* reported the dehydration of 2-amino-5-chlorobenzamide using P_2O_5 to form 2-amino-5-chlorobenzonitrile and phosphoric acid (Scheme 3).³⁴

Konwar and coworkers developed a method for the dehydration of oximes and amides in hydrated media to their corresponding nitriles in good yields using an $AlCl_3 \cdot 6H_2O/KI/H_2O/CH_3CN$ system (Scheme 4).³⁵ The same group has also reported *anhydrous* $AlCl_3$ -KI mediated dehydration of amides. In the presence of water, aluminium is coordinated with

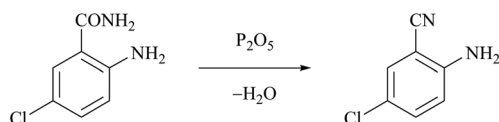
amide-nitrogen and in anhydrous medium it is coordinated with amide-oxygen and facilitates the removal of water from the amide to form the corresponding nitrile (Scheme 4).

In the preparation of oxathiazol-2-one, an inhibitor of threonine protease, using chlorocarbonylsulfonyl chloride as a coupling reactant and sodium carbonate as a base, the starting carboxamide in the presence of dioxane as solvent afforded the expected piperidin-3-yl-oxathiazol-2-one compound in 68% yield.³⁶ However, in the absence of sodium carbonate as a base, the reaction of carboxamide with chlorocarbonylsulfonyl chloride alone using pyridine as solvent initially involved coupling which is followed by a rapid elimination of sulphur oxide to form its corresponding nitrile. Thus the combination of chlorocarbonylsulfonyl chloride and pyridine constitutes a dehydrating system and facilitates the dehydration of the amide to give the corresponding nitrile by destabilizing the intermediate, piperidin-3-yl-oxathiazol-2-one (Scheme 5).³⁶

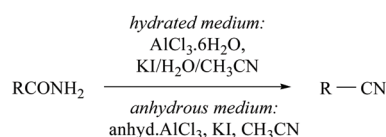
Unnikrishnan and coworkers disclosed a one-pot synthesis of nitriles from amides and oximes using an *in situ*-generated Burgess-type reagent as the dehydrating agent (Scheme 6). The treatment of chlorosulfonyl isocyanate with β -aminoalcohol [1-(2-hydroxyethyl)-piperidine] yielded in a single step a cyclic Burgess-type reagent which without further purification was used for the dehydration of aromatic amides.³⁷ Itabashi and coworkers reported the dehydration of benzamide, acetamide, thiobenzamide and nicotinamide using phosphorus tris(diethylamide) [$P(NEt_2)_3$] under reflux conditions in THF for



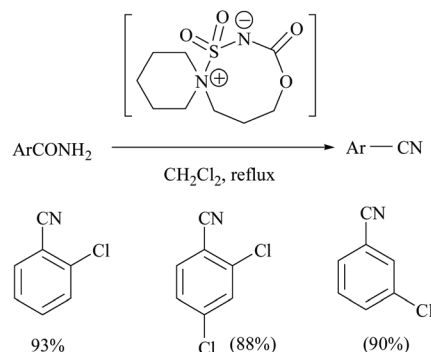
Scheme 5 Chlorocarbonylsulfonyl chloride and pyridine mediated dehydration.



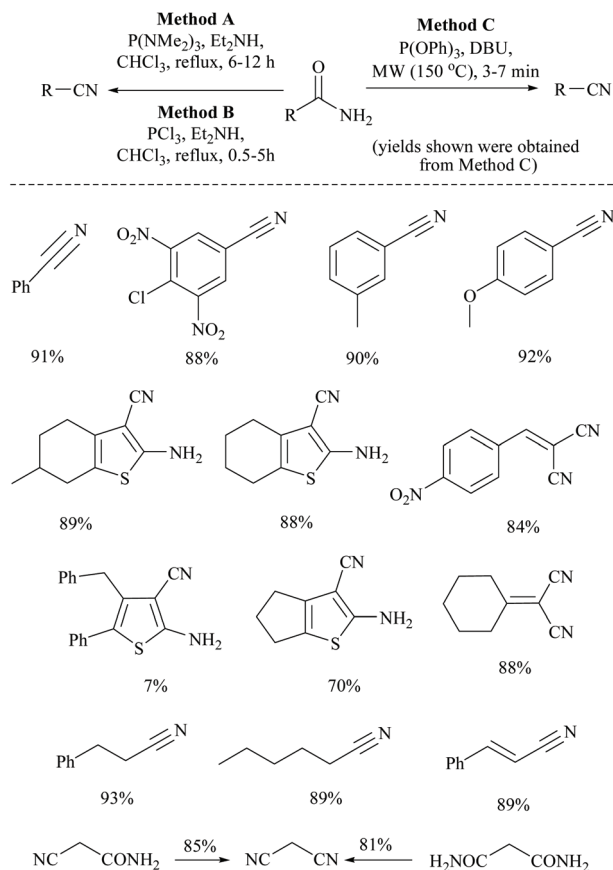
Scheme 3 P_2O_5 as a dehydrating agent.



Scheme 4 $AlCl_3$ as a dehydrating agent.



Scheme 6 Burgess-type reagent as the dehydrating agent.

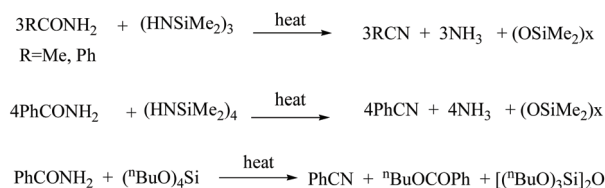


Scheme 7 Use of electrophilic phosphorus-III reagents.

3 hours, to give the corresponding nitriles in 89.6%, 98.5%, 90.4% (in benzene) and 70.5% yields, respectively (Scheme 6).³⁸

Touil and coworkers developed three different methods for the conversion of primary amides into the corresponding nitriles, mediated by the highly electrophilic phosphorus-III reagents, tris-(dimethylamino)phosphine [P(NMe₂)₃], phosphorus trichloride (PCl₃), or triphenylphosphite [P(OPh)₃], respectively (Scheme 7).³⁹ The primary amides coupled with P(NMe₂)₃, PCl₃, or P(OPh)₃ followed by rapid elimination with a base: diethylamine or DBU to form the corresponding nitriles under MW irradiation or reflux conditions. Among these three methods, method C with P(OPh)₃ and DBU under MW irradiation exhibited the best result. Notably, aromatic amides bearing electron-donating groups on the aromatic rings produced higher yields and the reaction conditions were found to be compatible with cyanoamides, malonamides, α,β-unsaturated amides, an unprotected amine functionality, the existing nitrile functionality, and a thiophene ring.

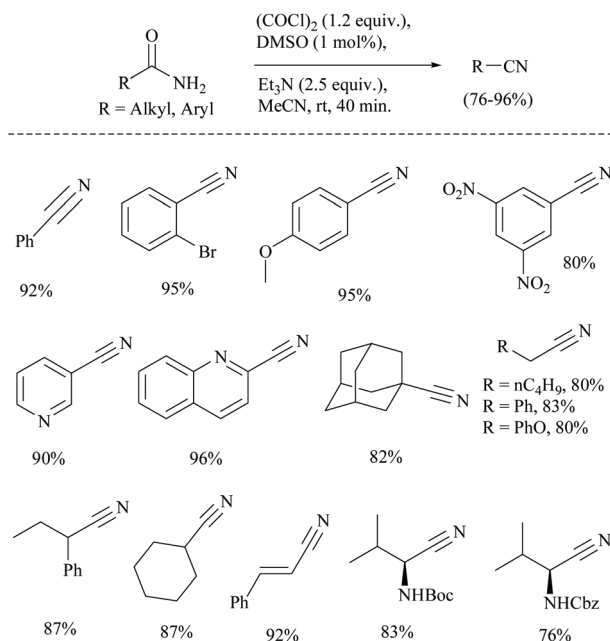
Dennis reported the dehydration of amides to nitriles with the help of silazanes, aminosilanes, alkoxy silanes, and chlorosilanes as dehydrating agents at high temperatures.⁴⁰ For example, hexamethyltricyclo-trisilazane with acetamide or benzamide underwent dehydration at a temperature range of 180–200 °C, forming acetonitrile and benzonitrile in 95 and 85% yields, respectively (Scheme 8). Moreover, with benz-



Scheme 8 Silyl compounds as dehydrating agents.

amide, silazanes formed a siloxane polymer of molecular weight *M_w* 120 000 as the by-product. Under these conditions, *N*-methylbenzamide also gave the benzonitrile when heated to above 200 °C.

The conversion of primary amides into nitriles has also been found to use oxalyl chloride as an activating agent only along with other reagents.⁴¹ In 1978, Swern used for the first time a combination of DMSO and oxalyl chloride for the oxidation of primary and secondary alcohols.⁴² The combination has been used for many other conversions.⁴³ Sun *et al.* wished to use the Swern reaction conditions catalytically for the dehydration of amides which often involve the usage of stoichiometric amounts of DMSO and (COCl)₂.⁴¹ The authors successfully carried out the catalytic dehydration of amides to obtain nitriles in good yields, using a catalytic amount of DMSO in the presence of (COCl)₂ and Et₃N (Scheme 9). The proposed mechanism for the catalytic Swern oxidation involved initial formation of chlorodimethylsulfonium chloride from the reaction of DMSO and (COCl)₂, followed by nucleophilic substitution of amide-oxygen in the presence of the Et₃N-base. In the last step, elimination of HCl and DMSO led to the formation of the corresponding nitrile. The replacement of the base Et₃N with other bases such as DBU and NMM was not



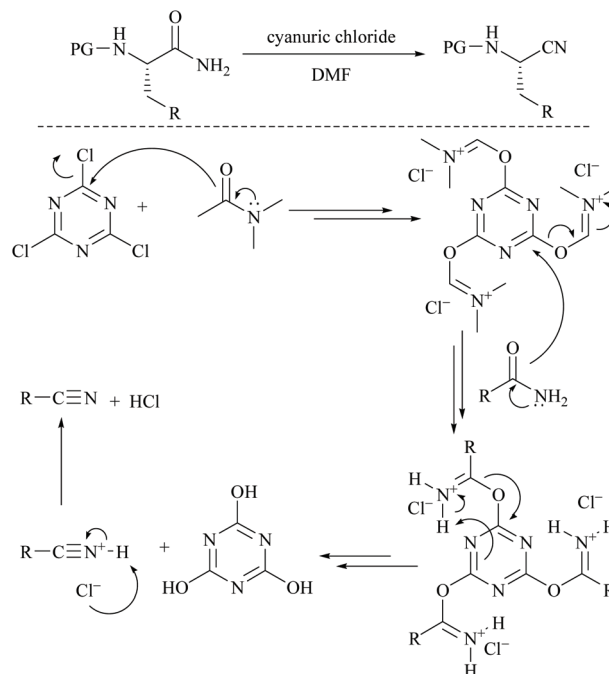
Scheme 9 Oxalyl chloride with a DMSO catalyst.

successful. Notably, the yields obtained were found to be comparable with those of the catalytic Appel type reaction (discussed in a later section). The Boc protected substrate was found to be compatible under catalytic Swern oxidation, whereas under catalytic Appel's reaction, Boc-protected amino acids did not undergo the conversion. Moreover, the present method was superior in terms of higher yield than the one reported by Nakajima and Ubukata⁴¹ using Swern oxidation with stoichiometric DMSO.

