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Synthesis and properties of seven- to nine-membered ring nitrogen heterocycles. Cyclic amidines and cyclic amidinium salts

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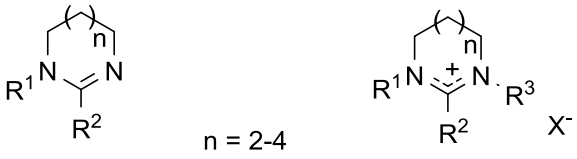
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Abstract

Medium-sized ring nitrogen heterocycles are an important class of compound which occurs in a range of natural and synthetic products. This review summarizes the current methods for the synthesis and chemical properties of cyclic amidines and cyclic amidinium salts of medium-sized rings (n= 2-4) with a fully saturated backbone.



Keywords: Medium-sized nitrogen ring heterocycles, cyclic amidines, cyclic amidinium salts

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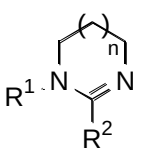
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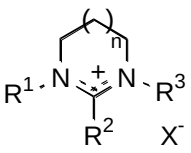
1. Introduction

Monocyclic medium-sized ring nitrogen heterocycles are an extremely important class of compounds which occur in a range of natural and synthetic products. The term “medium-sized ring” is usually applied to cyclic compounds having eight to eleven members;[1] however, seven- and twelve-membered rings are frequently included.

Cyclic amidines (**I**) represent a heterocyclic core of chemical, biological and pharmacological interest due to the nitrogen function present in these molecules (Figure 1). They are of considerable interest in drug discovery and have been proposed as potential agents for the treatment of several diseases. They are also important as synthetic intermediates.



I



II

I, n = 0, 1*H*-4,5-dihydroimidazole (imidazoline)
n = 1, 1,4,5,6-tetrahydropyrimidine
n = 2, 1*H*-4,5,6,7-tetrahydro-1,3-diazepine
n = 3, 1,4,5,6,7,8-hexahydro-1,3-diazocine
n = 4, 1*H*-4,5,6,7,8,9-hexahydro-1,3-diazonine
II, n = 0, 1*H*-4,5-dihydroimidazolium salt (imidazolinium)
n = 1, 1,4,5,6-tetrahydropyrimidinium salt
n = 2, 1*H*-4,5,6,7-tetrahydro-1,3-diazepinium salt
n = 3, 1,4,5,6,7,8-hexahydro-1,3-diazocinium salt
n = 4, 1*H*- 4,5,6,7,8,9-hexahydro-1,3-diazoninium salt

Figure 1

Five- and six-membered cyclic amidines, namely imidazolines and tetrahydropyrimidines (**I**, n = 0, 1), have been the most studied compounds and are present in many biologically active compounds.[2-14] However,

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higher homologues, such as 1*H*-4,5,6,7-tetrahydro-1,3-diazepines and 1,4,5,6,7,8-hexahydro-1,3-diazocines (**I**, n = 2, 3) have been less studied. It is known that medium-sized rings are generally more difficult to synthesize than their lower counterparts,[15-18] since the synthetic strategies employed have to overcome unfavorable transannular interactions leading to large enthalpies of activation[19,20] and the possibility of obtaining products of intramolecular condensation.[21]

Several tetrahydro-1,3-diazepines (**I**, n = 2) are especially interesting compounds due to their pharmacological activities. The antispasmodic,[22,23] hypoglycemic,[24,25] antiinflammatory,[25] diuretic,[24,25] and antitumor activities[26] of these compounds have been assessed. More recently, some of these compounds have been investigated as *N*-methyl-D-aspartate (NMDA) receptor antagonists,[27] dopamine D4 receptor[28] and muscarinic agonists,[29] as well as for the prophylaxis and protection of human skin against premature aging.[30]
Seven- and eight-membered substituted cyclic amidines are also useful precursors for the synthesis of heterocyclic ring systems as amidinium salts by alkylation,[31-33] and selectively substituted alkylenediamines by either reduction or alkaline hydrolysis.[34-36]

In recent years, cyclic amidinium salts (CAS) with a fully saturated backbone (**II**) have attracted a great deal of attention. 1*H*-Dihydroimidazolium and tetrahydropyrimidinium salts (**II**, n = 0, 1) have been the most investigated CASs. These salts have been studied in the past century as models of the coenzyme N^5,N^{10}-methenyltetrahydrofolic acid, which is involved in the biochemical transfer of one carbon unit at the oxidation level of formic acid.[37-46] CASs **II** with different patterns of substitution have been employed as synthetic intermediates for the preparation of cyclic and acyclic compounds carrying the alkylenediamine unit (>N-$(CH_2)_n$-N<).[42,43] Imidazolinium and tetrahydropyrimidinium salts are chemical precursors of N-heterocyclic carbenes (NHCs) that, either alone or as a metal complex, are efficient catalysts for chemical transformation reactions.[47-49] 4,5-Dihydro-1*H*-imidazolium salts have also been investigated as surfactants,[50] due to their potential for chiral molecular recognition,[51] and as catalysts for several chemical reactions.[52,53]

Unlike the compounds mentioned above, medium-sized cyclic amidinium salts have been less explored. The most important application of these salts is based on their capacity to act as precursors of expanded ring NHCs, which are stronger σ-donating ligands.

The present review will therefore focus on the synthesis and the main properties of medium-sized cyclic amidines and cyclic amidinium salts (**I** and **II**, n= 2-4).

2. Synthesis of Cyclic Amidines

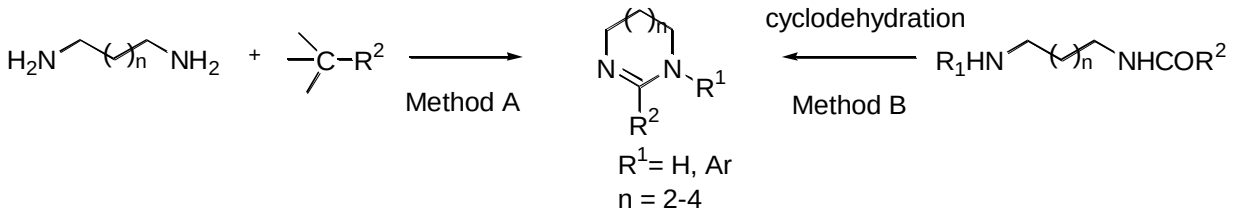
Synthetic methods of medium-sized cyclic amidines are generally extensions of the methods employed for lower homologous cycles such as imidazolines and tetrahydropyrimidines. Literature data on the higher homologues, 1,3-diazepines, diazocines and diazonines, are scarce.

There are basically two methods for the synthesis of cyclic medium-sized amidines which involve acyclic compounds as precursors (Scheme 1) :

Method A: from 1,*n*-alkylenediamines as precursors

Method B: from ω-aminoamides as precursors

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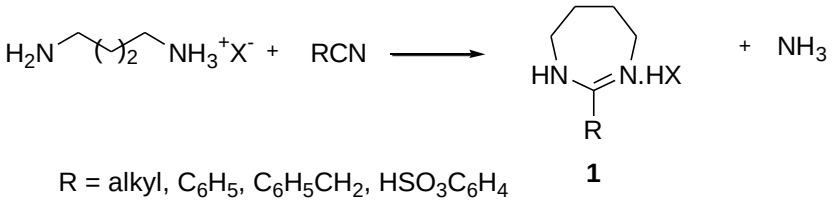


Scheme 1

2.1 Synthesis of amidines from 1,*n*-alkylenediamines (Method A)

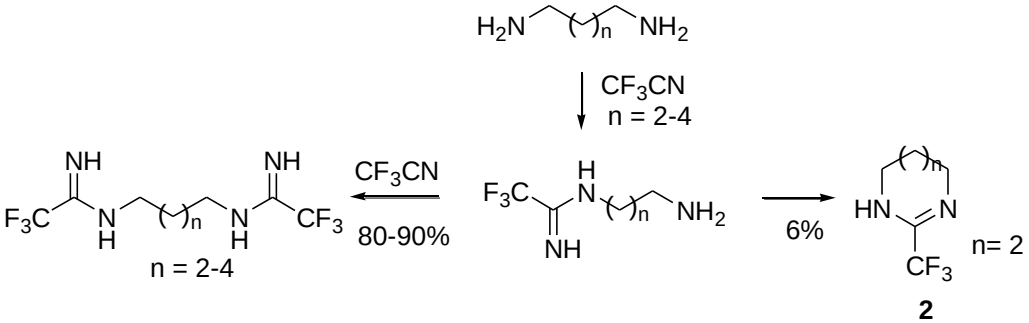
The construction of the amidine nucleus involves the coupling of a 1,*n*-diamine with an appropriate condensation partner that provides the C-2 of the amidine ring. The synthesis of tetrahydro-1,3-diazepines and hexahydro-1,3-diazocines from tetramethylenediamines (putrescine) and pentamethylenedimine (cadaverine) was the first class of methods developed in chemistry. Generally, this method was applied to the synthesis of cyclic amidines C-2-substituted with alkyl or aryl groups but without N-substitution.

Cyanides can be an adequate source of C-2. Oxley *et al.* have described the synthesis of some 2-substituted tetrahydrodiazepines **1** in good yields from a mixture of a nitrile and tetramethylenediamine/ tetramethylenediammonium bistoluene-*p*-sulfonate at 200 ^{o}C (Scheme 2).[54] The practical limitation of this method appears to be that ocurring with the synthesis of tetrahydrodiazepines, since attempts to produce hexahydrodiazocines and diazonines (**I**, n= 3,4) by condensation of penta- and hexamethylenediamines or their salts with cyanides, resulted in a mixture from which no homogeneous solid derivative could be isolated.



Scheme 2

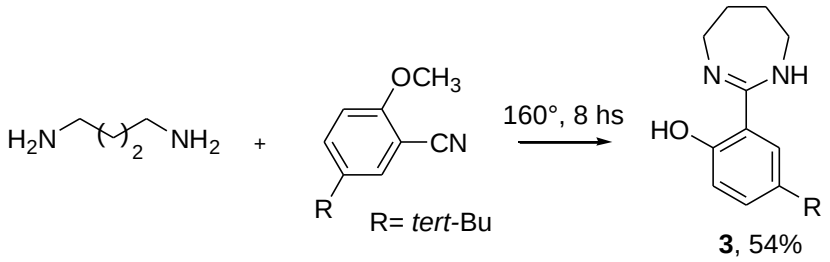
Johnson and Woodburn have used more electrophilic nitriles for the synthesis of cyclic amidines of five to seven members. Thus, the authors accomplished the reaction between trifluoroacetonitrile with aliphatic diamines to yield carboxamidines and, where n = 0-2, with cyclic amidines as well.[55] The reactions with tetramethylenediamine yielded carboxamidine as the major product but only 6 % of the cyclic compound, namely 2-trifluoromethyl-4,5,6,7-tetrahydro-1*H*-1,3-diazepine **2**, was isolated (Scheme 3). On the other hand, penta- and hexa-methylenediamine only produced carboxamidines when treated with trifluoroacetonitrile.



