SYNTHESIS OF PYRIDOACRIDINES

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Abstract - The synthesis of pyridoacridines is reviewed.

A review of quinoline/isoquinoline-containing sea alkaloids revealed that more than 30 compounds, reported thus far, have as a common structural feature a pyrido[2,3,4-kl]acridine (dibenzo[f,ij][2,7]naphthyridine) nucleus. This group contains substances with a variety of biological activities: Ca-releasing, antiviral, antimicrobial activity, and cytotoxicity to L1210 murine leukemia cells. Norsegoline is representative of the structurally simplest of the group; amongst pentacyclic examples, amphimedine, ascididemine, and meridine include an additional pyridine (pyridone) ring, dercitine has an additional thiazole, the kuanoniamines an isomerically fused thiazole, and shermilamine A an additional thiazine. The segolines incorporate an isoprene unit, generating hexacyclic molecules, and eilatine is a 'dimer' of the nucleus.
In view of the considerable interest in such polyheterocyclic substances, the range of biological activities displayed, and the expectation that further natural, polyheterocyclic, poly-nitrogen-containing substances will be uncovered having variations on the structural types already known, we review here the available methods for the construction of pyrido[2,3,4-kl]acridines and, in addition, of all the other, currently known, isomeric pyridoacridines. The review concentrates on the methods which have been used to generate the pyridoacridine nuclei, and generally does not give complete natural product syntheses, even where these have been reported.

**1H-Pyrido[2,3,4-kl]acridines**

Necatorone (1), which has considerable mutagenic activity in the Ames test, has been synthesised. The tetracyclic ring system was established in an interesting oxidative process (Scheme 1), believed to involve a quinone imine: isoquinoline (2, R=NH₂), assembled in routine manner, was exposed to air in alkaline solution, however the ring closure could only be achieved at all efficiently on a 10 mg scale. An attempt to close the nitro compound, (2, R=NO₂) with triethyl phosphite, resulted in cyclisation of nitrene onto the isoquinoline ring nitrogen atom.

**Scheme 1**  
Reagent: i, O₂, 5% aq. NaOH (67%).
Cyclisation of 9-acetamido- (R₁=Me) and 9-formamido- (R₁=H) -benzo[a]acridines (R₁=H) (3) in a melt of AlCl₃/NaCl was employed in the synthesis (Scheme 2) of pentacyclic benzo-pyrido[2,3,4-k]acridines (R₁= H or Me, R₂=H or Me) (4)\(^{15}\).

\[ 
\begin{align*}
\text{R}^2 & \quad \text{NH}_2 \\
\text{N} & \quad \text{R}^2 \\
\text{N} & \quad \text{NHCO}R^1 \\
\text{N} & \quad \text{R}^1 \\
\text{N} & \quad \text{N} \\
\text{R}^2 & \quad \text{NHCO}R^1 \\
\end{align*}
\]

\[ 
\begin{align*}
\text{R}^2 & \quad \text{NH} \\
\text{N} & \quad \text{R}^2 \\
\text{N} & \quad \text{R}^1 \\
\text{N} & \quad \text{N} \\
\text{R}^2 & \quad \text{NHCO}R^1 \\
\end{align*}
\]

Scheme 2  Reagents: i, Ac₂O, 100°C (92%) or DMF, NaOMe, reflux (52%); ii, AlCl₃, NaCl, 220°C (58-64%).

**1H-Pyrido[3,4,5-k]acridines**

There are no reported examples of this ring system, either natural or synthetic.

**1H-Pyrido[2,3,4-k]acridines**

This, the ring system common to more than 35 marine alkaloids, has been the subject of considerable synthetic effort, and five distinct routes to the nucleus have been described.

Oxidation of the methyl group in 2-methoxy-9-methylacridine and condensation with methyl azidoacetate gave an azide, (5), which on thermolysis generated a nitrone for an insertion (Scheme 3) producing the tetracyclic nucleus\(^{16}\).

\[ 
\begin{align*}
\text{CO}_3\text{Me} & \quad \text{N}_3 \quad \text{OME} \\
\text{N} & \quad \text{OME} \\
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\end{align*}
\]

Scheme 3  Reagent: i, xylene, 140°C (97%).