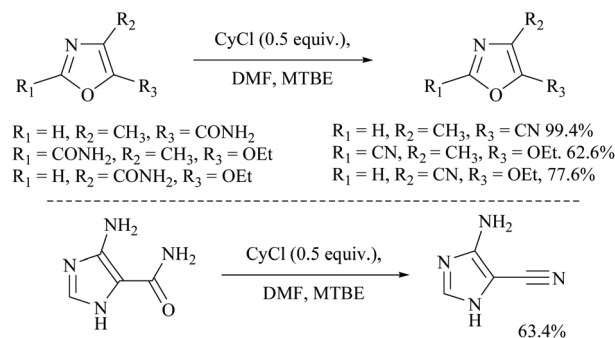
Monson and Priest reported the dehydration of aliphatic and aromatic carboxamides to their corresponding nitriles in hexamethylphosphoric triamide (HMPT) at 220–240 °C in good yields *viz.*, propionamide (94%), *n*-butyramide (75%), hexanamide (78%), benzamide (67%), phenylacetamide (75%) and adipamide (49%) (Scheme 10).⁴⁴ The plausible mechanism for this reaction involves the evolution of dimethylamine, the formation of a phosphorodiamidate derivative of the amide, and the elimination of the phosphorodiamidate leaving group to form the nitrile (Scheme 10).

Cyanuric chloride (2,4,6-trichloro-*s*-triazine) has also been used as a dehydrating agent for carboxamide. Under these reaction conditions, only a one third molar equivalent of cyanuric chloride is needed to convert an amide group into its nitrile.⁴⁵ The mechanism involves initially simple nucleophilic substitution of chloride in cyanuric chloride with formamide which is then substituted by carboxamide. For example, Rodriguez and coworkers reported the dehydration of *N*-protected α -amino-carboxamides to their corresponding nitriles in good yields using cyanuric chloride in DMF as the dehydrating agent (Scheme 11).⁴⁶ Using similar conditions, Bonrath and coworkers reported the dehydration of heterocyclic carboxamides to the corresponding nitriles (Scheme 12). The oxazole amides underwent dehydration smoothly to form their corresponding nitriles.⁴⁷ They also found that the nature of the formamide is not critical and any conventional *N,N*-disubstituted formamide can be used but the formamide must be water insoluble. However, nicotinic acid amide did not react under these conditions to 3-cyano-pyridine.

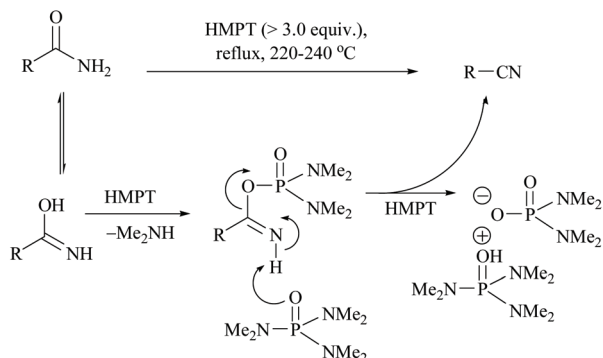
Shia and coworkers disclosed a simple procedure for the conversion of primary amides to the corresponding nitriles, using ethyl dichlorophosphate as the dehydrating agent along



Scheme 11 Cyanuric chloride mediated dehydration.



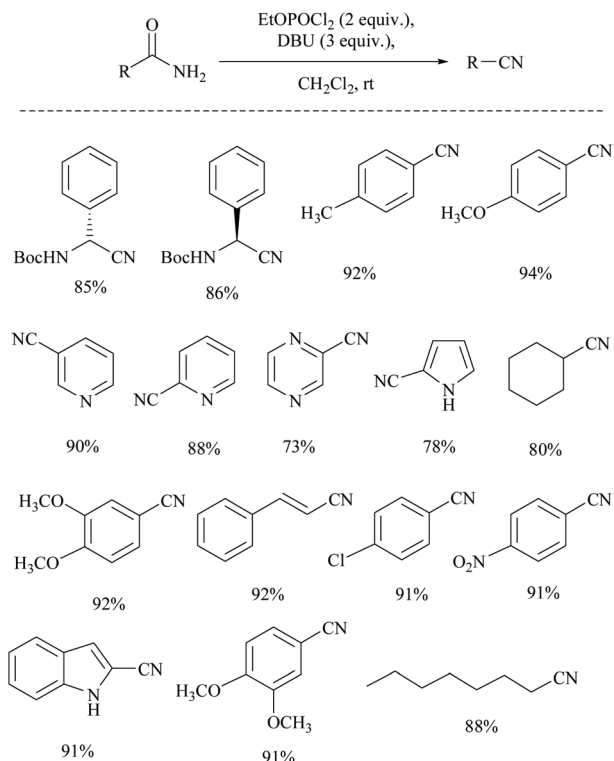
Scheme 12 CyCl with MTBE for dehydration.



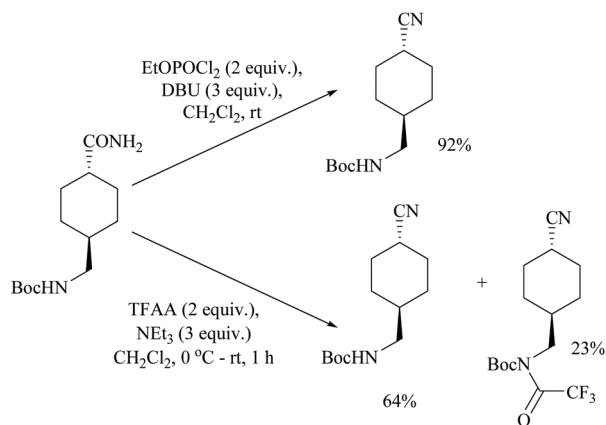
Scheme 10 HMPT mediated dehydration.

with DBU (Scheme 13).⁴⁸ Moreover, for comparison, the authors also carried out the dehydration of cyclohexanyl carboxamide distinctly using their own reagent system ethyl dichlorophosphate/DBU as well as with a trifluoroacetic anhydride (TFAA)/Et₃N system which had already been reported for the dehydration of primary amides.⁴⁹ (Scheme 14). They observed that their reagent afforded the nitrile in 92% yield; however with the TFAA system the nitrile was formed in only 64% yield along with trifluoro-acylated nitrile as a side product in 23% yield.

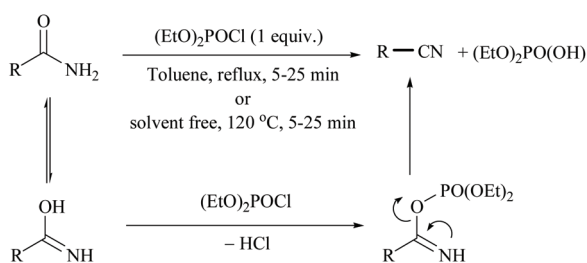
In 2011, Sardarian and coworker⁵⁰ reported a better method using “diethyl chlorophosphate” (EtO)₂POCl as a dehydrating agent (Scheme 15), compared to the previous one with “ethyl dichlorophosphate” (Scheme 13).^{48,49} Moreover, the present method has an advantage that the dehydration could also be carried out in the absence of a solvent and base, without compromising the yield (Table 1).



Scheme 13 Ethyl dichlorophosphate as a dehydrating agent.



Scheme 14 Ethyl dichlorophosphate vs. TFAA.



Scheme 15 Diethyl chlorophosphate as a dehydrating agent.

Table 1 (EtO)₂POCl vs. (EtO)POCl₂: comparison of reaction time and yields^{48–50}

R	(EtO) ₂ POCl				(EtO)POCl ₂	
	Toluene		Solvent free		Yield, %	Time, min
R = H	Yield, %	Time, min	Yield, %	Time, min	Yield, %	Time, min
R = <i>p</i> -Me	83	20	89	15	98	60
R = <i>p</i> -OMe	82	10	90	15	92	180
R = <i>p</i> -OMe	85	5	91	5	94	180
R = <i>p</i> -NO ₂	89	15	91	25	93	120
Ph-CH=CH-CN	92	10	94	10	92	180
Cyclohexyl-CN	72	10	74	10	80	180

The proposed mechanism involves the reaction of the imidic form of primary amide with diethyl chlorophosphate to form the corresponding diethyl phosphate imide followed by the release of diethyl hydrogen phosphoric acid and nitriles. In order to study the effect of solvent, they carried out the dehydration in toluene as solvent under reflux conditions and the solvent method was found to produce relatively low yields. Notably, 2-cyanopyridine (2-CP) has been used as a dehydration agent in the synthesis of dimethylcarbonate (DMC) from CO₂ and methanol.⁵¹ In order to improve the recyclability, the end product, 2-picolinamide (2-PA), was dehydrated back to 2-CP. In this context, many attempts have been made to achieve the dehydration of 2-PA to 2-CP.^{49,52,53} Some of them such as ethyldichlorophosphate,⁴⁹ (*N*-methyl-*N*-trimethylsilyl) trifluoroacetamide,⁵² trifluoromethane sulfonic acid anhydride and trimethylamine⁵³ are non-renewable and expensive too.

3. Catalyzed dehydration of amides

Recent years have seen an upsurge in the publications of catalytic dehydrations using a variety of metals and organic small molecules as catalysts in the presence of dehydrating agents. Transition metal catalyzed amide-dehydration reactions typically utilize acetonitrile^{41,54} or silane^{55,56} including *N*-methyl-*N*-(trimethylsilyl)trifluoroacetamide (MSTFA), even though a very high temperature can also cause dehydration but with poor yield. A catalyzed water shuffling method between amides and nitriles has also been gaining reasonable attention for converting amides to nitriles and *vice versa*.⁵⁷ Metal based catalysts reported for the dehydration of primary amides to nitriles include complexes of transition metals *viz.*, Cu,^{41,58} In,⁵⁹ Fe,^{56,60} Pd,⁶¹ Sn,¹⁹ Re,⁶² Rh,⁶³ Ru,^{56,64} U,⁵² V,⁶⁵ W,^{52,66} Zn,^{52,66,67} and Zr.⁶⁸ The available literature reports are reviewed on the basis of the catalyst/dehydrating agent: (1) catalyzed dehydration at high temperature; (2) catalyzed dehydration using acetonitrile as a dehydrating agent, (3) transition metal catalyzed silylative dehydration, and (4) non-transition metal catalyzed, (5) organo-catalyzed, and (6) photocatalyzed dehydration.