Scheme 3

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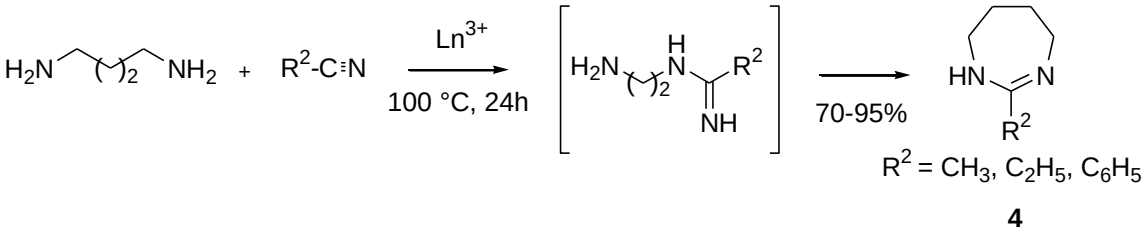
Working at high temperatures, 2-*o*-hydroxyphenyl-1,3-diazepines **3** with antihypertensive activity have been obtained from putrescine and 2-methoxybenzonitriles (Scheme 4).[56]



Scheme 4

The use of catalysts has, in some cases, improved the results of reactions of diamines with nitriles. In 1974, three 1,3-diazepines containing α-alkoxybenzyl groups on C-2, with potential hypoglycemic and natriuretic activity, were synthesized using 2-alkoxy-2-arylacetonitriles as source of C-2.[57] The conversion of the nitrile into the corresponding amidine was readily accomplished by heating the reaction mixture with an excess of diamine using a few drops of CS_2 as catalyst.

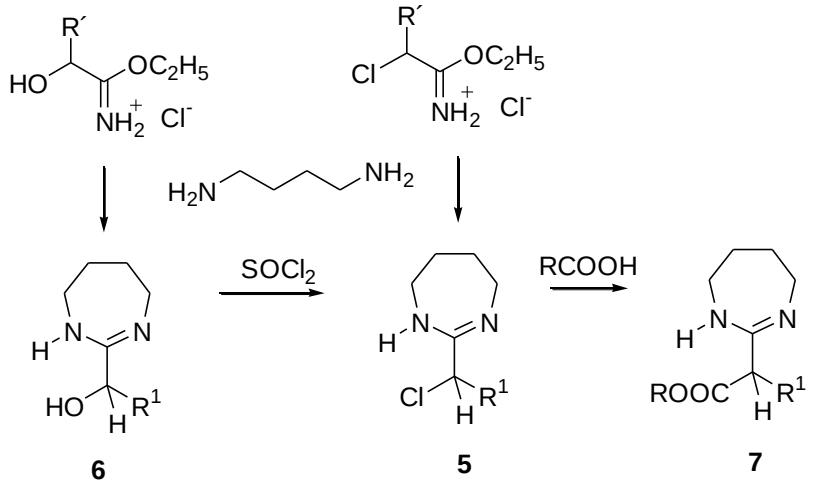
Forsberg *et al.* have published the lanthanide(III)-catalyzed addition of amines to nitriles for the construction of amidines.[58] Ln^{3+} ions activate nitriles through a predominantly electrostatic ion-dipole interaction. This interaction enhances polarization of the cyano group, thereby facilitating the attack by the nucleophilic amine. The reactions are quite facile and progress to completness (yields 75-95%) when equimolar amounts of amine and nitrile are heated with 1 mol % Ln^{3+} for 24 h. By means of this strategy, 2-methyl, ethyl and phenyl 1*H*-4,5,6,7-tetrahydro-1,3-diazepines **4** could be synthesized (Scheme 5).



Scheme 5

As activated equivalents of carboxylic acids, **imidic esters** (imidates), have been used in the synthesis of cyclic amidines. The reaction of an imidic ester with alkylendiamines generally requires milder reaction conditions than those in which the corresponding nitriles are employed. Sahyun *et al.* have obtained among other cyclic amidines 2-chloro- and 2-hydroxy-alkyltetrahydro-1,3-diazepines **5,6** from the corresponding imidic ester hydrochlorides. The compounds were subsequently transformed into esters **7** with antispasmodic activity (Scheme 6).[22]

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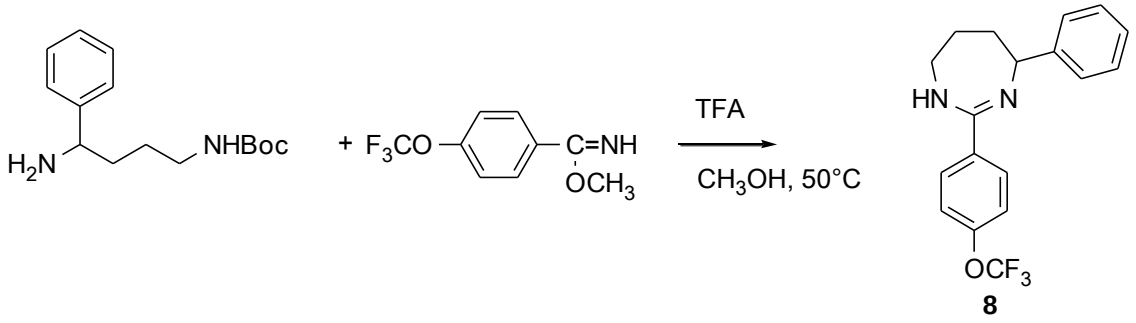
Scheme 6

White *et al.* have patented the synthesis of a series of five to eight-membered amidines (imidazolines, tetrahydropyrimidines, tetrahydro-1,3-diazepines and hexahydro-1,3-diazocines) from imidic esters with the corresponding diamines.[59,60]

Many of the synthesized compounds have pharmacological activity *e.g*. diuretic, anti-inflammatory, hypoglycemic and cardiovascular activity.[59] Miller *et al.* have obtained other tetrahydrodiazepines and hexahydrodiazocines with antifungal activity using the same methodology.[61]

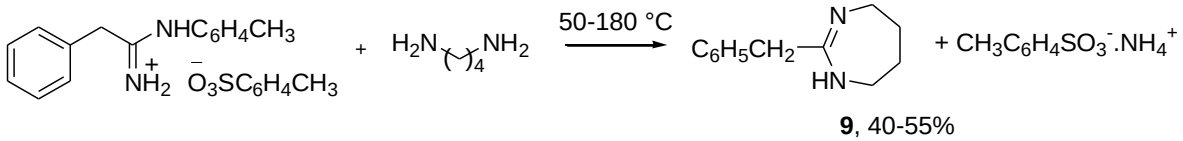
Another example of the use of an imidic ester as source of C-2 is the formation of the 2-phenyl-1,3-diazepines by condensation of 1,4-diaminobutane derivative with methyl benzimidate under moderate reaction conditions. These compounds have been studied as potential dopamine D4 receptor agonists.[28]

More recently, a series of cyclic amidines including a 2,4-diaryl-1,3-diazepines **8** with selective NMDA antagonist activity has been synthesized by reaction of an imidoester with 2-phenylputrescine (Scheme 7).[27]



Scheme 7

The synthesis of cyclic amidines from alkylenediamines and **amidinium** salts was reported in 1950 by Oxley *et al.*,[62] who obtained 2-benzyl-1*H*-4,5,6,7-tetrahydro-1,3-diazepine **9** from N-substituted amidinium salts and putrescine (Scheme 8).

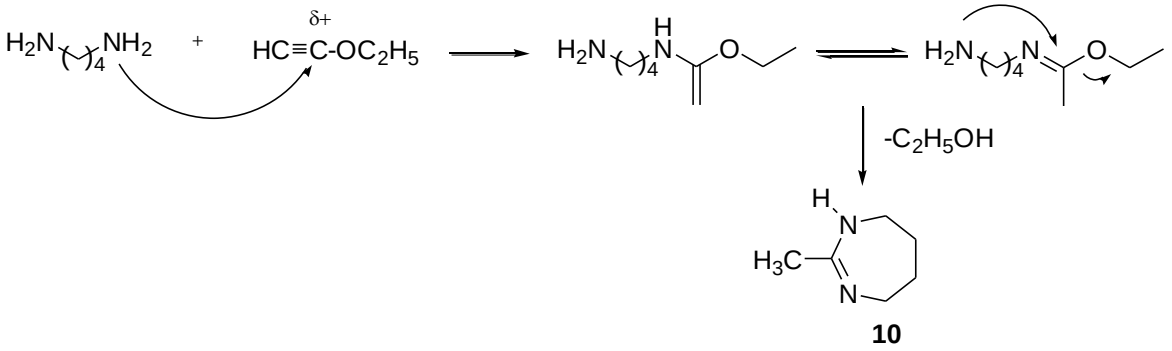


Scheme 8

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Desmarchelier reported the synthesis of 2-methyl-1*H*-4,5,6,7-tetrahydro-1,3-diazepine from 1,4-diaminobutane with acetamidine hydrochloride under mild experimental conditions.[63]

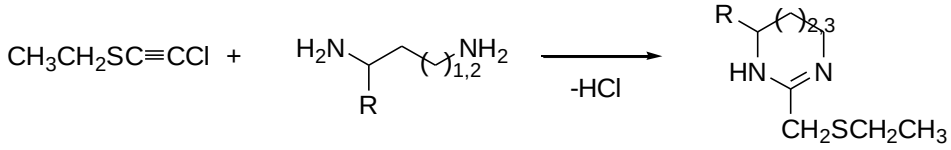
Ethoxyacetylene was used as electrophilic precursor to synthesize 2-methyl-1*H*-4,5,6,7-tetrahydro-1,3-diazepine **10** (Scheme 9).[64]



10

Scheme 9

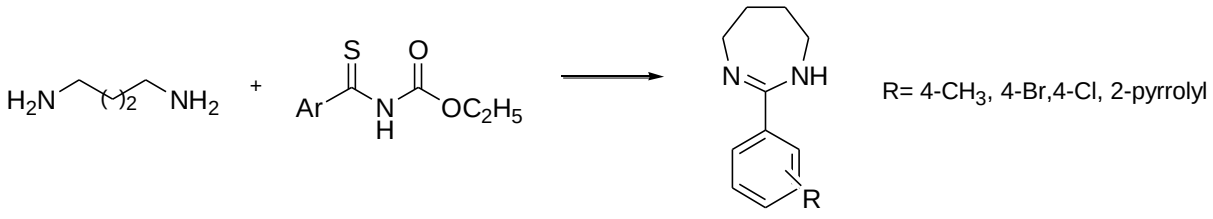
Organylthiochloroacetylenes react easily with aliphatic diamines such as putrescine and cadaverine, at low temperatures in benzene, with the formation of 2-(ethylthiomethyl)-1,3-diazepine **11** and the corresponding hexahydrodiazocine (Scheme 10).[65]



11

Scheme 10

Thioamides such as N-ethoxycarbonylthioamides have been used to obtain 2-arylderivatives of 4,5,6,7-tetrahydro-1*H*-1,3-diazepines **12** to obtain moderate yields (Scheme 11).[66]