In two total syntheses of amphimedine, (6), a simple N/C bonding from advanced *para*-quinonoid intermediates completed the nucleus (Scheme 4). Quinone (7)\(^{17}\) was
prepared by coupling 5,8-dimethoxyquinolin-4-yl triflate with a 2-trifluoroacetylamino phenyltrimethyltin, and the pyridone added using a Diels-Alder cycloaddition with Ghosez' diene; quinone (8) was generated using a standard quinolone ring synthesis producing a 4-(2-nitrophenyl)quinolin-2-one, the pyridone again being added later using Ghosez' diene.18

![Scheme 4](image)

Scheme 4  Reagents: i, HCl, THF, 80°C (86%); ii, Me2SO4, K2CO3, DMF, 23°C (96%); iii, H2, 10% Pd-C, Et3N, MeOH, room temperature (13%).

In a synthesis of 'isoascididemine', (9), cycloaddition between a quinoline-5,8-dione and the dimethylhydrazone of acrolein added a pyridine in an orientation isomeric with both ascididemine and amphimedine; again the final ring closure involved a straightforward amine/carbonyl condensation (Scheme 5).19

![Scheme 5](image)

Scheme 5  Reagents: i, CH3CN, Et2O, 23 → 80°C; ii, 6M HCl, 80°C (92%).

Intramolecular Friedel-Crafts type acylation of the 2-pyridone, at its C-3, in intermediate (10), in a third total synthesis of amphimedine, is a quite different method for the generation of a pyrido[2,3,4-kl]acridine (Scheme 6).20 Substrate (10) was produced in a short route which neatly utilised the Schmidt reaction of a pyridyl-aza-fluorenol with migration of the most electron-rich ring.
Nitrene insertion was also central to a route (Scheme 7) by which cystodytines A and B were synthesised, thus thermolysis of the ortho-quinone (11) produced 12 (R=OH); a better route involved cyclisation at lower oxidation level, for example in the conversion of 13 into 12 (R=H).21

The pyrido[2,3-b]acridine (14) was cyclised to afford ascididemine, (15), using the side-chain enamine, formed by reaction of a precursor methyl group with dimethylformamide dimethyl acetal (DMFDMA), as an acetaldehyde synthon (Scheme 8).22
The same alkaloid has also been constructed in a quite different manner, by the photo-catalysed cyclisation of the iodo-quinone imine (16) (Scheme 8), produced in only three steps from phenanthroline.

**Benzof[\textbf{j}][1,7]\textit{phenanthroline (Pyrido[2,3-\textbf{b}]acridine)}**

The reaction of phloroglucinol with anthranilic acid, \textit{via} three Bucherer-type reactions, leads to a dibenzo[1,7]\textit{phenanthroline} (Scheme 9).

![Scheme 9](image)

**Scheme 9**  \textbf{Reagent}: i, heat.

A triple Ullman-amine coupling of 1,3,5-tribromobenzene, then three intramolecular Friedel-Crafts cyclising acylations generated ‘triquinolonobenzene’, (17) (Scheme 10). The compound was prepared for comparison with linear ‘quinacridone’ (see later) systems, organic pigments with outstanding light fastness, believed to be associated with a high degree of intermolecular hydrogen bonding. Heptacycle (17) is only weakly coloured.

![Scheme 10](image)

**Scheme 10**  \textbf{Reagents}: i, K$_2$CO$_3$, spongy Cu, CuCl, Cu(OAc)$_2$.H$_2$O, n-C$_5$H$_{11}$OH, reflux (92%);
ii, PPA, 140-150°C, 4.5 h (98%).
Ullmann-amine coupling with 7-aminoquinoline provided the carboxylic acid (18), which was converted into acid chloride, cyclised selectively to the quinoline 8-position, and acridone thus produced converted into the corresponding halo-heterocycle, all in one pot (Scheme 11). This type of approach has been used to produce several isomeric pyridoacridines: Schemes 14, 27 and compound 37.

Scheme 11 Reagents: i, Cu-bronze, EtC(Me)2OH, 150°C, 6 h; ii, POCl3, PCl5, 150°C.

The dibenzo[1,7]phenanthroline, (19), was constructed (Scheme 12) from meta-phenylenediamine, which on reaction with 2-formylcyclohexanone gave 20, and this, on strong acid treatment, cyclised to give 21.