3.1. Metal catalysed dehydration at high temperature

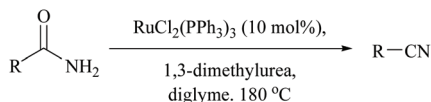
Dehydration of amides in the absence of a dehydrating agent, at high temperature either with or without a catalyst, has been found to give the corresponding nitrile. However, in order to improve the yield, to make it suitable for thermally unstable amides possessing other functional groups, and to avoid the usage of stoichiometric reagents including dehydrating agents, it is wise to make use of catalysts. Catalyzed dehydrations by bringing down the actual required temperature and time are found to be compatible for a variety of amide substrates. One such condition was accomplished by Campbell and coworkers for non-catalyzed dehydration of primary amides under flash vacuum pyrolysis (FVP) conditions requiring heating to >800 °C; however the addition of WO_3 to the furnace tube reduced the temperature by half to 400 °C for yielding benzonitrile in 97%.^{52,66}

Indeed, the first method in this context, using Wilkinson's heterogeneous catalyst, $\text{RhCl}(\text{PPh}_3)_3$, also required a very high temperature to remove the water molecules from the amide to form benzonitrile.⁶³ Also, under such harsh conditions, dehalogenation was observed with *p*- and *m*-chlorobenzamides. Thus, Watanabe *et al.* replaced the catalyst with a ruthenium (II) complex for the dehydration of the amide using, notably, an equimolar amount of *N,N'*-dialkylurea which gets converted into isocyanate and finally into alkyl amines and CO_2 (Scheme 16).⁶⁴

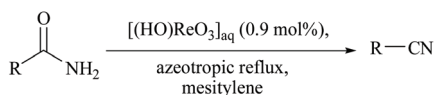
Yamamoto and coworkers reported a functional group-tolerable and large-scalable method using ~1 mol% of perhenic acid under azeotropic reflux of mesitylene for the dehydration of amides including vinyl and aromatic amides (Scheme 17).⁵⁸

Kaneda and coworkers reported that monomeric vanadium oxide supported on hydrotalcite (V/HT) acted as a reusable solid heterogeneous catalyst for the dehydration of amides into the corresponding nitriles. Under these reaction conditions, 16 different nitriles were obtained in good yields (83–94%) (Scheme 18).⁶⁵

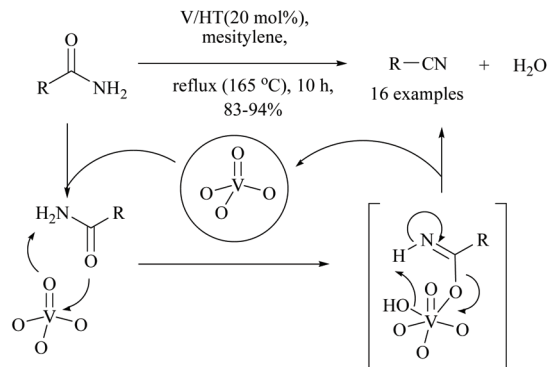
In 2004, Bergman and Ruck⁶⁸ reported a zirconium-mediated conversion of amides to the corresponding nitriles and thus joined the club consisting of the lone transition metal-mediated dehydration by TiCl_4 ⁶⁹ (Scheme 19). In the present method, primary amides were converted into their corresponding nitriles through the formation of the corresponding *N*-acylimidozirconocene complex.



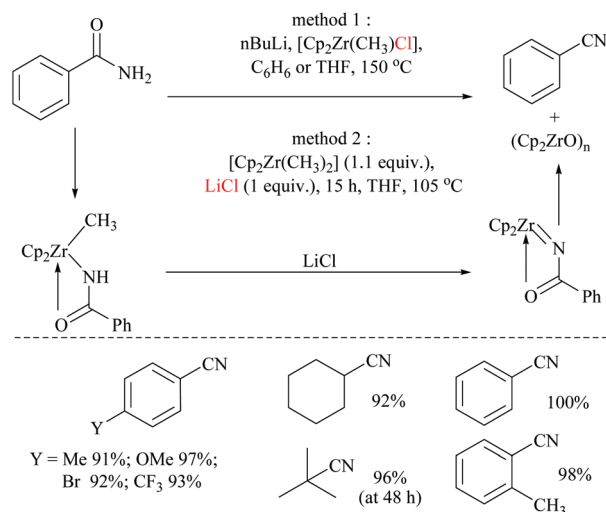
Scheme 16 Ruthenium(II) complex for dehydration.



Scheme 17 Perrhenic acid mediated dehydration.



Scheme 18 Vanadium catalyst for dehydration.



Scheme 19 Zirconium catalyst with LiCl for dehydration.

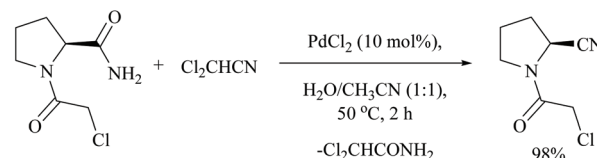
Dimethylzirconocene reacted with amides to form a methyl-zirconium amide complex. This methyl-zirconium amide complex, with the help of a chloride source, underwent 1,2-hydro methyl elimination to form an *N*-acylimidozirconocene complex which reacted intra-molecularly to give the corresponding nitrile (Scheme 19). Mechanistic studies reveal that chloride coordination to zirconium is required for this transformation to occur which was proved by carrying out dehydration with two different methods. In method 1, $[\text{Cp}_2\text{Zr}(\text{CH}_3)\text{Cl}]$ with an inbuilt chloride neatly formed the corresponding nitrile. However in method 2, with dimethylzirconocene, initially *N*-benzoylimidozirconocene was formed and left intact even after prolonged heating. The formed *N*-benzoylimidozirconocene continued to react only after the external addition of LiCl to form an *N*-benzoylimidozirconocene complex. In method 1, the residual LiCl by-product formed from the reaction between the lithium salt of benzamide and $[\text{Cp}_2\text{Zr}(\text{CH}_3)\text{Cl}]$. These observations indicated that the dehydration was dependent upon the chloride-anion additive effect.

3.2. Metal catalysed dehydration using acetonitrile as a dehydrating agent

Acetonitrile, the relatively inert solvent, gets activated in the presence of a metal catalyst and acts as a dehydrating agent for the conversion of primary amides to form the corresponding nitriles along with an equimolar amount of acetamide. However, dehydrations using nitriles as solvents are typically reversible, and the catalyst can promote the reaction in either direction depending on the type of solvent involved in that direction. Palladium-catalyzed dehydration of amides using acetonitrile as a dehydrating agent is known for its chemo-selectivity and compatibility over many other functionalities.^{54,61,70}

In 2005, Maffioli *et al.* reported for the first time PdCl₂ catalysed reversible conversion of primary amides to nitriles in aqueous acetonitrile solvent (Scheme 20).^{61,70} The dehydration of primary amides involved “water transfer” from the amide to acetonitrile. They revealed three important findings: (i) the dehydration of amides using a palladium-catalyst requires the addition of water; no reaction occurs under anhydrous conditions, (ii) water was not generated during the reaction but transferred directly from the amide to acetonitrile, and (iii) PdCl₂ and Pd(OAc)₂ alter the pH of the reaction solutions to the ranges 2.8–3.5 and 4.7–5.0, respectively; thus Pd(OAc)₂ can be a better choice of catalyst for the substrates which are sensitive to an acidic environment. The mechanism of palladium catalyzed dehydration involves the coordination of the palladium metal with nitrogen, and a H⁺-shift between the amide and nitrile (Scheme 20). The reversibility of this reaction *i.e.*, conversion of the nitrile to amide by hydration under the same reaction conditions was validated. Benzonitrile (PhCN) was allowed to react with MeCONH₂ under the same reaction conditions in which PhCN was used as a co-solvent instead of MeCN along with water. For making a homogeneous solution, a small amount of THF was also added and complete hydration of PhCN to benzamide was observed in 6 hours at room temperature.

In order to drive the reversibility of this reaction towards the complete conversion of amides to nitriles, Okabe *et al.* very recently carried out the palladium-catalyzed dehydration of primary amides to nitriles using dichloroacetonitrile as a water acceptor (Scheme 21).⁷¹ The authors initially searched for a

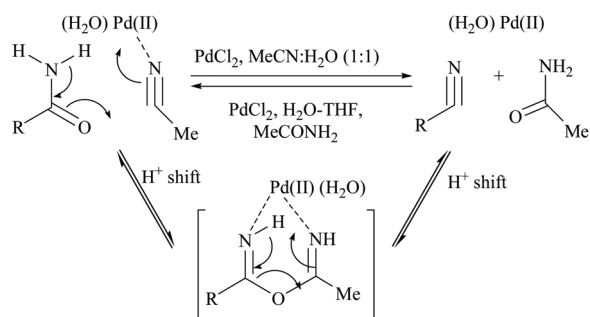


Scheme 21 Dichloroacetonitrile as a water acceptor.

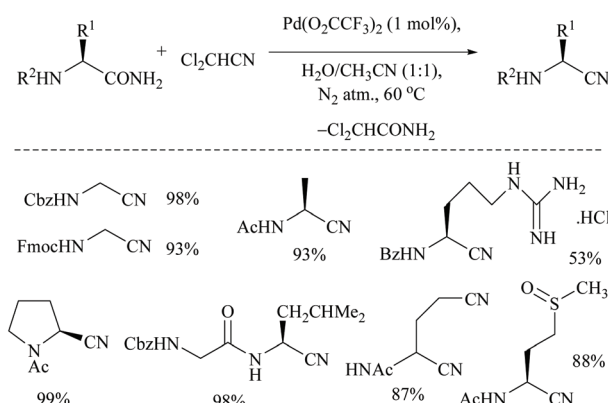
better water acceptor than MeCN in order to overcome the fact that the dehydration of electron-deficient amides with MeCN is energetically unfavourable. In this context, they focused on the use of electron-deficient nitriles such as dichloroacetonitrile as water acceptors to make the dehydration thermodynamically more favourable and also to obtain nitriles with the retention of the stereochemistry. The driving force to push the reaction towards nitrile formation is the energy required for hydration of the water acceptor. It was found that the hydration of dichloroacetonitrile was more exergonic than that of acetonitrile.

Using dichloroacetonitrile, a palladium trifluoroacetate-catalyzed dehydration method was effectively carried out for various α -aminoamides derived from natural amino acids in which many protecting groups were found to be intact. Notably, only the amide group with free –CONH₂ gets dehydrated under these conditions (Scheme 22).

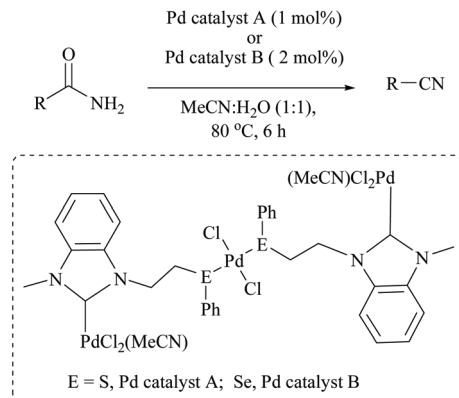
Singh and coworkers used a very low catalyst loading (0.5 mol%) of a trinuclear palladium(II) complex in aqueous acetonitrile solvent for the dehydration of amides to the corresponding nitriles in good yields (72–96%) (Scheme 23).⁷⁰ They used two NHC-Pd complexes with NHC containing soft donors *viz.*, sulphur and selenium and found that Pd catalyst A (with S) was more efficient than Pd catalyst B. They found that the reaction conditions were compatible with aliphatic amide, valeramide, pyridyl and aryl groups with various halogen substituents. During the optimization the authors screened two other catalysts *viz.*, Pd(CH₃CN)₂Cl₂ and Pd(PhS(CH₂)₂SPh)Cl₂, and even with 1 mol% loading these failed to achieve conversion, whereas in contrast, Maffioli and coworkers succeeded with 0.5 mol% for PdCl₂. Since direct comparison is not avail-



Scheme 20 Palladium catalyst for dehydration.



Scheme 22 Dehydration of α -aminoamides.