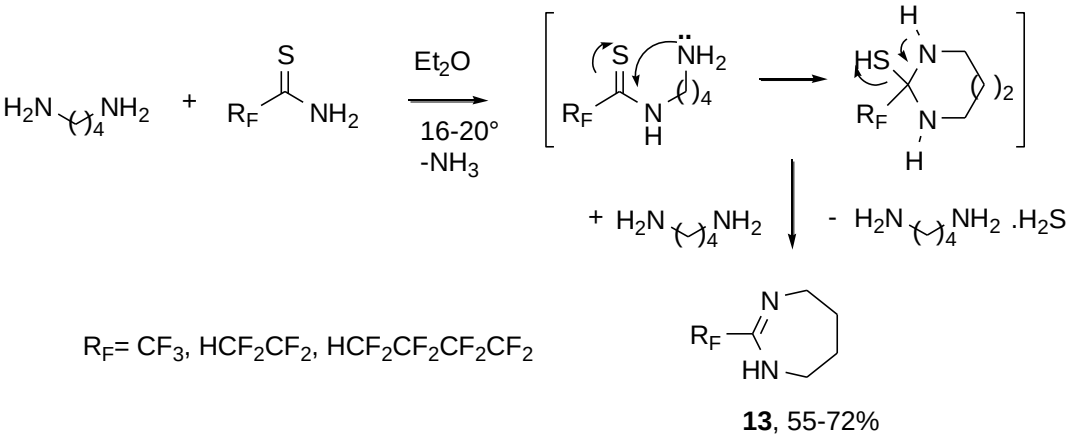


12, 25-57%

Scheme 11

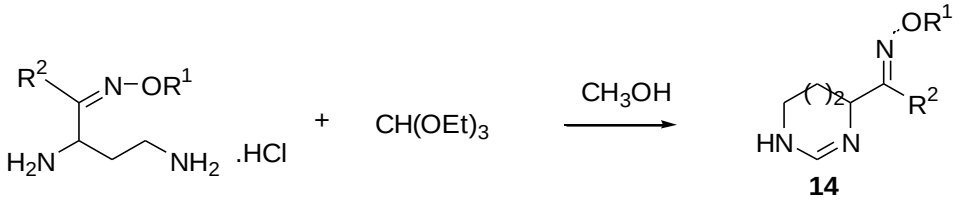
One of most intensively developing fields in the chemistry of biologically active heterocycles is the synthesis of fluorinated analogues. 4,5,6,7-Tetrahydro-1*H*-1,3-diazepines **13** bearing a perfluoroalkyl group on C-2 have been obtained by reactions of tetramethylenediamine and a polyfluoroalkylthioamide.[67] One possible reaction mechanism assumes the intramolecular cyclization of the re-amidation product with the subsequent separation of hydrogen sulfide from the product of cyclization. The reactions have been performed under mild conditions to obtain good yields of the corresponding heterocycles (Scheme 12).

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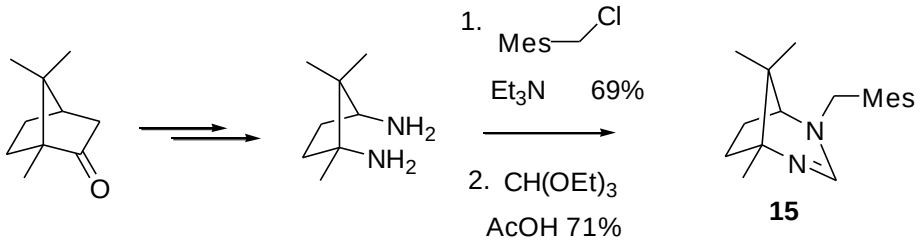
Scheme 12

Orthoesters have also been used in the synthesis of cyclic amidines. Plate *et al.* have synthesized a series of 1,3-diazacycloalkyl carboxaldehyde oxime derivatives with potential muscarinic activities.[29] Among them, 4,5,6,7-tetrahydro-1,3-diazepine-4-carboxaldehyde oxime **14** was obtained using triethyl orthoformate as reagent (Scheme 13).



Scheme 13

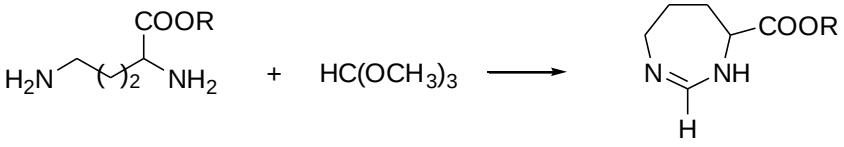
In 2009, Wilhelm *et al.* synthesized a tetrahydro-1,3-diazepine **15** containing a bicyclic core derived from camphor from 1,3-diamino-1,2,2-trimethylcyclopentane, which is a compound easily obtained from camphor that is a very useful material to construct chiral compounds. The method involves the N-alkylation of the camphoric diamine followed by reaction with triethyl orthoformate.[68]



Scheme 14

Lentzen has patented the synthesis of five- to seven-membered 2-substituted cyclic amidines **16**. The ring closure to the generate the cyclic amidines was accomplished by reactions of C-substituted alkylenediamines with orthoesters (Scheme 15).[69]

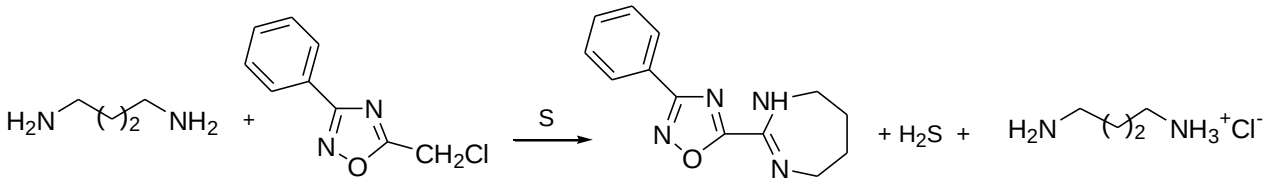
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16

Scheme 15

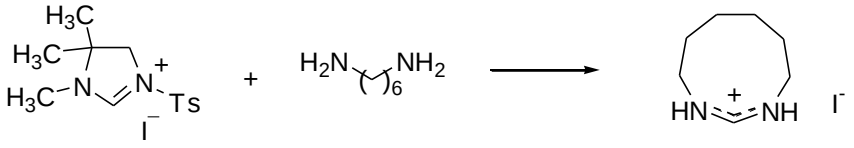
Alkyl halides have also been used as source of C-2. The 3-substituted 5-(chloromethyl)-1,2,4-oxadiazole reacts with putrescine in the presence of sulfur as dehydrogenating agent to yield 2-heteroaryltetrahydro-1,3-diazepines **17** (Scheme 16).[70,71]



17

Scheme 16

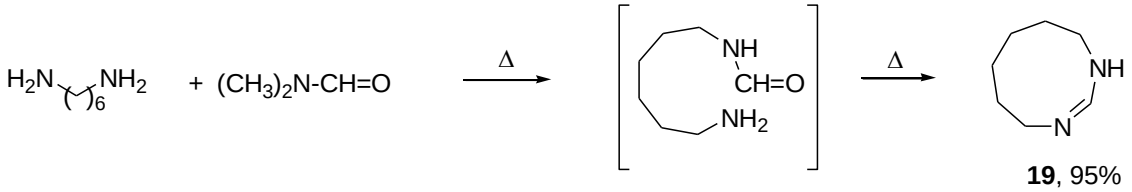
Bieraugel has studied the carbon unit transfer from cyclic **amidinium salts** to bifunctional nucleophiles as α,ω-diaminoalkanes.[72,34] Thus, protonated 4,5,6,7,8,9-hexahydro-1,3-diazonine **18** was obtained using an imidazolinium salt as C-2 donor to hexamethylenediamine (Scheme 17). The process results from the ability of such imidazolinium salt to transfer a formyl equivalent to a variety of nucleophiles. However, the product was only characterized by ^{1}H-NMR through the presence of the N = CH signal at 8.00 ppm.



18

Scheme 17

Recently Simion *et al.* have reported an unexpected synthesis of the same diazonine **19** using hexamethylenediamine and **dimethylformamide** as source of C-2.[73] The cyclization was explained as a two-step process involving formylation and subsequent intramolecular condensation (Scheme 18).

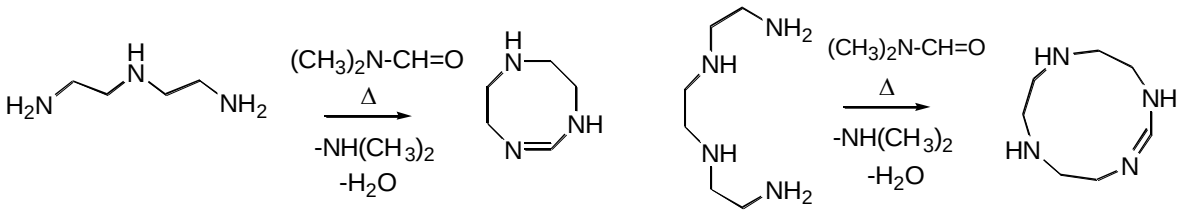


19, 95%

Scheme 18

The usefulness of this process has been demonstrated through the synthesis of two other nitrogen-containing macroheterocycles (Scheme 19).

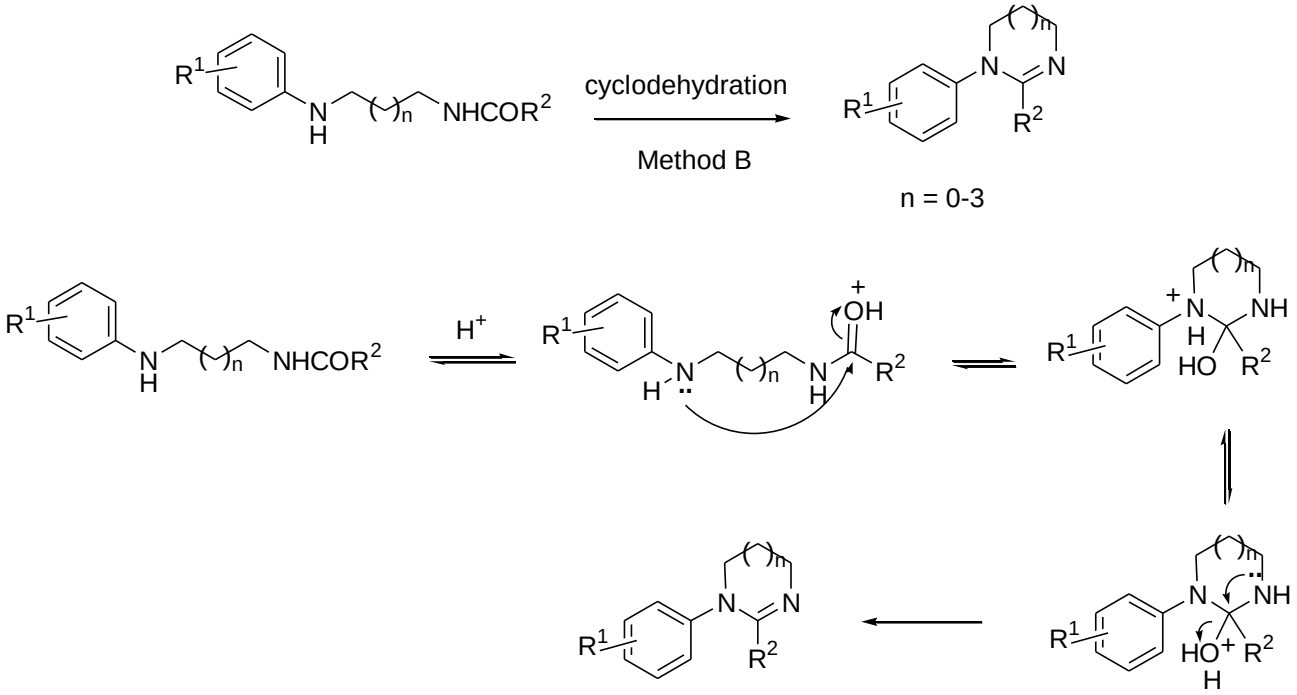
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Scheme 19

2.2 Synthesis of amidines from ω-aminoamides (Method B)

This general method involves the cyclodehydration of *N*-aryl-*N*′-acylalkylenediamines **20** to the corresponding cyclic amidine by heating with a cyclizing agent. The proposed mechanism is given below. This is a general method for the synthesis of five to eight-membered N-aryl substituted cyclic amidines, and the variations depend on the synthetic route to generate the precursor aminoamide and the cyclizing agent used (Scheme 20).