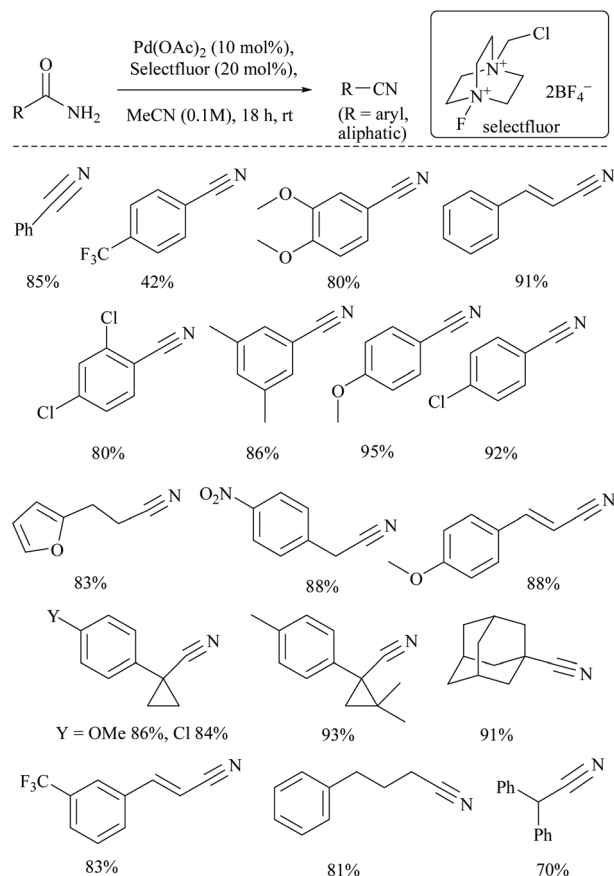
Scheme 23 Palladium catalyst for dehydration.

able, we shall assume that this discrepancy in the observation of two groups is due to the different sources of Pd(II), PdCl₂ versus Pd(CH₃CN)₂Cl₂.

Croatt and coworkers, while carrying out their research on improving the biological activities of Alamethicin F50, a 20-mer peptaibol, by fluorination using Pd(OAc)₂ and Selectfluor®. However, instead of fluorination, only dehydration of amide functionalities to form the corresponding tri-cyano product, along with semi-dehydrated analogues *viz.*, mono- and di-cyano products, was detected (Scheme 24).⁷⁰ This observation led them to explore further the application of this catalytic system to alkyl and aryl primary amides to obtain their corresponding nitriles in good yields. The role of the additive (20 mol% Selectfluor® in acetonitrile) was to act as an oxidant and to avoid the requirement of water for palladium catalysed dehydrations. The added Selectfluor® modified the Pd-catalytic pathway that generally uses either Pd(0) or Pd(II), by the formation of Pd(IV) which no longer requires the addition of water. Several non-, mono-, and disubstituted benzamides and aliphatic amides were converted to their corresponding nitriles in good yields. Notably, under these catalytic conditions, the cyclopropyl moieties were found to be intact.

Dai and coworkers reported two different palladium-catalyzed reaction conditions using non-aqueous acetonitrile for the conversion of α,β,γ,δ-unsaturated amides (Scheme 25) and aryl-alkyl amides (Scheme 26) into the corresponding nitriles.⁵⁴ The general requirement of water for palladium-catalysed dehydration of amides was exempted by the addition of metal acetate, which takes different pathways to assist the palladium catalyst. However, metal acetate alone without the palladium catalyst did not result in the conversion. The optimized reaction conditions for the conversion of unsaturated amides were achieved with PdCl₂, LiCl, and AgOAc in MeCN, whereas for the alkyl and aryl amides, the conditions with PdCl₂, (PPh₃)₂, and Cu(OAc)₂ in MeCN were used.

Notably, both reaction conditions were found to be successful at moderately high temperatures (80 °C). In general, the role of AgOAc is to regenerate Pd(II) from Pd(0) in the catalytic cycle. However, in the present method, AgOAc and Cu(OAc)₂



Scheme 24 Palladium catalyst with Selectfluor®.

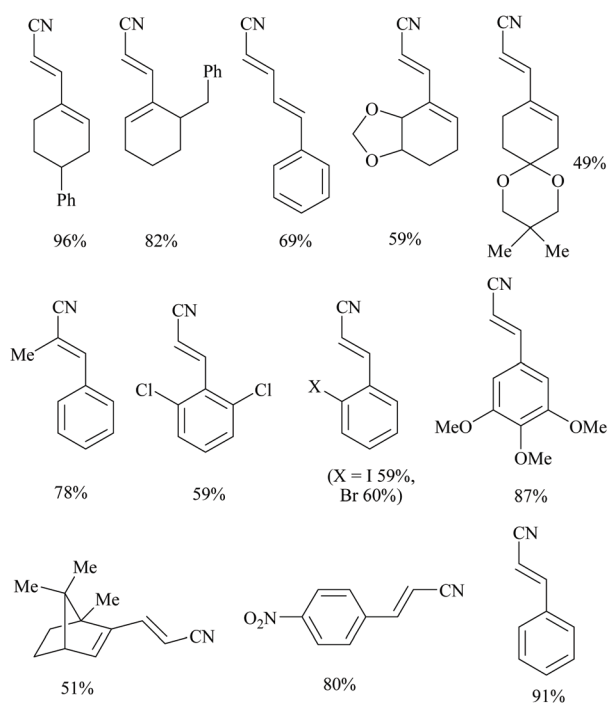
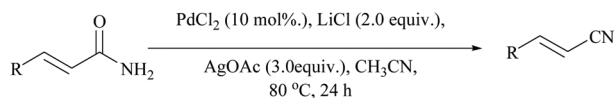
are merely the source of acetate counter anions which can be realized from the observation that the replacement of AgOAc with NaOAc did not affect the yield very much. But, the presence of acetate is highly required for the conversion, as it acts as a base throughout the mechanism. Being a dehydrating agent, when MeCN was replaced with DMF or toluene, it reduced the yield to less than 15%. Moreover, during the dehydration, MeCN is present as solvent in excess so as to drive the equilibrium towards the nitrile product formation by getting converted itself into acetamide.

A rapid method for the conversion of primary amides to nitriles and *vice versa* using ZnCl₂ and a ZnCl₂-H₂O-THF system in the presence of aqueous acetonitrile and acetamide, respectively, under microwave irradiation was reported by Manjula and Pasha (Scheme 27).⁶⁷ Zn(II) facilitates the transfer of water from the amide to acetonitrile and *vice versa*.

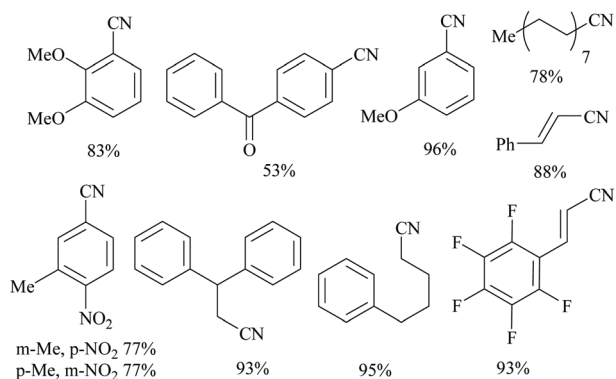
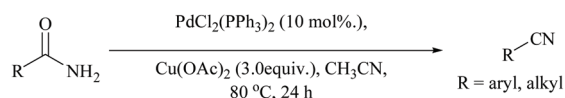
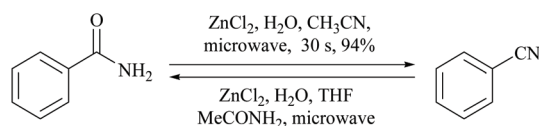
3.3. Catalyzed silylative dehydration – theoretical aspects

In silylative dehydrations of primary amides, transition metals get inserted into the Si-H bond and form hydrogen gas and silyl ether (Scheme 28). Indeed, silicon compounds have been used as silylating reagents, and they actively induce dehydration of the amides only in the presence of a catalyst.

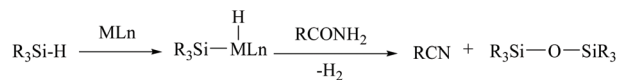
However, there have been recent reports that either the *high* energy barrier *syn*-elimination or *low* energy barrier



Scheme 25 Palladium catalyst with Ag(OAc) for dehydration.

Scheme 26 Palladium catalyst with Cu(OAc)₂ for dehydration.

Scheme 27 Zinc catalyst for dehydration.

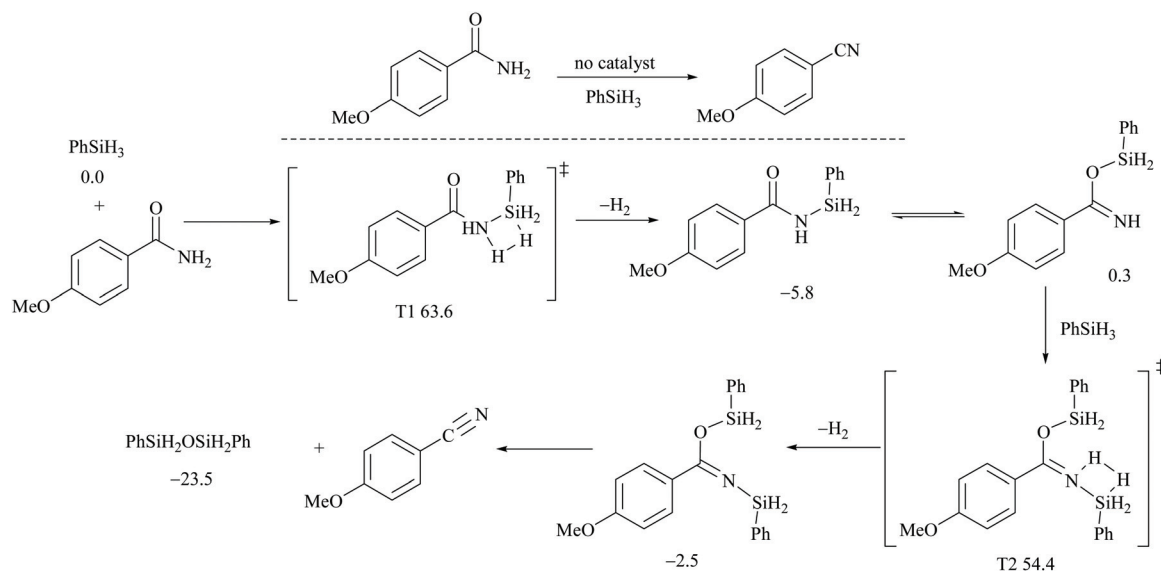


Scheme 28 Silicon-based dehydration of amides.