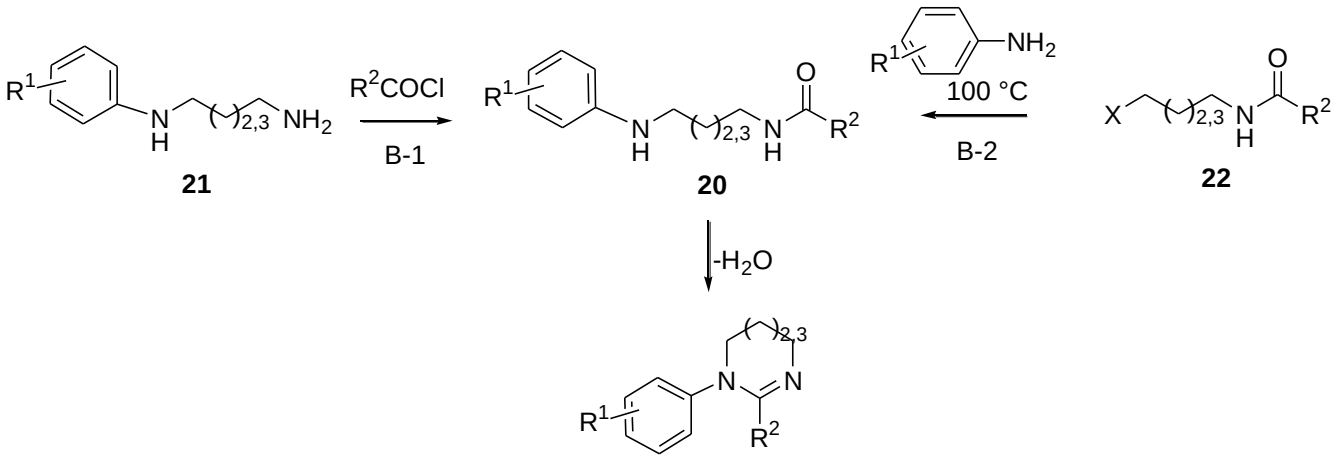


Scheme 20

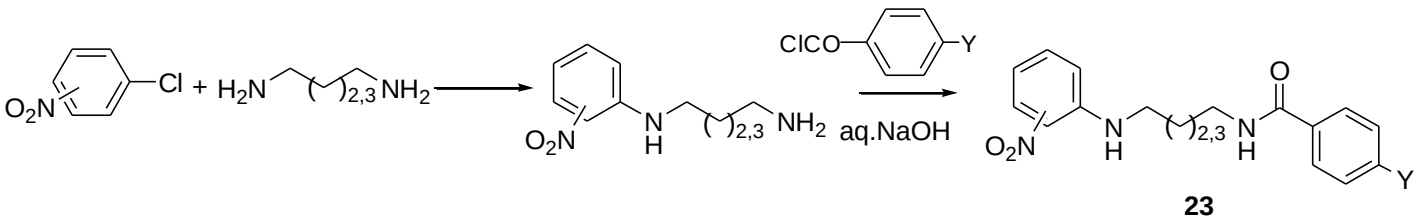
Two methods have been employed to synthesize the precursor aminoamides **20** of tetrahydrodiazepines and hexahydrodiazocines: N-acylation of the corresponding N-arylalkylenediamines **21** (Method B-1) and from the reaction of *N*-4-halobutyl or 5-halopentyl benzamides **22** with arylamines (Method B-2) (Scheme 21).

In the first method (Method B-1), the precursor ω-aminoamides can be obtained by aminolysis of the appropriate chloronitrobenzene with tetra- or penta-methylenediamine, followed by benzoylation under Schotten-Baumann conditions. Thus, in 1977 Perillo *et al.* reported the use of *N*-nitroaryl-*N'*-aroyltetra- and penta-methylenediamines **23** as precursors of 1,2-diaryl-1,3-diazepines and diazocines having a nitrophenyl substituent on N-1 (Scheme 22).[74]

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Scheme 21



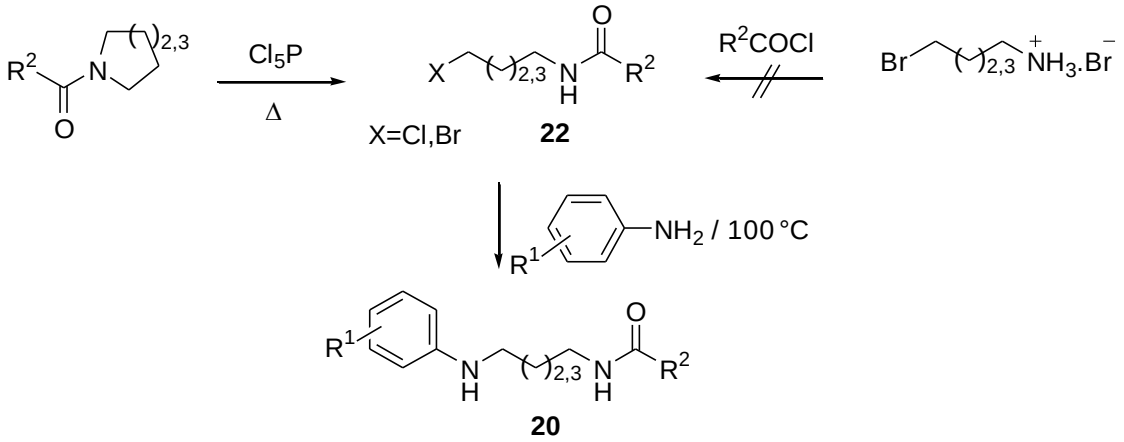
Scheme 22

However, when the aryl group of the *N*-arylalkylenediamine is not substituted or when it is substituted with electron donor or slightly electron withdrawing groups, the reaction with acyl chlorides under Schotten-Baumann conditions led to the corresponding *N,N'*-diacyl derivatives.[32] Selective monoacylation was achieved using aliphatic carboxylic acid anhydrides working at 0 ^{o}C in a biphasic system (Cl_3CH/aqueous Na_2CO_3)[75] or working in homogeneous phase with DCM as solvent and TEA as acceptor of hydrogen chloride at -10 ^{o}C (55-60%).[32]

Another suitable synthetic strategy to obtain the precursor aminoamides **20** involves the synthesis of *N*-4-halobutyl or *N*-5-halopentylbenzamides **22** as key synthetic intermediates and subsequent reaction with amines (Method B-2)**.** Attempts to obtain the haloalkylamides **22** by acylation of the corresponding ω-haloalkylamines in basic medium have failed, because in such reaction medium intramolecular aminolysis occured.[33] Conversely, 4-chlorobutyl- and 5-chloropentyl-benzamides could be obtained by reaction of *N*-benzoylpyrrolidine or piperidine with phosphorus pentachloride by the von Braun reaction (Scheme 23).[76] The subsequent reaction with arylamines leads to the expected aminoamides **20** (R^1 = Ar).[31,77] A drawback of this strategy is that the procedure is restricted to N-acyl derivatives without α-hydrogens.[76]

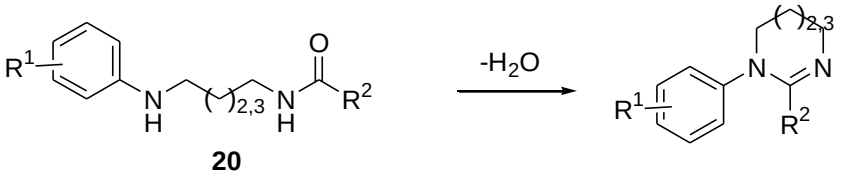
Phosphorus oxychloride, polyphosphoric acid (PPA), ethyl polyphosphate (PPE) and trimethylsilyl-polyphosphate (PPSE) have been used as dehydrating agents for the synthesis of N-aryl five- to eight-membered cyclic amidines.[35,78-80] The synthesis of medium-sized cyclic amidines through the cyclization of aminoamides was first reported in 1977 (Table 1, entries 1-6),[74] when a series of 1,2-diaryl-1,3-diazepines and diazocines having a nitrophenyl substituent on N-1 were synthesized through the ring closure of the corresponding *N*-nitroaryl-*N*'-aroyltetra- and penta-methylenediamines, respectively, employing a chloroform solution of PPE or phosphorus oxychloride as cyclizing agent for the synthesis of diazepines and neat PPE to obtain the corresponding diazocines. Similarly in 2000 Hedrera *et al.* synthesized 1,3-diazepines employing the same cyclizing agent (Table 1, entries 7-10).[33,35]