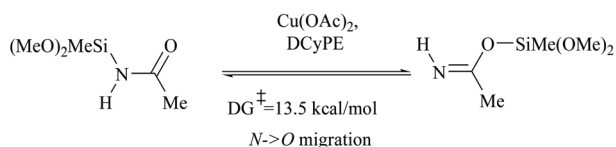
β -elimination in the presence of catalysts was the responsible pathway for the silylative dehydration to occur. Before reviewing the literature available for the metal catalyzed silylative dehydration of carboxamide to form the corresponding nitrile, we believe that knowing the mechanism of amide-dehydration will be very useful in designing the reaction conditions. In addition, Density Functional Theory (DFT) studies are found to be very helpful in determining the structure–reactivity relationships for the catalytic dehydration of primary amides using silanes. Mandal and coworkers⁷² reported the DFT calculations for amide to nitrile conversion using dehydrating agents such as phenyl silane. They compared the energy profile of the catalyzed reaction with that of the non-catalyzed one by taking *p*-methoxy benzamide as an example. Uncatalyzed silylative dehydration undergoes *N*-silylation, *N*- to *O*-silyl migration, a second *N*-silylation to form *N,O*-disilylimide, and finally elimination of silylether to form the nitrile (Scheme 29). Each silylation is followed by dehydrogenation to give relatively low energy silylated imide intermediates. Notably, both the silylation and elimination of silylether were accompanied by large energy barriers (Scheme 29). Hence, the addition of a catalyst can greatly help in the silylative dehydration process. For example, aNHC catalyzed mono-silylation and *N*- to *O*-silyl migration have energy barriers of 27.0 (T1, Scheme 32) and 15.7 kcal mol^{−1} (calculated), respectively, which were found to be lower than those for uncatalyzed conditions. Buchwald's group⁵⁸ also obtained similar results using a copper catalyst that *N*- to *O*-silyl migration needs only 13.5 kcal mol^{−1}; hence *O*-silyl and *N*-silyl imide forms equilibrium rapidly at room temperature (Scheme 30).

The catalyzed silylative dehydration follows either *syn*(1,2)-elimination or β -elimination. The *syn*-elimination pathway involves mono-*N*-silylation, *N*- to *O*-silyl migration, a second *N*-silylation to form *N,O*-disilylimide, and finally *syn*-elimination of silylether to form the nitrile with the regeneration of the catalyst (Scheme 32). In the case of the β -elimination pathway, differently, the amide substrate underwent mono-*N*-silylation, *N*- to *O*-silyl migration, and finally β -elimination of the catalyst along with silyl oxide as an ion-pair to form the nitrile. In order to regenerate the catalyst, a second silylation is required to dissociate the ion-pair into silyl ether (Scheme 32). It is interesting to note that in the catalyzed dehydrations, the addition of a second mole of silane is entirely utilized for regenerating the catalyst and eliminating the silylether. Thus, the role of a catalyst can be observed until the second silylation which substitutes the catalyst.

DFT calculations for the copper catalyzed elimination process in disilyl imide derived from the reaction of acetamide with DMMS showed that the barrier for *syn*-elimination



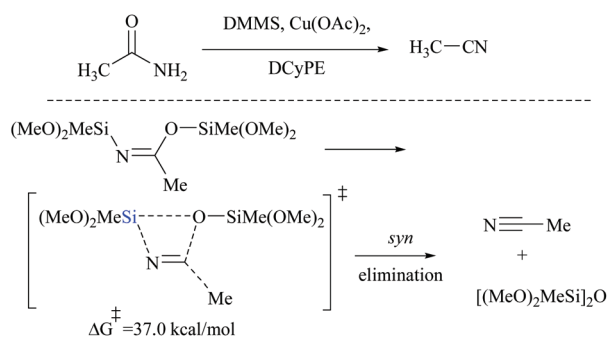
Scheme 29 DFT calculations and Gibbs free energy ΔG (kcal mol^{-1}): uncatalyzed dehydration.



Scheme 30 DFT calculations: Cu catalysed interconversion of *N*-silyl and *O*-silyl imidate forms.

was too high to be surmountable at ambient temperature ($+37.0 \text{ kcal mol}^{-1}$, Scheme 31). Also, the formation of *N,O*-disilylimidate itself requires a high energy barrier to be surmounted ($37.7 \text{ kcal mol}^{-1}$), making *syn*-elimination energetically least viable and therefore not feasible.

Appreciably, for the β -elimination, aNHC and copper/DCyPE catalysts have low energy barriers of 6.6 and 4.8 kcal mol^{-1} , respectively (Schemes 32 and 33). In copper catalysed dehydration, β -elimination of the silyl group was facilitated by reduced strain in the four-membered transition state (Scheme 33). This can be explained with the help of the differ-



Scheme 31 DFT calculations: Cu-catalyzed *syn*-elimination.

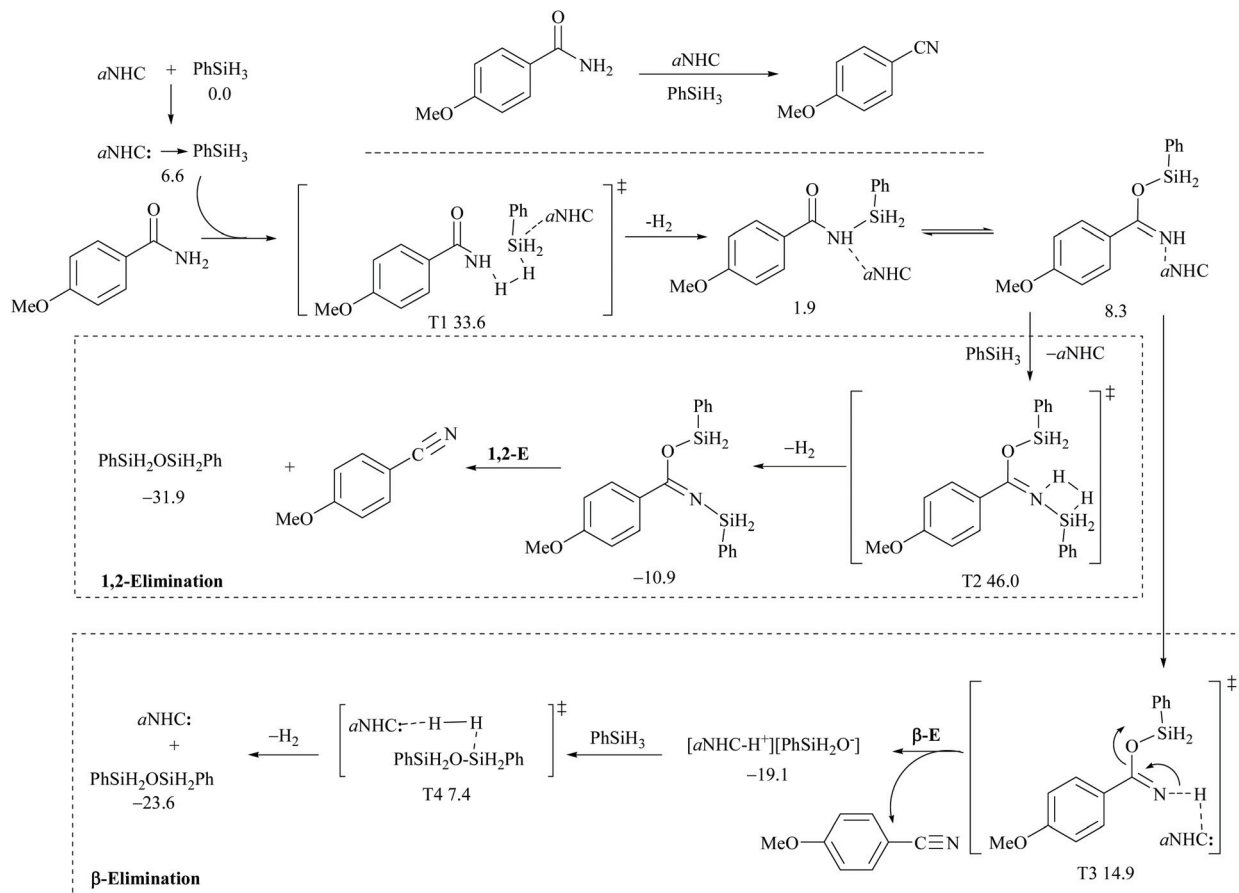
ence in bond distances in their respective transition states. The β -elimination has a transition state with bond lengths, *viz.* Cu–O and Cu–N bond lengths (251 and 199 pm), longer than those of the *syn*-elimination transition state (Si–O, Si–N: 187, 200 pm). Due to the larger size of Cu compared to Si, the reduced strain and the thermodynamically favourable $\text{C}\equiv\text{N}$ bond make the β -elimination comparatively easy and rapid.

3.4. Metal catalysed silylative dehydration

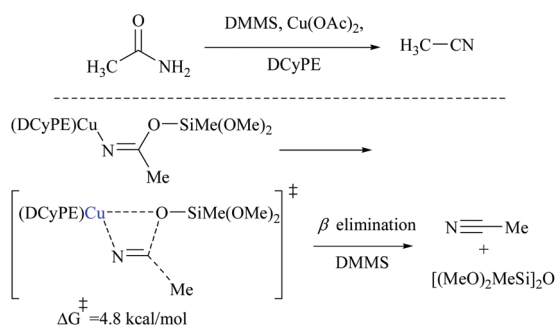
In 2008, Nagashima and coworkers reported the first silicon-based dehydration of primary amides using a triruthenium carbonyl cluster (Scheme 34).⁵⁶ They showed that the yield of nitriles depended on the type of silane mediating the dehydrations and that poly(methylhydroxysiloxane) (PMHS), triethyl and dimethoxymethyl silane were obtained in poor yields (Scheme 34).

The highest yield was observed with a bifunctional organosilane, 1,2-bis(dimethylsilyl)ethane, in the presence of 2.5 mol% of triruthenium catalyst. The reaction conditions were found to tolerate several functional groups including an alkene but not α,β -unsaturated amides. The authors also deduced the mechanism for the ruthenium catalyzed dehydration that the initially formed *N,N*-bis(trimethylsilyl)amide, at high temperatures, underwent *N*- to *O*-silyl migration followed by *syn*-elimination of silyl groups as its ether to give the corresponding nitrile and to regenerate the catalyst.

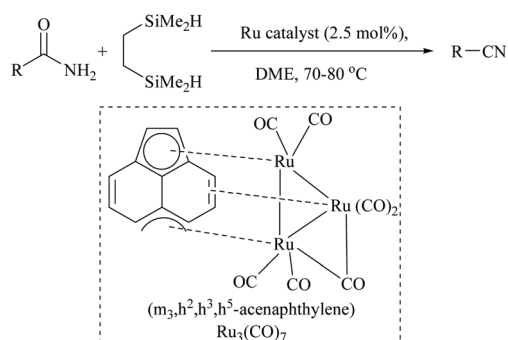
Silylative dehydrations of amides to nitriles catalyzed by iron hydrides have also been reported. For example, Beller and coworkers reported the first iron-catalyzed dehydration of primary amides to nitriles using silanes as the dehydrating agents (Scheme 35).⁵⁶ In the presence of silane, $(\text{EtO})_2\text{MeSiH}$, the dehydration did not proceed without a catalyst and also even with 5 mol% of $\text{Fe}(\text{CO})_5$, FeCl_2 or FeCl_3 . However, on the addition of various iron carbonyl clusters $[\text{CpFe}(\text{CO})_2]_2$,



Scheme 32 DFT calculations and Gibbs free energy ΔG (kcal mol⁻¹): aNHC catalyzed 1,2-elimination and β -elimination.



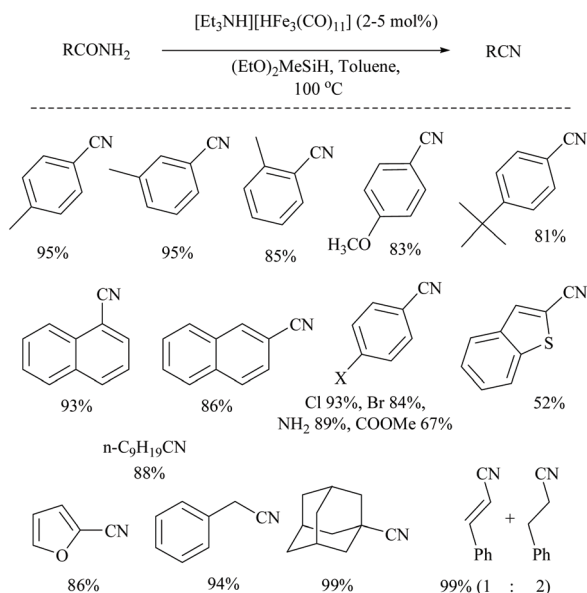
Scheme 33 DFT calculations: Cu-catalyzed β -elimination.