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Scheme 23

Table 1. Synthesis of ω-aminoamides and their cyclization conditions



Entry	**20** Synthetic method	R^1	R^2	n	Cyclization conditions	Yield (%)	Ref.
1	B-1	4-NO_2	C_6H_5	2	PPE/$CHCl_3$, $POCl_3$	92, 76	74
2	B-1	4-NO_2	4-NO_2-C_6H_4	2	PPE/$CHCl_3$, $POCl_3$	89, 65	74
3	B-1	2-NO_2	C_6H_5	2	PPE/$CHCl_3$, $POCl_3$	80, 62	74
4	B-1	2-NO_2	4-$NO_2C_6H_4$	2	PPE/$CHCl_3$, $POCl_3$	75, 59	74
5	B-1	4-NO_2	C_6H_5	3	PPE, $POCl_3$/SF[a]	39, 24	74
6	B-1	4-NO_2	4-$NO_2C_6H_4$	3	PPE, $POCl_3$/SF	42, 30	74
7	B-2	H	C_6H_5	2	PPE/$CHCl_3$	64	33,35
8	B-2	4-CH_3	C_6H_5	2	PPE/$CHCl_3$	68	33,35
9	B-2	4-OCH_3	C_6H_5	2	PPE/$CHCl_3$	60	33,35
10	B-2	4-Cl	C_6H_5	2	PPE/$CHCl_3$	55	33,35
11	B-1	2-NO_2	C_2H_5	2	PPE/$CHCl_3$/MW[b]	93	81
12	B-1	4-Br	C_6H_5	2	PPE/$CHCl_3$/MW	89	81
13	B-1	3,4-$(CH_2)_4$	$C(CH_3)_3$	2	PPE/$CHCl_3$/MW	84	81
14	B-1	H	C_6H_5	3	PPE/$CHCl_3$/MW	83	81
15	B-1	4-CH_3	C_6H_5	3	PPE/$CHCl_3$/MW	79	81
16	B-1	4-Cl	C_2H_5	2	PPE/$CHCl_3$/MW/100°	79	75
17	B-1	4-Cl	CH_3	2	PPE/$CHCl_3$/MW/100°	85	75
18	B-1	4-Cl	*iso*-C_3H_7	2	PPE/$CHCl_3$/MW/100°	75	75
19	B-1	4-CH_3	CH_3	2	PPE/$CHCl_3$/MW/100°	87	75
20	B-1	4-CH_3	C_2H_5	2	PPE/$CHCl_3$/MW/100°	71	75
21	B-1	4-CH_3	*iso*-C_3H_7	2	PPE/$CHCl_3$/MW/100°	90	75

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Table 1. Continued

Entry	20 Synthetic method	R^1	R^2	n	Cyclization conditions	Yield (%)	Ref.
22	B-1	4-Br	iso-C_3H_7	2	PPE/$CHCl_3$/MW/100°	72	75
23	B-1	4-F	iso-C_3H_7	2	PPE/$CHCl_3$/MW/100°	63	75
24	B-1	4-CH_3	iso-C_3H_7	3	PPE/$CHCl_3$/MW/100°	42	75
25	B-1	4-CH_3	C_2H_5	3	PPSE/SF[b]/ MW/90°	58	82
26	B-1	4-CH_3	iso-C_3H_7	3	PPSE/SF/ MW/90°	60	82
27	B-1	4-Cl	CH_3	3	PPSE/SF/ MW/90°	50	82
28	B-1	4-Cl	C_2H_5	3	PPSE/SF/ MW/90°	61	82
29	B-1	4-Cl	iso-C_3H_7	3	PPSE/SF/ MW/90°	84	82
30	B-1	4-Br	CH_3	3	PPSE/SF/ MW/90°	52	82
31	B-1	4-Br	C_2H_5	3	PPSE/SF/ MW/90°	61	82
32	B-1	4-Br	iso-C_3H_7	3	PPSE/SF/ MW/90°	81	82
33	B-1	H	H	3	PPSE/SF/ MW/90°	27	82
34	B-1, B-2	H	C_6H_5	3	PPE/$CHCl_3$/MW/120°	95-100	32
35	B-1, B-2	H	4-ClC_6H_4	3	PPE/$CHCl_3$/MW/120°	95-100	32
36	B-1, B-2	4-CH_3	C_6H_5	3	PPE/$CHCl_3$/MW/120°	95-100	32
37	B-1, B-2	4-CH_3	4-ClC_6H_4	3	PPE/$CHCl_3$/MW/120°	95-100	32
38	B-1, B-2	4-Cl	C_6H_5	3	PPE/$CHCl_3$/MW/120°	95-100	32
39	B-1, B-2	3,4-Cl_2	4-ClC_6H_4	3	PPE/$CHCl_3$/MW/120°	95-100	32
40	B-2	4-Cl	C_6H_5	2	PPE/$CHCl_3$/MW/70°	96	31
41	B-2	H	4-ClC_6H_4	2	PPE/$CHCl_3$/MW/70°	95	31
42	B-2	2-Cl	C_6H_5	2	PPE/$CHCl_3$/MW/70°	98	31
43	B-2	H	2,4-$Cl_2C_6H_3$	2	PPE/$CHCl_3$/MW/70°	96	31
44	B-2	4-Cl	2,4-$Cl_2C_6H_3$	2	PPE/$CHCl_3$/MW/70°	97	31

[a] SF = solvent-free conditions. [b]MW = microwave heating.

In general, cyclization reactions usually required long reaction times and high temperatures, resulting in lower product yields in some cases. The traditional methods have been modified to improve their efficiency, optimizing the routes of synthesis with the use of new technologies. Microwave irradiation has emerged as an efficient technique for reagent activation in organic reactions. The remarkable advantages of this methodology are the simple experimental procedures, high yields of products, short reaction times, mild conditions and easy work-ups. In this context, a large number of organic reactions can be carried out under microwave irradiation and compared with classical synthesis procedures.[83-87]

Using this methodology, Orelli *et al.* have presented a simple and efficient microwave-based protocol for the synthesis of cyclic amidines through an PPE-promoted cyclodehydration of *N*-aryl-*N*′-acylalkylenediamines, using a modified domestic microwave[81] (Table 1, entries 11-15). Employing microwave heating and a chloroformic solution of PPE, 1-aryl-1,3-diazepines and diazocines could be obtained with satisfactory (240 W, 2.5 min and 320 W, 6 min respectively.)

The microwave-assisted ring closure of *N*-aryl-*N*'-acyltetramethylenediamine derivatives promoted by PPE allowed the synthesis of 1-aryl-2-alkyl-1,3-diazepines (Table 2, entries 16-23).[75] The cyclodehydration was

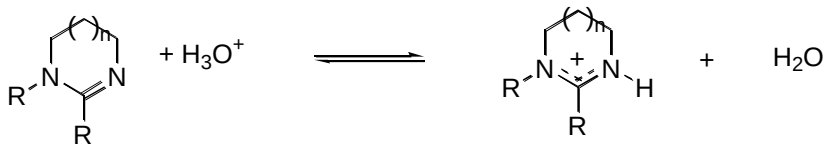
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carried out in a Monowave 300 monomode reactor. The reactions were completed in 8 min at 100°C with 63-90% yield. Under similar conditions, however, considerably lower yields of homologous 1-(4-methylphenyl)-2-alkyl-1,3-diazocine were obtained (Table 1, entry 24).[75] Alternatively, the use of PPSE as cyclodehydrating agent under solvent-free conditions in the microwave-assisted ring closures of *N*-acyl-*N*′-arylpentamethylenediamines allowed obtaining acceptable yields of 1-aryl-2-alkyldiazocines (Table 1, entries 25-33).[82] On the other hand, 1,2-diarylderivatives of 1,3-diazepines and diazocines were obtained by cyclization of either the corresponding *N*-aryl-*N*′-aroyltetra- or penta- methylenediamine (Table 1, entries 34-38).[31,32]

3. Chemical Properties of Cyclic Amidines

3.1 Basicity

Cyclic amidines are monobases which, upon protonation on N-3, become cyclic amidiniums salts that are strongly stabilized by mesomeric effect (Scheme 24).[35,36,74]



Scheme 24

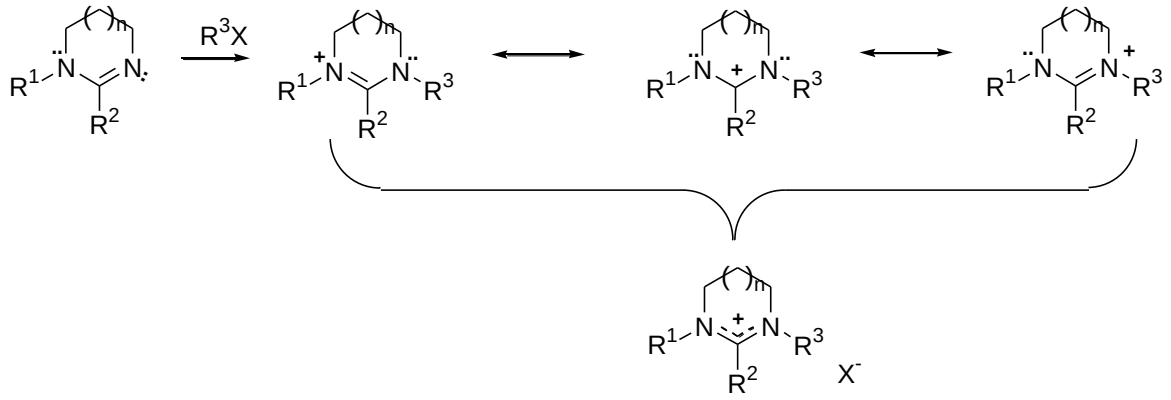
Upon comparing the basicity of the seven and eight-membered 1,2-diaryl substituted cyclic amidines (n=2,3) with that of the lower amidine (n=0,1), it was observed that basicity decreases in the order tetrahydropyrimidines (n=1) > tetrahydrodiazepines (n=2) > hexahydrodiazocines (n=3) > imidazolines.[35,36,74] This phenomenon was attributed to the possible torsion of the seven and eight-membered rings that may result in a less favored delocalization of the amidinium charge, and consequently in a decrease in basicity.

The effects of the sustituents on the N-1 aryl, were analyzed in a series of 1-aryl-2-phenyltetrahydro-diazepines.[35,36]

3.2 Nucleophilic character

Like other cyclic amidines,[88-91] 1-substituted tetrahydrodiazepines and hexahydrodiazocines have a strong nucleophilic character due to the N-3 lone electron pair. The reaction with alkyl halides leads to the corresponding resonance-stabilised cyclic amidinium salts **24**.[31-33,35] Since the reaction is a typical S_N2 displacement, it is adequate for the introduction of primary alkyl groups (Scheme 25).