Scheme 34 Ruthenium catalyzed silylative dehydration.

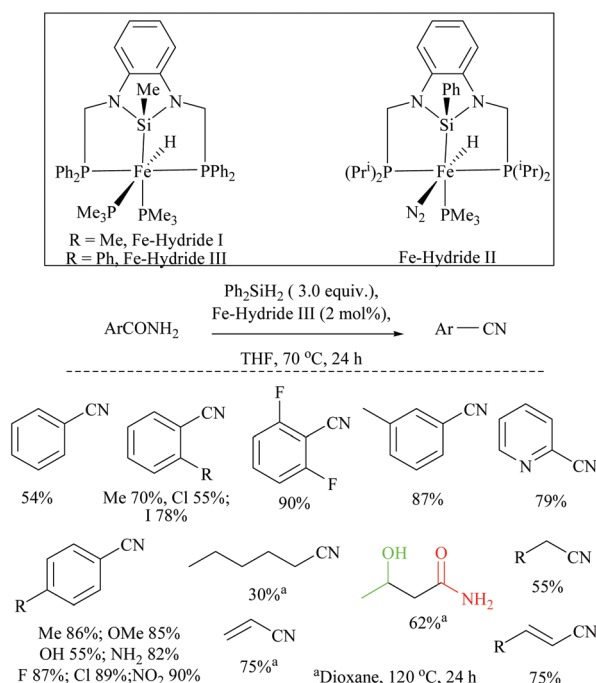
$\text{Fe}_3(\text{CO})_{12}$ and $[\text{Et}_3\text{NH}][\text{HFe}_3(\text{CO})_{11}]$, $\text{Fe}(\text{acac})_3$ and $\text{Fe}_2(\text{CO})_9$, a good conversion of amides to nitriles (83–97%) was observed. The optimization of the reaction for dehydration of benzamide resulted in conditions with the iron catalyst $[\text{Et}_3\text{NH}][\text{HFe}_3(\text{CO})_{11}]$ which gave a better result (90%) in a short time (3 h). Also the silane, $(\text{EtO})_2\text{MeSiH}$, along with toluene solvent showed better conversion in comparison with other silanes such as PMHS, PhSiH_3 , Ph_2SiH_2 , Et_3SiH , TMSOSiMeH , and $(\text{EtO})_3\text{SiH}$ as dehydrating agents. The result that PMHS gave poor yield was similar to that of Nagashima's method⁵⁶ reported earlier. Notably, the dehydration of cinnamamide

using four equivalents of silane in 1,4-dioxane as solvent gave the corresponding cinnamonitrile along with a reduced product 3-phenylpropionitrile in a 1 : 2 ratio. Also, no dehydration was observed with nicotinamide under these reaction conditions. They proposed a mechanism for the iron-catalyzed dehydration and found it to be similar to that reported earlier for the ruthenium catalyst (trirutheniumcarbonyl cluster, $(\text{ACE})\text{Ru}_3(\text{CO})_7$) by Nagashima and co-workers that involved metal catalysed *N*-silylation followed by its elimination as silyl ether.



Scheme 35 Iron-hydride catalyzed silylative dehydration.

Li and coworkers reported an air-stable N-heterocyclic PSiP pincer iron hydride and an analogous nitrogen iron hydride for the dehydration of primary amides to nitriles (Scheme 36).⁷³ They optimized the reaction conditions using three hydride catalysts and found that iron hydride III was better than other hydrides in terms of its air-stability and amide to nitrile conversion. Among different silanes, *viz.* (EtO)₂SiMeH, Ph₃SiH, Et₃SiH, Ph₂SiH₂ and (EtO)₃SiH that were tested, Ph₂SiH₂ was found to be a promising dehydrating

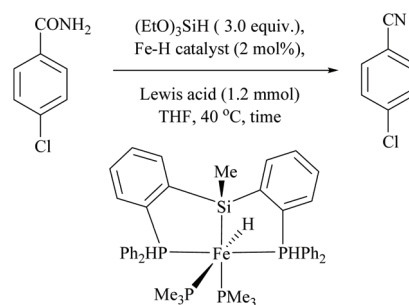


Scheme 36 Iron-hydride catalyzed silylative dehydration.

agent due to its excellent conversion and lower toxicity. The proposed mechanism for the conversion of amides to nitriles involved the iron-hydride complex assisted *O*-silylation followed by its elimination in ether form. They noted that the presence of an electron withdrawing group on the phenyl ring had raised the yields of the nitrile product. Moreover, the catalytic system did not affect the double bonds in acrylonitrile and cinnamamide. The dehydration of acetoacetamide, however, yielded 3-hydroxybutyramide, and the carbonyl group of the acetyl group was selectively reduced but not that of the amide group. The authors also reported the use of silyl iron hydride bearing [PSiP]-pincer ligands as catalysts along with a Lewis acid as the promoter for dehydration of primary amides to nitriles (Scheme 37).⁷³ Lewis acids accompanied by higher temperature, longer reaction time, and less yield have already been utilized for the dehydration of amides to nitriles.^{52,67,74,75}

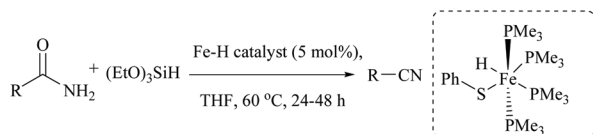
Thus, the authors wanted to study the influence of Lewis acids on the metal-hydride catalytic activity. The significance of a Lewis acid as an additive in the iron-hydride catalytic reaction was extensively studied and it was found that ZnBr₂ was the best under these catalytic conditions. However, the Lewis acid ZnBr₂ alone without the catalyst gave very poor yield. The same group has also achieved the reductive dehydration of amides to nitriles in good yields at lower temperature (60 °C) using hydrido thiophenolato iron(II) complexes [*cis*-Fe(H)(SAr)(PMe₃)₄] as catalysts with (EtO)₃SiH as the dehydrating agent (Scheme 38).⁵⁶ Though the reaction conditions with low catalyst loading (5 mol%) can tolerate aliphatic, acrylic and aromatic amides possessing nitro, halo and methoxy substituents, the keto-group of the amide was found to be reduced much faster than dehydration.

Similarly, the same group reported the applications of phosphorus-chalcogen chelated hydrido iron(II) complexes [(*o*-(R'₂P)-*p*-R-C₆H₄Y)FeH(PMe₃)₃, Y = O, S, Se] and also *cis*-sele-



Lewis Acid	Time (h)	Yield (%)
MgCl ₂	12	<10
FeCl ₃	12	<10
CuI	12	52
CuBr	12	<10
CuCl	12	<10
Cu(acac) ₂	12	<10
ZnCl ₂	12	77
ZnBr ₂	12	81
ZnBr ₂	24	85
ZnBr ₂	30	89
None	30	34

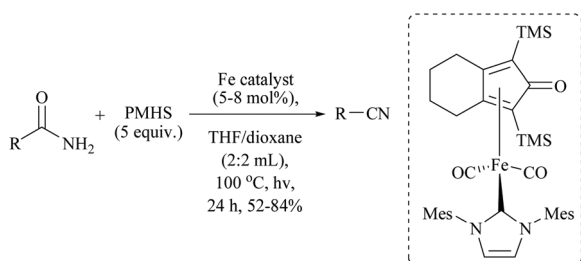
Scheme 37 Iron-hydride catalyzed silylative dehydration.



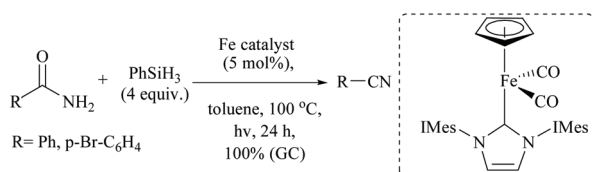
Scheme 38 Iron-hydride catalyzed silylative dehydration.

nophenolato iron hydrides (*cis*-[(ArSe)FeH(PMe₃)₄]) as catalysts and (EtO)₃SiH as a reducing agent for the dehydration of primary amides to nitriles.⁷⁶ In 2011, Lemaire and coworkers reported the use of the inexpensive polymethylhydrosiloxane (PMHS) as a reducing agent for the conversion of primary amides to the corresponding amines along with nitriles as side products.⁷⁷

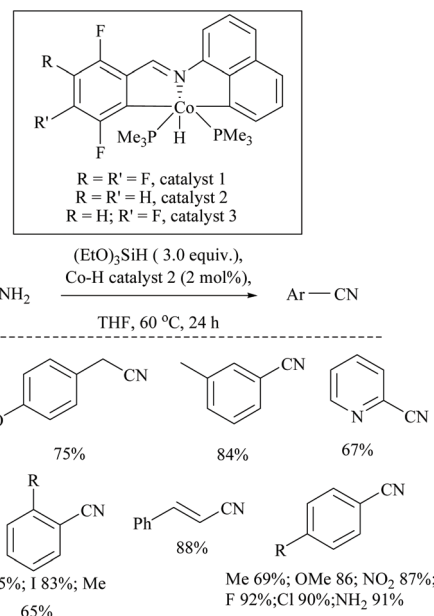
Iron based N-heterocyclic carbene complexes as homogeneous catalysts have been used in many transformations including silylative dehydrations.⁷⁸ Basic NHC with or without transition metal has also been used for silylative dehydrations.^{56,60,72} For example, when Darcel and coworkers⁵⁶ carried out the dehydration at high temperatures (100 °C) using a light-activated iron Knölker-type complex bearing an NHC ligand, nitriles were formed in moderate yields (Scheme 39). However, like Beller's observation (Scheme 35),⁵⁶ when they changed the catalytic system to [Et₃NH][HFe₃(CO)₁₁] and (EtO)₂MeSiH, cinnamamide gave cinnamyl nitrile (65% conversion, 61% isolated yield) under these conditions, along with reduced product 3-phenylpropionitrile in very small amounts (~2%). The group has also published the iron-catalyzed dehydration conditions but with different pairs of catalysts and silanes, *viz.* a cyclopentadienyl NHC-iron complex and phenylsilane (Scheme 40).⁵⁶ Primary amides were smoothly dehydrated into nitriles in quantitative yields; however, under this system, secondary and tertiary



Scheme 39 Iron Knölker-type complex catalyzed silylative dehydration.



Scheme 40 NHC-iron complex catalyzed silylative dehydration.

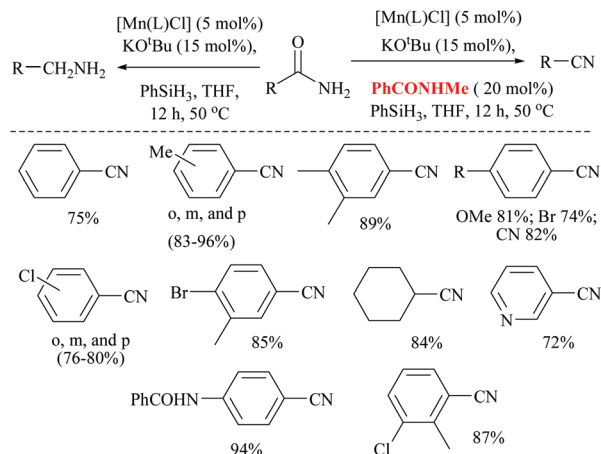


Scheme 41 Cobalt-hydride catalyzed silylative dehydration.

amides were found to be reduced to the secondary and tertiary amines, respectively.