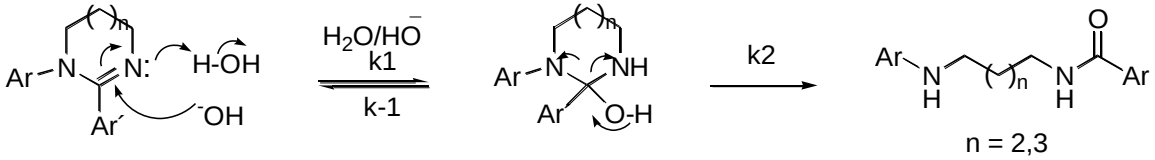
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Scheme 25

3.3 Hydrolysis

1,2-Diaryltetrahydrodiazepines and hexahydrodiazocines are resistant to acid hydrolysis due to the high stability of the amidinium ion.[36] However, due to their amidinic nature, they are hydrolyzed in alkaline solutions affording N-acyl derivatives of the corresponding tetra- and pentamethylenediamines. The observed regioselectivity was analyzed in the light of the stereoelectronic control theory (Scheme 26).[35,36]



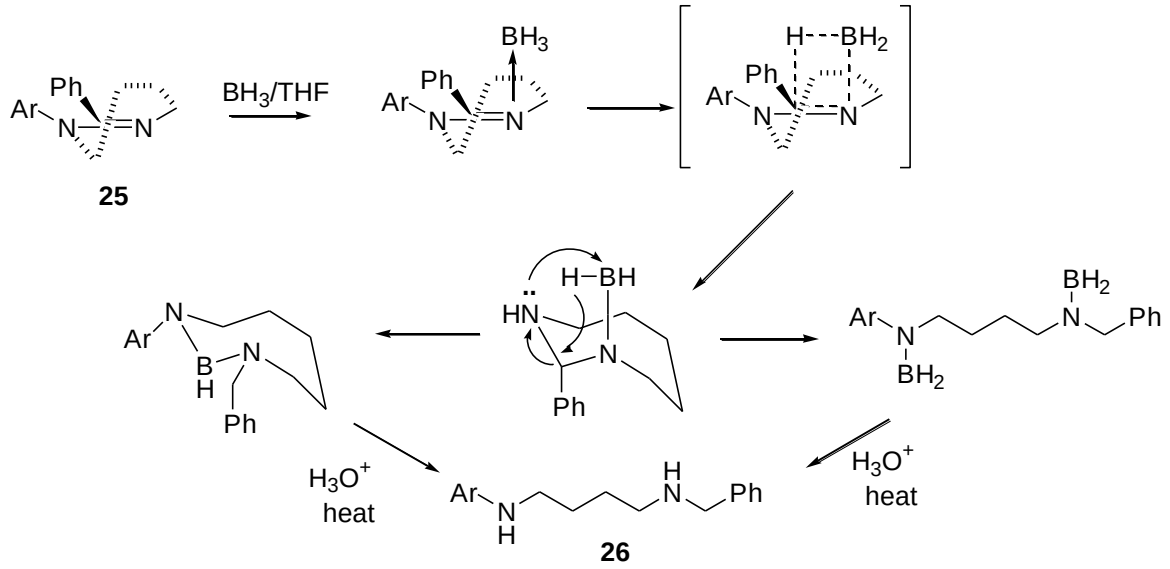
Scheme 26

The resistance of cyclic amidines to alkaline hydrolysis depends on the cycle size. A comparison of half-lives of 1,2-diaryl derivatives has demonstrated that the degree of stability is: imidazolines (n=0) < tetrahydropyrimidines (n=1) << tetrahydrodiazepines (n=2) < hexahydrodiazocines (n=3).[74] The greater stability of the larger rings was attributed to conformational factors that prevent the attack of the nucleophile OH^- on C-2.

3.4 Reduction

Reduction of 1,2-diaryltetrahydrodiazocines **25** with borane/THF leads to the regiospecific asymmetrical *N*-aralkyl-*N*'-aryltetramethylenediamines **26** with good yields (78-81%)[33,35] (Scheme 27). The following reaction mechanism explains the observed regioselectivity (Scheme 27) :

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Scheme 27

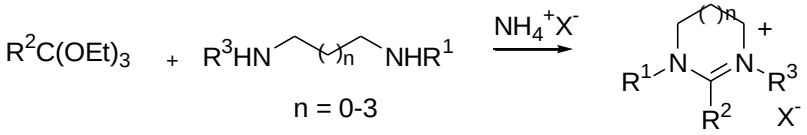
4. Synthesis of Cyclic Amidinium Salts

As indicated above (3.2) cyclic amidinium salts are typical salts where the cation is resonance-stabilized and the positive charge can be delocalized either on the nitrogen atoms or on the C-2 (Scheme 25).

As in the case of cyclic amidines, the methods of synthesis for cyclic amidinium salts of medium size, are in general, extensions of the methods employed for lower cyclic amidinium salts. These methods employ both cyclic and acyclic compounds as precursors.

4.1 Synthesis of amidinium salts from acyclic precursors

The treatment of **N,N'-disubstituted α,ω-alkanediamines with trialkyl orthoesters** in the presence of a source of protons and anions (ammonium tetrafluoroborate, ammonium hexafluorophosphate, ammonium chloride), leads to the corresponding cyclic amidinium salts. The drawback of this method is that it is not applicable to acid-sensitive substrates. This method employing alkyl orthoformates as C-1 building block is the method of choice for obtaining 2-unsubstituted salts (R^2 = H, Scheme 28), which are important as NHC precursors.



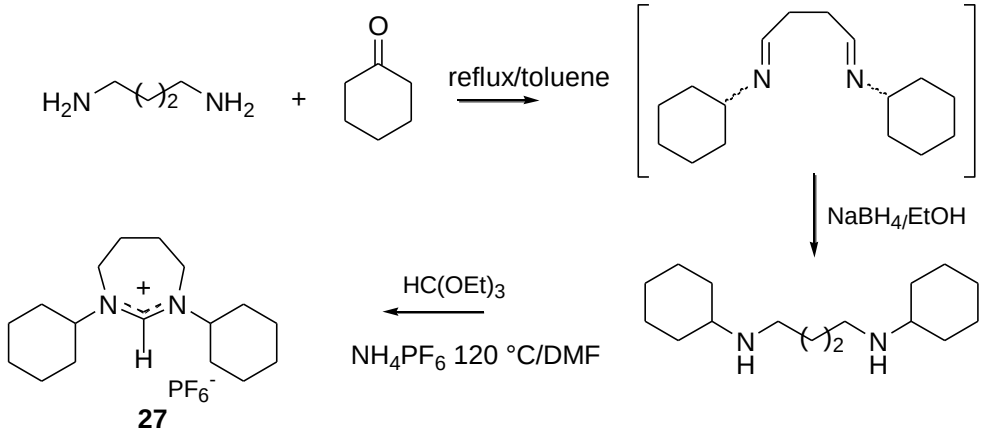
Scheme 28

The synthetic routes for obtaining the precursor diamines vary according to the nature of the linking backbone, the nitrogen substituents and the presence of chirality. This issue has been extensively addressed by César *et al.*[92]

The first report on the synthesis of seven-membered cyclic amidinium salts dates from 1991, when Saba *et al.*[93] prepared a series of cyclic amidinium salts by the reaction of trialkyl orthoesters with various *N*,*N*'-dialkyl-α,ω-alkanediamines in the presence of ammonium tetrafluoroborate or hexafluorophosphate. Among others, 2-methyl, 2-ethyl and 2-isopropyl substituted tetrahydro-1,3-diazepinium salts were obtained.

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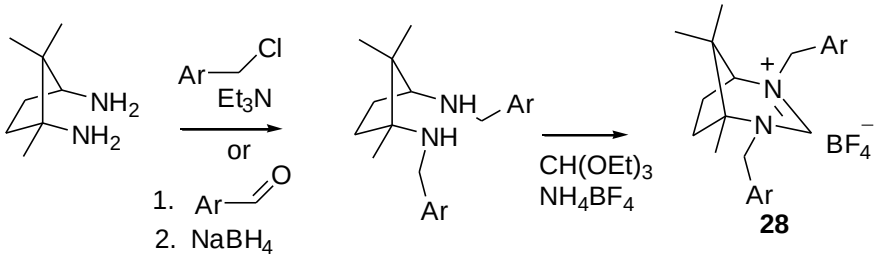
Several diazepinium salts were synthesized later to be used as precursors in the synthesis of seven-membered *N*-heterocyclic carbenes (NHC). In this sense, Iglesias *et al.* have obtained 1,3-dicyclohexyl derivatives **27** through the reaction of *N,N'*-dicyclohexylputrescine with triethyl orthoformate in the presence of ammonium hexafluorophosphate.[94] The diamine was obtained in high yields by condensation of 1,4-diaminobutane followed by reduction of the formed di-imine with sodium borohydride. Overall yields of 70% after recrystallization were obtained for the formation of the amidinium salt (Scheme 29).



27

Scheme 29

Çetinkaya *et al.* have reported the synthesis of 1,3-dibenzyl and 1,3-diheteroarylmethyl diazepinium salts with good yields.[95,96] More recently, Wilhelm *et al.* has reported the synthesis of 1,3-dibenzyl and 1,3-di-(α-phenylethyl)diazepinium salts to be employed as organocatalysts[97] Diazepinium salts embedded in a rigid bicyclic system containing a core derived from camphor **28** were obtained by cyclization of 1,3-diamino-1,2,2-trimethylcyclopentane (camphoric diamine) with triethyl orthoformate.[68] These salts were employed as precursors of enantiopure NHCs (Scheme 30).



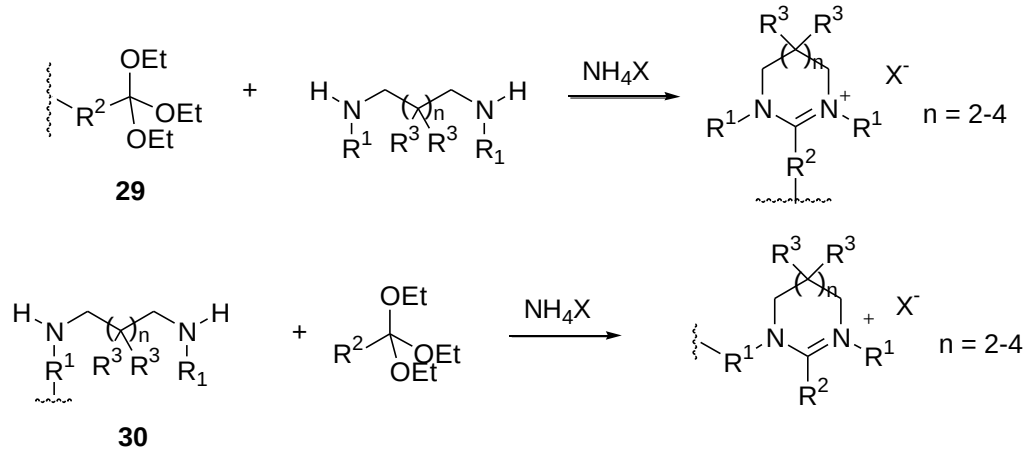
28

Scheme 30

Employing a similar methodology, Newman *et al.* have synthetized salts containing aryl groups bearing electron donor groups as precursors of tridentate ligands.[98,99]

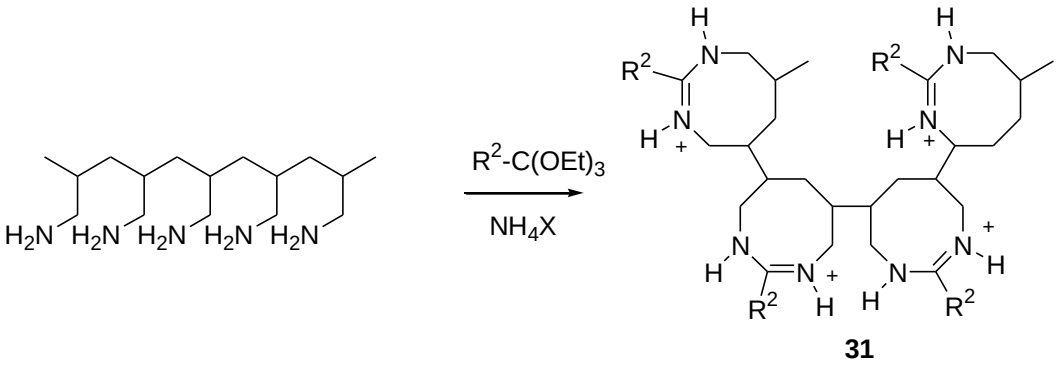
In 2005, a method was patented in which orthoesters were used as precursors to prepare cationic polymers bearing cyclic non-aromatic units containing an amidinium group, such as tetrahydrodiazepinium salts, among others.[100] One strategy to introduce a cyclic amidinium group into a side chain of the polymer is either to start out from a polymer which bears an orthoester group **29**, preferably an ethyl orthoester, in the side chain and allow it to react with an *N,N'*-dialkyl-α,ω-alkanediamine, or to start out from a polymer which bears the diamine function **30** in the side chain and allow it to react with an orthoester, preferably an ethyl orthoester (Scheme 31).

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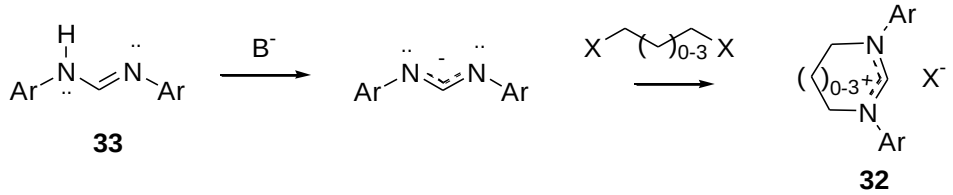
Scheme 31

In the same patent, polymers in which cyclic amidinium cations are located in the main chain and are linked to it via C atoms are described. Thus, for example, the reaction of polyamine with an orthoester leads to the formation of polymers **31** eight-membered rings (cyclic diazocinium ions) in the main chain (Scheme 32).



Scheme 32

The reaction of an **N,N'-disubstituted amidine with an α,ω-dihalo compound** in basic medium has been successfully used for the synthesis of N,N'-diaryl substituted cyclic amidinium salts **32**. From a mechanistic point of view, in the basic medium the deprotonation of the amidine **33** generates an 1,3-diazaallyl anion, which reacts with a dielectrophile such as the α,ω-dihalo compound (Scheme 33).

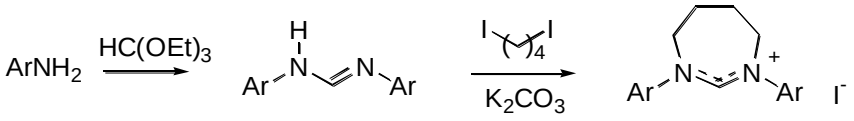


Scheme 33

Thus, 2-unsubstituted tetrahydrodiazepinium salts have been synthesized by Cavell through the reaction of the appropriate *N,N'*-diarylformamidine with 1,4-diiodobutane in refluxing acetonitrile in the presence of a mild base such as potassium carbonate.[101,102] The reaction proceeds rapidly for the larger ring sizes and less congested amidines. Formamidine precursors are easily obtained through the reaction of triethyl orthoformate with anilines. This synthetic strategy has been especially employed for N-substituted

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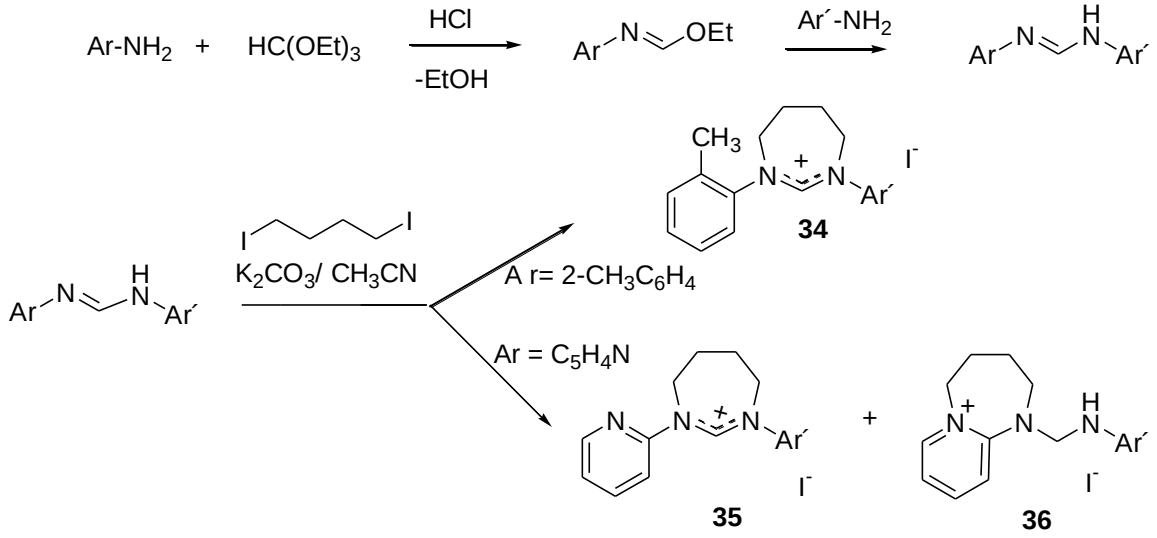
compounds bearing bulky aryl groups (Scheme 34). This method has been used with good results for the synthesis of seven-membered salts with different N-aryl substituents.[103-106]



Scheme 34

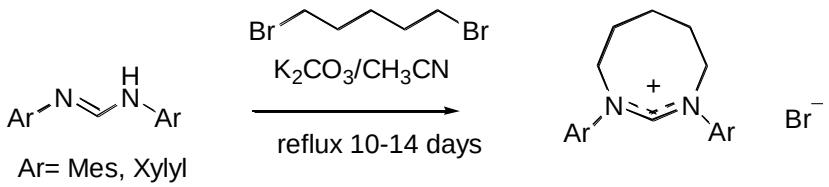
Similarly, Nechaev *et al.* have synthesized six- and seven-membered ring salts with bulky aryl groups by reaction of neat *N,N'*-diarylformamidine with 1,4-dibromobutane in the presence of diisopropylethylamine (DIPEA).[107,108] This method has recently been extended to the synthesis 1,3-dialkyl diazepinium salts.[109] The formamidine precursor was obtained through the reaction of the alkyl amine with triethyl orthoformate and one equivalent of acetic acid.

Asymmetrically 1,3-substituted rings **34** may be generated in good to high yields using a formamidine obtained by a step-wise reaction sequence.[106] During the synthesis of the seven-membered salts containing a pyridine substituent **35**, an alternative ring closure, via of the pyridine ring nitrogen, was observed, giving rise to a novel ionic fused ring product **36** (Scheme 35).



Scheme 35

Employing this method, Cavell *et al.* have recently synthesized the first eight-membered cyclic amidinium salts to be employed as NHC precursors (Scheme 36). The reaction was slow but yields were good (≥75%).[110]

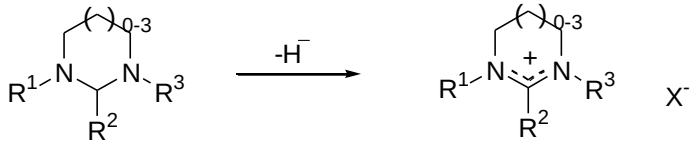


Scheme 36

4.2 Synthesis of amidinium salts from cyclic precursors

One of the method involves the **dehydrogenation of cyclic aminals** (Scheme 37).

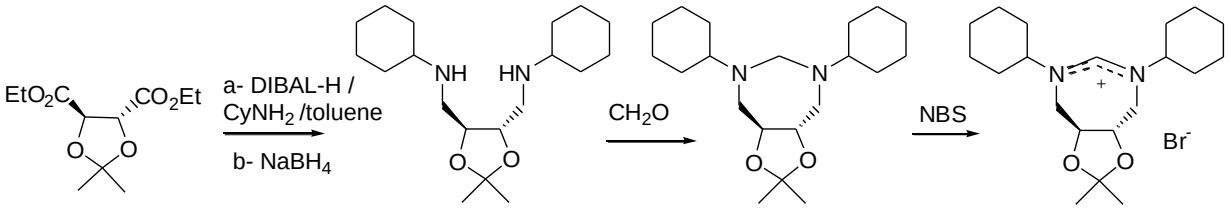
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37

Scheme 37

The cyclic aminal **37** is commonly obtained from the corresponding α,ω-diaminoalkane and an aldehyde. Formaldehyde is used as C-1 building block to obtain 2-unsubstituted salts. In this method, the cyclization step requires neutral conditions, therefore, it is applicable to acid-sensitive substrates. Probably, for these reasons, Iglesias *et al.*[94] have used this route for the synthesis of a seven-membered ring salt **38** containing a strained 5,6-dioxolane moiety using NBS as a dehydrogenating agent (Scheme 38).

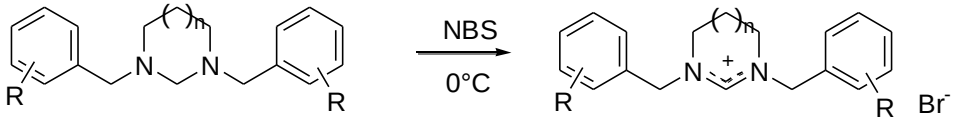


38

Scheme 38

Similarly, Wilhelm[97] has reported the synthesis of analogous 1,3-dibenzyl derivatives using NBA as dehydrogenating agent.