In 2019, Li and coworkers, similar to their hydrido iron complexes along with (EtO)₃SiH as the reducing agent,^{56,79} reported for the first time, the hydrido cobalt(III) catalyst to dehydrate aryl and aliphatic primary amides to the corresponding nitriles in 67–92% yields using (EtO)₃SiH as the dehydrating agent at 60 °C (Scheme 41).⁶⁰ The optimization of the reaction resulted in findings that as the catalyst [CNC]-pincer hydrido cobalt(III) had decomposed at very high temperature, the conversion of the amide to nitrile was found to be reduced. They also found that THF was the best solvent system over toluene, DMF, DMSO, dioxane and acetonitrile. Compared to other silanes such as PhSiH₃, Ph₃SiH, Et₃SiH, TMDS, and PMHS, the highly flammable and toxic silane (EtO)₃SiH as a reducing agent was found to give good conversion and was selective over double bonds. The electron withdrawing groups afforded nitriles in good yields; however electron donating groups showed moderate results.

Mandal and coworkers reported the reduction of amides by a manganese catalyst with dual activity in transforming a primary amide into an amine or nitrile depending on the reaction conditions such as in the presence of a catalytic amount of a secondary amide (Scheme 42).⁸⁰ They optimized the reaction conditions with KO^tBu as a base, PhSiH₃ as a silylating agent, *N*-methyl-benzamide as an inhibitor, and a Mn catalyst for the conversion of various aryl, hetero-aryl and aliphatic primary amides to the corresponding nitriles in good yields (72–94%). Interestingly, the catalyst exhibited excellent selectivity and only the primary amide was selectively reduced in the presence of a secondary amide. In the substrate 4-cyanobenzamide, only the amide group was selectively reduced into the nitrile, sparing the existing nitrile. The mechanism that they

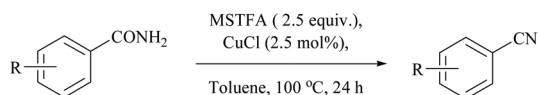


Scheme 42 Manganese with PhCONHMe for dehydration.

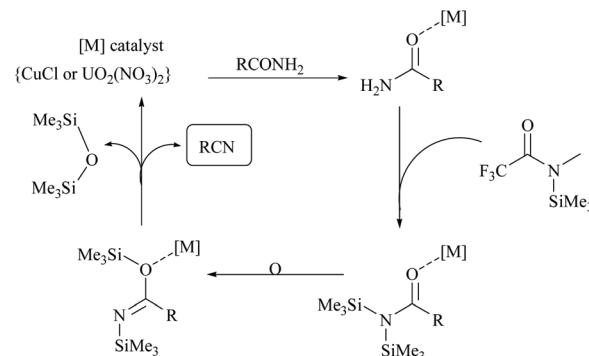
proposed was a non-radical, metal alkoxide transformation into a metal-hydride complex by σ -bond metathesis with the silane, followed by the formation of a primary amide coordinated complex with a vacant Mn(III) site. The activated amide can be converted into nitrile *via* *N*-silylation by *tert*-butoxide activated silane with the evolution of hydrogen gas followed by the elimination of the silyl group as its ether and regeneration of the catalyst. The same reaction conditions in the absence of PhCONHMe reduced the amide to the corresponding amine. Thus the single catalyst was either able to completely reduce the amide into amine or could give nitrile in the presence of the inhibitor PhCONHMe (Scheme 42).

Enthaler's research group has extensively worked on catalyzed silylative dehydrations of amides by elegantly overcoming the problems associated with the selection of silanes and catalysts.^{58,60} Instead of simple silanes, an excellent silylating reagent MSTFA, in the presence of various metal salts, iron(II) chloride, copper(I) chloride, zinc(II) triflate, and uranyl nitrate, was utilized for the dehydration of amides.⁵⁸ The addition of MSTFA meticulously ensured the safe handling of the reaction and elicited the scope of silylative dehydration by avoiding the release of hydrogen gas, but still followed through the 1,2-elimination of silyl groups. With copper(I) chloride as a metal catalyst in toluene, primary amides were dehydrated into their corresponding nitriles using MSTFA as a silylating reagent (Scheme 43).⁵⁸ The mechanism involved first the copper metal coordination with oxygen, and then silylation of nitrogen followed by *N*- to *O*-silyl migration yielded nitrile upon elimination of silylether as the by-product (Scheme 44).

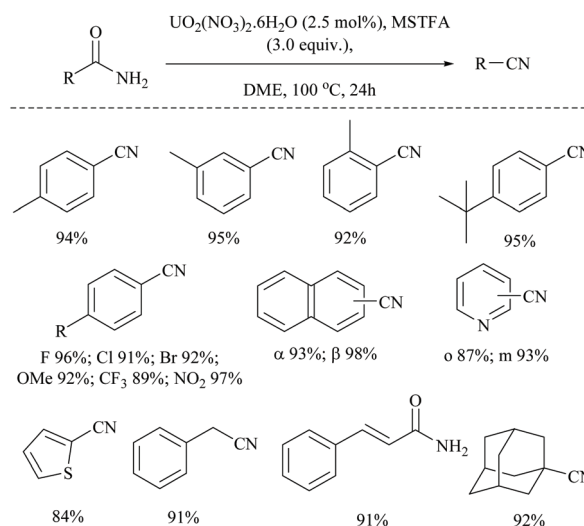
The same group has also reported uranium catalyzed (Scheme 45) and zinc catalyzed dehydrations of primary



Scheme 43 Copper catalyst for dehydration.



Scheme 44 Mechanism of Cu and U-catalyzed dehydrations.

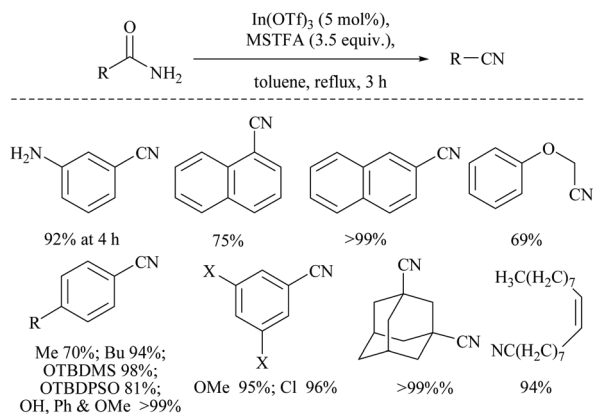


Scheme 45 Uranium catalyzed silylative dehydration.

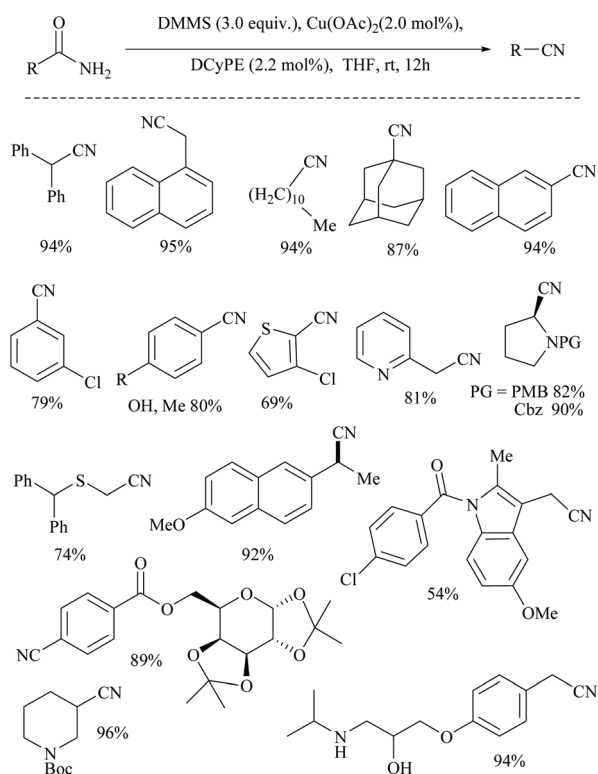
amides using MSTFA as the silylating reagent.⁵² They found that the catalyst alone did not give the product without the help of MSTFA; however on increasing its concentration, the yield of the nitriles increased. The temperature of the reaction conditions below 70 °C did not give appreciable yields of the product and at least 100 °C was required (Scheme 45). Also, MSTFA in THF as a solvent with *N*-heterocyclic compounds did not give appreciable yields. The silylating agent MSTFA and hence the mechanism were expected to work well with other metals too.

In 2014, Mineno and coworkers reported an efficient MSTFA dehydration method using indium(III) triflate (5 mol%) (Scheme 46).⁵⁹ The reaction with 3.5 equivalents of MSTFA in toluene under refluxing conditions was found to tolerate mono- and di-substituted aryl compounds, α,β -unsaturated amides and aliphatic amides. *O*-Silyl protected groups, *viz.* TBDMS and TBDPS groups, and also double bonds in oleamide were found to be resistant to the reaction conditions.

Buchwald and coworkers reported a copper hydride (CuH)-catalyzed silylative dehydration of primary amides to nitriles using a DCyPE ligand (Scheme 47).⁵⁸ Notably, in contrast to



Scheme 46 Indium catalyzed silylative dehydration.



Scheme 47 Copper catalyzed silylative dehydration.

other silylative dehydrations, under these conditions, the reaction was carried out at room temperature with 3.0 equivalents of dimethoxymethylsilane (DMMS) as a dehydrating agent. Dehydrations were performed on a variety of amide structures *viz.*, α -primary, secondary, and tertiary amides, aromatic amides with electron-withdrawing and -donating groups, and heterocyclic amides, including a thiophene, pyridine, pyrrolidine, and piperidine to give their corresponding nitriles in good yields. The optimization of the dehydration resulted in a copper(II) salt, a chelating diphosphine ligand (DCyPE), and monomeric DMMS. The addition of ligands was found to be

Table 2 Comparison of yields under various reaction conditions^a

Method				
Method 1	97%	85%	80%	93%
Method 2	98%	27%	38%	42%
Method 3	86%	48%	13%	68%
Method 4	<5%	<5%	24%	<5%
Method 5	85%	23%	75%	49%
Method 6	7%	<5%	16%	5%
Method 7	87%	8%	62%	47%

^a Method 1: Cu(OAc)₂, DCyPE, DMMS, temp, 12 h; method 2: POCl₃, Et₃N, rt, 12 h; method 3: propylphosphonic anhydride (T3P), toluene, 100 °C, 12 h; method 4: diiron nonacarbonyl, toluene, rt, 12 h; method 5: diiron nonacarbonyl, toluene, 100 °C, 12 h; method 6: phenylsilane, TBAF, rt, 30 min; method 7: phenylsilane, TBAF, 100 °C, 30 min.

necessary as in the absence of any ligand, the net dehydration reaction did not yield the nitrile product, even though it could catalyze the dehydrogenative silylation of acidic groups (Scheme 47). In addition, they chose four amide-substrates of different skeletons and subjected them to dehydration using six different classical and modern methods to obtain their corresponding nitriles using each of the methods described in Table 2. These six methods were followed as per the literature reported already and were compared with the one that the authors developed (copper-catalyst, DCyPE-ligand and DMMS – silylating agent). The yields of the corresponding nitrile products are shown in the following table. The results showed that the method developed by the authors performed better than other methods in terms of yield. In order to deduce a plausible mechanism, the authors have performed a computational study for the dehydration reaction of acetamide using DMMS as the silylating agent and copper(II) acetate as the catalyst (Schemes 30, 31, and 33). The DFT-optimized transition state structures followed the dehydration through β -elimination of monosilyl imidate in the presence of copper (II) acetate with a low energy barrier (+4.8 kcal mol⁻¹) using a DCyPE supporting ligand and resulted in the formation of the nitrile (Scheme 33). They also proved that the dehydration did not follow a high energy barrier (+35 kcal mol⁻¹) *syn*-elimination of *N,O*-disilyl imidate by carrying out the dehydration using the same reaction conditions but with a large excess (5 equiv.) of the silylating agent. The additional silanes under these reaction conditions resulted in only a trace amount of the nitrile product, which means that disilyl imidate did not lead to the formation of the nitrile product anyway.