The synthesis of 1,3-dibenzyltetrahydrodiazepinium and hexahydrodiazociniun salts **39** through the dehydrogenation of aminals with NBS has recently been described (Scheme 39).[31]



39

n =2, R = H; 2,3-Cl_2; 3,4-Cl_2; 4-Cl, 4-CH_3

n =3, R = H; 2,3-Cl_2; 3,4-Cl_2; 3-Cl

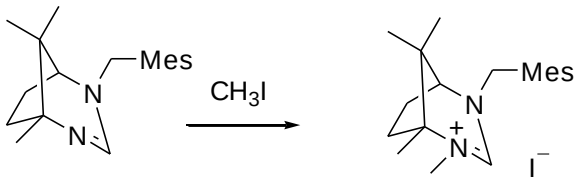
Scheme 39

As mentioned in Section 3.2, the **alkylation of 1-substituted cyclic amidines** affords the corresponding cyclic amidinium salts (Scheme 25). Since the reaction is a typical S_N2 displacement, it is adequate for the introduction of primary alkyl groups, being a method of choice for the synthesis of asymmetrically substituted salts. Thus, series of 1,2-diaryl-3-methyl-1*H*-4,5,6,7-tetrahydro-1,3-diazepinium iodides have been obtained through the reaction of the corresponding diazepines with methyl iodide in anhydrous THF under reflux, obtaining yields of 81-90 % in reaction times of 1-2 h.[33,35] More recently, this methodology has been optimized by the use of MW irradiation, the reaction times were dramatically decreased (3-6 min) using a Microwave Digestion System in chloroform solution at 90 °C and 400 W.[31] However, when the reaction of 1,3-diazocines were conducted under the same conditions, the major product obtained was the corresponding hexahydro-diazocine hydrohalide.[32] The protonation of the amidine could be avoided by using a mixture of DCM-DMSO (10:1) as solvent. Working under reflux conditions with conventional heating, 2-4 h were required for total

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conversion (71-95%), while the reaction times were reduced to 6-15 min when MW irradiation was employed (85-96% yields).[31,32]

Diazepinium salts with a rigid bicyclic system **40** (camphor skeleton*)* were also obtained by alkylation (Scheme 40).[68]



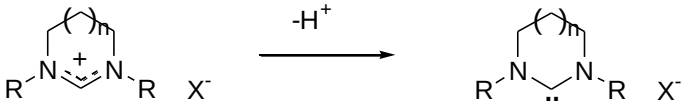
40, 78%

Scheme 40

5. Chemical Properties of Cyclic Amidinium Salts

5.1 Reaction with bases

5.1.1 Cyclic amidinium salts as precursors of N-heterocyclic carbenes (NHC). The deprotonation of 2-unsubstituted cyclic amidinium salts leads to the generation of NHCs **41**.[92] These compounds are of special interest due to their electron richness. Consequently, they have been widely applied as ligands in transition-metal catalysts and organometallic chemistry,[111-115] and as organocatalysts in their own right (Scheme 41).[116-120]



41

Scheme 41

In particular, tetrahydrodiazepinium salts have been synthesized to be employed as precursors of ring expanded NHCs (RE-NHCs), which are stronger σ-donating ligands.[120] Structurally, they also have unique features: the saturated seven-membered ring is flexible, highly twisted, which provides an opportunity to design novel chiral ligands, and of considerable interest due to the large heterocyclic rings with large N-C_{NHC}-N angles. Key features of these ligands are the presence of an increased basicity and a a high "steric" pressure on the metal center with respect to the more traditional five-membered NHCs.

Different bases have been employed to generate ER-NHCs: $KN(SiMe_3)_2$ (potassium hexamethyl-disilylamide, KHMDS) in THF;[101-103,106,109,121] $LiN(iPr)_2$ (lithium diisopropylamide, LDA) in toluene,[101] t-BuOK in water/DMF, [96,104,105,122] Ag_2O (silver oxide) in DCM[102,107] and potassium carbonate.[95]

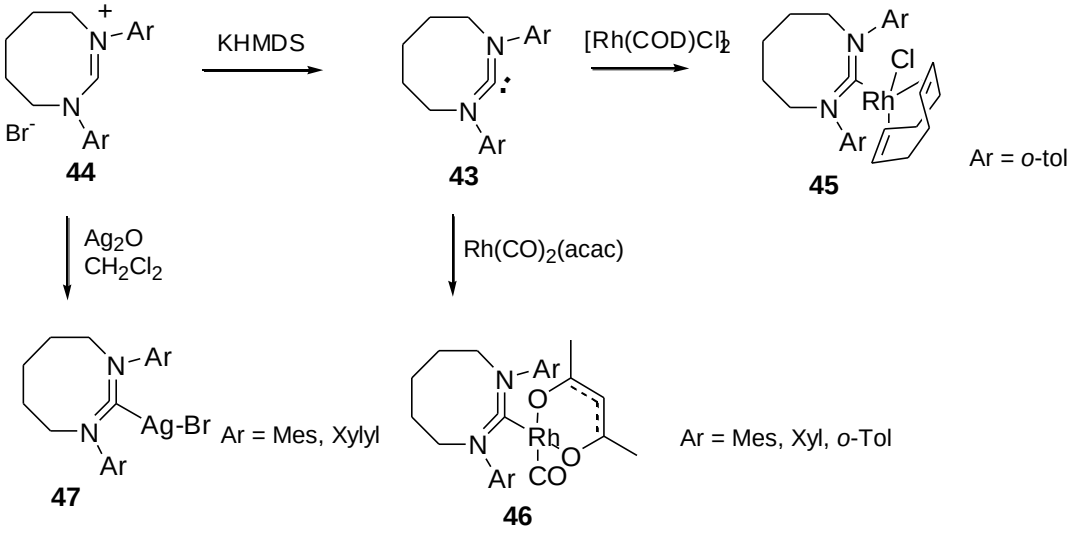
In some cases, seven-membered free carbenes have been isolated and characterized. [102,108,121] However, they are generally obtained as metal complexes by direct reaction of the *in situ* generated carbenes with suitable metal salts. Thus, the generation of Pd-containing complexes have been reported.[95,96,105,122] Other complexes with Pt,[104] Au,[123] Rh and Ir,[98,103,106,109] have also been reported. Seven-membered NHC complexes of Cu and Pd have been obtained by transmetallation of the corresponding NHC-Ag (I) complex with suitable metal salts.[107,108]

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The use of seven-NHC metal complexes in catalytic transformations have provided encouraging results. The first example has been reported by Cetinkaya *et al.,* who demonstrated that *in situ* generated 1,3-dibenzyltetrahydrodiazepin-2-ylidene palladium complexes are very effective in Suzuki-Miyaura coupling reactions of deactivated aryl chlorides.[95] Since then, seven-membered NHC complexes have been tested for catalytic transformations such as the Heck type cross-coupling,[96,105,122] Suzuki type reactions,[10,99] hydration of internal alkynes,[123] hydrosilylation,[99,104] catalytic hydrogenation[106] and transfer hydrogenation.[124,125]

The important results achieved with complexes bearing expanded ring NHC ancillary ligands in catalytic transformations have been attributed to the strong binding of the electron-rich carbene to the metal center that helps the metal retain its ligand, which provides the compound a longer catalyst life time, thus affording enhanced activity[106, 126]

Cavell *et al.* have reported the synthesis of the first eight-membered ring (diazocanylidene) NHCs **43** (8-NHCs) through the reaction of the corresponding cyclic amidinium salt **44** with KHMDS.[110] In general, the free RE-NHCs could be isolated, and in one case, the molecular structure was elucidated. Rh complexes **45, 46** have been formed through the treatment of the *in situ* formed free carbene with the appropriate Rh precursor complex. Silver complexes **47** have been prepared through direct reaction of Ag_2O with the diazocanylidinium salts (Scheme 42).

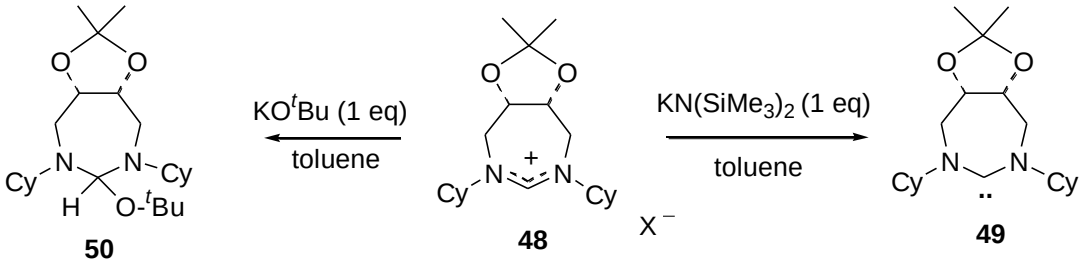


Scheme 42

Key features of these novel RE-NHCs are the extreme steric strain they impose on the metal center and their high electron donor capacity, being some of the most basic NHCs currently available.

5.1.2 Adduct formation. In some cases, the reaction of 1,3-diazepinium salts with alkoxides leads to the addition products (2-alkoxyaminals). In this sense, while the 1,3-dicyclohexyl salt **48** reacts with strong bases affording the expected NHC **49**, the corresponding adduct **50** was obtained when the salt was treated with potassium *tert*-butoxide in toluene (Scheme 43).[101]

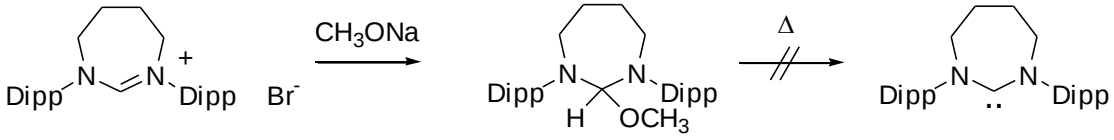
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50 **48** **49**

Scheme 43

Similarly, the adduct (7-dipp)(H)(OMe) **51** has been obtained by the reaction of equimolar amounts of the salt and NaOMe in absolute THF (Scheme 44). Attempts to obtain the NHC by vacuum thermolysis have failed.[107]



[(7-Dipp)H][Br] **51** (7-Dipp)(H)(OCH$_3$) 7-Dipp

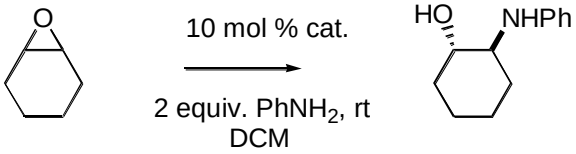
Dipp= diisipropylphenyl

Scheme 44

5.2 Catalytic activity

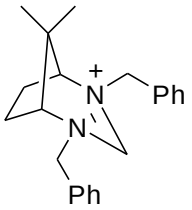
Cyclic amidinium salts have low Lewis acid character due to the contribution of the mesomeric structure with the positive charge on the C-2 (Scheme 25). These salts have been used as organocatalysts.[127-129]

Wilhelm *et al.* have reported the application of several cyclic amidinium salts as catalysts in the ring opening of epoxides (Scheme 45).[97] They tested four 1,3-diazepinium salts (**52**-**55**) and their activity was compared with the imidazolinium salt **56** (Figure 2). While the salt **56** displayed very low catalytic activity (12%), diazepinium salts **53** and **55** gave the product in 78% and 99% yield, respectively.

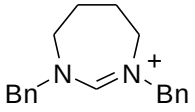


Scheme 45

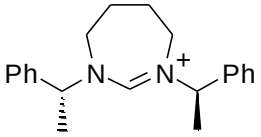
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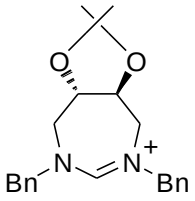
52 $[3,5\text{-}(CF_3)_2C_6H_3]_4\bar{B}$



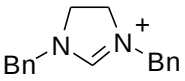
53 $[3,5\text{-}(CF_3)_2C_6H_3]_4\bar{B}$



54 $[3,5\text{-}(CF_3)_2C_6H_3]_4\bar{B}$



55 $[3,5\text{-}(CF_3)_2C_6H_3]_4\bar{B}$



56 $[3,5\text{-}(CF_3)_2C_6H_3]_4\bar{B}$

Figure 2

These results were attributed to the special geometry of the 1,3-diazepinium cation which does not allow the planar conformation to be kept, thus the positive charge is less delocalized over the NCN atoms. Instead, there is a better delocalization of the positive charge in the planar sp^2-centered imidazoline scaffold in the imidazolinium salts. The camphor-based salt **52** and the salt **54**, with a larger steric environment around the amidinium unit, gave yields of only 24% and 38%, respectively.

Acknowledgements

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