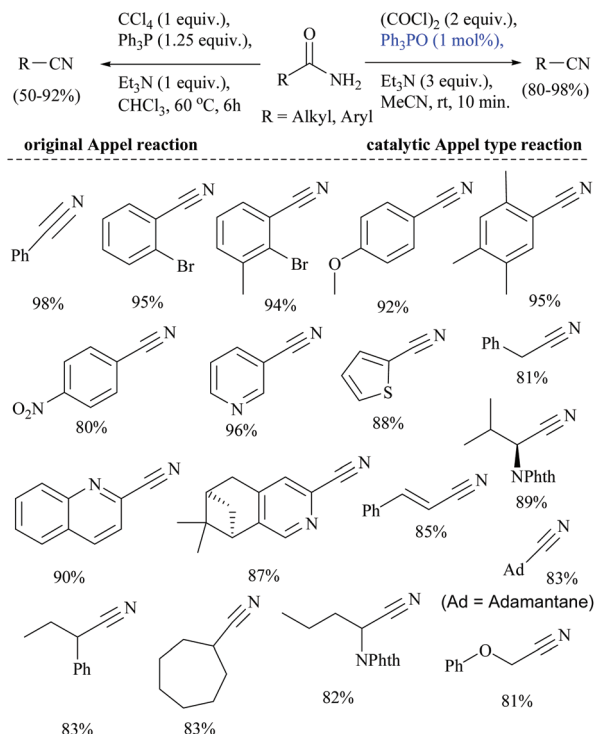
4. Non-transition metal catalyzed dehydration

Wang and coworkers searched for other catalytic dehydration methods without any additives such as a dehydrating agent and an activator, which led them to the use of silica supported potassium oxide acting as a heterogeneous catalyst.⁸¹ The K₂O/

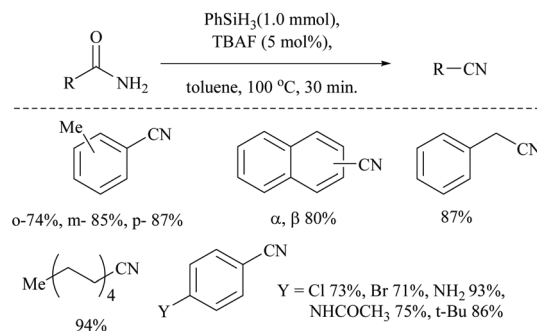
SiO₂ catalyst on calcination at 300 °C for 3 h afforded 2-CP in 23.6% yield.⁸¹ Huang and coworkers reported selective reduction of primary amides to nitriles with hydrosilane or hydrosiloxane using a catalytic system triethylborane in combination with an alkali metal base (Scheme 48).⁷⁴ They also studied the mechanism of BEt₃/base catalyzed amide hydrosilylation reaction and found that a modified Piers' silane Si-H...B activation mode facilitated the hydride abstraction by BEt₃ by the coordination of the Si-center with a hydroxide or an alkoxide. The reaction proceeded with the hydrosilylation reaction of primary amides using either PhSiH₃ at rt, or polymethylhydrosiloxane (PMHS) at 50 °C followed by reduction catalysed by BEt₃/base. The role of the base was also examined and it was found that the base enhanced the catalytic activity and facilitated the hydride transfer from Si to B through modified Piers' mode.

In 1971, the Appel reaction was applied for the synthesis of nitriles from amides in which the harmful CCl₄ and stoichiometric Ph₃P were utilized along with triethylamine.⁸² Shipilovskikh *et al.*, in 2018, developed a *catalytic* Appel-type dehydration of amides to nitriles by modifying the original Appel reaction in which the combination of Ph₃P, CCl₄, and Et₃N was replaced with a new system consisting of Ph₃PO, oxalyl chloride, and Et₃N (Scheme 49).⁸² The mechanisms of both the systems follow the same pathway through the initial formation of a common intermediate chlorophosphonium chloride.

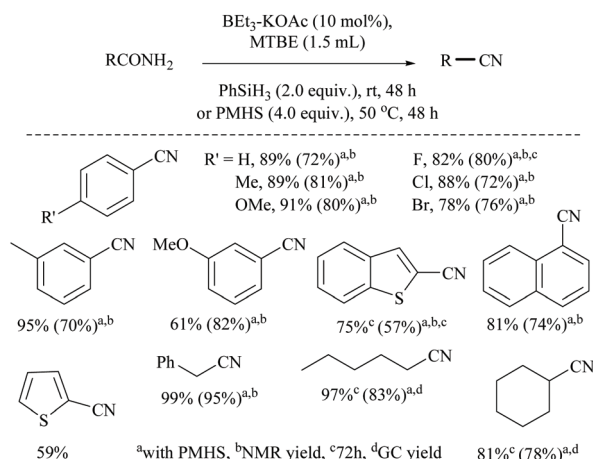
Optimization of the reaction conditions resulted in finding that the oxalyl chloride alone and the presence of Et₃N did not give the nitrile; however on the addition of Ph₃PO as a catalyst, the trio (oxalyl chloride, Ph₃PO and Et₃N) produced the nitrile product in the presence of MeCN solvent. Under these new catalyzed Appel type reaction conditions, aromatic, heteroaromatic, and aliphatic amides, including derivatives of α-hydroxy and α-amino acids, were converted to the corresponding nitriles. However, the exception was *N*-Boc protected amino acids which did not yield the nitrile, due to their reaction with oxalyl chloride. Beller *et al.* reported the fluoride-catalyzed de-



Scheme 49 Catalytic Appel type reaction.



Scheme 50 TBAF catalyzed silylative dehydration.



Scheme 48 BEt₃/base reagent for dehydration.

hydration of many aromatic and aliphatic amides to nitriles in good yields, using phenylsilane as the a dehydrating agent in the presence of a catalytic amount of TBAF (Scheme 50).⁵⁵

5. Organocatalytic dehydration of amides

Mandal and coworkers reported *N*-heterocyclic carbene (NHC) catalyzed silylative dehydration of primary amides (Scheme 51).⁷² They found that this NHC catalyzed reaction bypasses the energy need for the usual 1,2-siloxane (*syn*)-elimination step in metal/silane coupled dehydrations of amides to nitriles. The dehydration followed low energy barrier β-elimination to give nitriles. Optimization of the dehydration



Alternatively, without silylation, simply using an oxalyl chloride/DBU base combination in the presence of cyclopropanone as an organo-catalyst has also been found to be successful for the conversion of the amide group into its corresponding nitrile. Yadav and coworkers reported the conversion of aldoximes and primary amides of aromatic, heterocyclic, and aliphatic compounds into nitriles by employing cyclopropanone as an organo-catalyst (Scheme 52).⁸³ In the conversion of the amide to nitrile, oxalyl chloride along with the base DBU is used as an activating agent for the cyclopropanone to form the catalytically active species cyclopropenium cations (Scheme 52). Then, the cyclopropenium cation undergoes nucleophilic reaction with the amide to give the nitrile and regenerating the cyclopropanone catalyst. Under these reaction conditions, they found that the absence of either the cyclopropanone catalyst or the base DBU did not lead to the conversion even in the presence of oxalyl chloride.

Photocatalyzed dehydration of an amide is a sustainable approach as it uses a clean light source in the presence of an organo catalyst. In this context, Yadav and coworkers reported a photosensitization based approach for the dehydration of primary amides into the corresponding nitriles using eosin-Y



as an organophotoredox catalyst (Scheme 53).⁸⁴ The plausible mechanism proposed for this conversion involved light absorption by eosin-Y to reach the triplet excited state which transfers a single electron to CBr_4 to give a $\text{CBr}_3^\bullet$ radical. Upon reaction with DMF, $\text{CBr}_3^\bullet$ forms the radical species which in turn loses a single electron to the oxidatively quenched species EY^{+*} and forms the corresponding iminium ion completing the redox catalytic cycle. Then, the iminium ion loses COBr_2 to



form a Vilsmeier–Haack reagent which reacts with the added amide in its imidol form to give the corresponding nitrile. A similar method has also been applied for the dehydrosulfurization of thioamides.⁸⁵

7. Conclusions

Nitriles are important precursors or fine products in academia and in industries. Therefore, an efficient and green synthesis of nitriles has always been one of the major topics of interest for organic synthetic chemists. Dehydration of amides, an efficient, clean and fundamental route for the syntheses of nitriles, was started with stoichiometric amounts of metal as well non-metal based dehydrating reagents. With the progress in this field the stoichiometric reagents were replaced with catalysts to afford high conversion and selectivity towards nitriles. The more recent advancements require less than one equivalent of the metal catalyst, but the reactions require either very high reaction temperatures or mild reaction conditions using a stoichiometric dehydrating agent. Overwhelmingly, numerous research studies are available with respect to acetonitrile and silyl compounds as dehydrating agents either with or without a metal catalyst. It is clear from the review that attempts have been made to develop an efficient catalytic system for dehydration of amides.

In comparison with the metal catalyzed dehydrations, there are only a small number of publications available in the literature for non-transition metal and organo-catalyzed dehydrations. Moreover, a conclusion can be deduced based on the works by Buchwald and Mandal that the result outputs in terms of the reaction time and product yield of the organo-catalyzed silylative dehydrations are comparable with those of metal-catalyzed dehydrations. Hence, there is a better scope for the organo-catalyzed dehydration of amides. Indeed, the green and clean organo-photocatalyzed reaction has only a pair of research reports, each one on: the dehydration of carboxamides and the dehydrosulfurization of thioamides. Thus, in this direction, much more attention is needed in order to explore the sustainable dehydration methods.

Even though, in most of the cases, the dehydration of an amide gives a clean product, it can lead to multiple by-products depending on the other functionalities. Hence, in this context, new methods with the help of suitable reagents and catalysts for the selective dehydration of amides to nitriles are still needed to be explored. Moreover, most of the catalyzed dehydrations require inevitable use of non-renewable and expensive dehydrating agents. Additionally, these methods often require transition metals and/or stoichiometric reagents which might cause severe storage and handling problems as well as significant waste generation.

New strategies for promising methods, in order to address the problems associated with amide dehydration, *viz.* recyclability of reagents, selectivity, energy parameters and reaction mechanisms, are always in demand.

Conflicts of interest

There are no conflicts to declare.

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