Scheme 12 Reagents: i, EtOH, room temperature (65%); ii, PPA, 160-170 °C (40%); iii, Se, 300°C (58%).

**Benzol[f][2,7]phenanthroline (Pyrido[3,4-a]acridine)**
There are no reported examples of this ring system, either natural or synthetic.

**Benzol[f][3,7]phenanthroline (Pyrido[4,3-a]acridine)**
There are no reported examples of this ring system, either natural or synthetic.
Benzo[b][4,7]phenanthrol ine (Pyrido[3,2-a]acridine)

In what appears to be an extraordinary regiochemical preference, the diacid (22) closes to the dibenzo[4,7]phenanthrol ine (23) via two acid-catalysed Friedel-Crafts intramolecular acylations (Scheme 13).  

\[ \text{Scheme 13 Reagent : i, c. H}_2\text{SO}_4, 100^\circ\text{C (59%).} \]

Acids, (24) (R\(^1\)=H or Cl and R\(^2\)=H or Cl), produced via Ullmann-amine couplings with 6-aminoquinolines, were cyclised (Scheme 14), in one preparative step involving conversion of acid to acid chloride, intramolecular Friedel-Crafts closure, selective for the quinoline 5-position, and conversion of initial acridone into halo-heterocycle.  

\[ \text{Scheme 14 Reagents : i, Cu-bronze, reflux, EtC(Me)\(_2\)OH (50-60%); ii, POCl}_3, 150^\circ\text{C (70-80%).} \]

The interaction of para-phenylenediamine with 2-formylcyclohexanone to generate a bis-enamine, followed by acid-catalysed closure to a mixture of isomers, separate dehydrogenation of which produced (Scheme 15) the dibenzo[4,7]phenanthrol ine, (25) and the dibenzopyrido[2,3-b]acridine (26).  

As in Scheme 12, interchange of amino-aromatic, and net transfer of nitrogen to the ring carbonyl carbon must be involved.
1,4-Diiodobenzene reacted in a double Ullmann-amine coupling with 2-aminoacetophenone to give a diketone, (27) which underwent acid-catalysed cyclisation with what would have appeared to be an unlikely regioselectivity, for the hindered dibenzo[4,7]phenanthroline, (28), was obtained (Scheme 16). It is significant that the bis-benzoyl analogue of 27 only underwent one cyclisation, giving an acridine. This synthetic approach has been used to produce several isomeric pyridoacridines (Schemes 25 and 32).

Hexa- and heptacyclic molecules containing a pyrido[3,2-a]acridine nucleus were generated (Schemes 17 and 18) by ring closures of anthraquinone-derived precursors.
Pyrido[2,3-b]acridine

The organic pigment, quinacridone (29) can be synthesised from diethyl 2,5-dioxo-1,4-cyclohexanedicarboxylate (30) and aniline then dehydrogenation (Scheme 19)\textsuperscript{35}; many analogues have been synthesised, and the large field has been reviewed\textsuperscript{36} and is accordingly only briefly summarised here. Alternative routes to such systems include cyclisations of 2,4-diarylaminoterephthalic acids which give the fully aromatic systems directly, or cyclisation of 2,4-di-(2-carboxyphenylamino)-1,4-benzoquinones, which give quinones, reducible to the parent system.

Interaction of diketone (30) with 2-aminobenzophenone in a double quinoline Friedländer synthesis, and dehydrogenation, provided (31) (Scheme 20).\textsuperscript{37} Simply heating 32 in 1-chloronaphthalene brought about double intramolecular acylation and the formation of 33.

The synthesis of compound (26) has already been discussed.\textsuperscript{31}
Scheme 19  **Reagents**: i, heat; ii, *e.g.* chloranil; iii, *e.g.* PPA; iv, *e.g.* c. H$_2$SO$_4$; v, *e.g.* Zn dust, 70% H$_2$SO$_4$.

Scheme 20  **Reagents**: i, 6N HCl, EtOH, reflux (72%); ii, chloranil, methylcellosolve, reflux (92%); iii, C$_{10}$H$_7$Cl, reflux (83%).
Pyrido[3,4-b]acridine
There are no reported examples of this ring system, either natural or synthetic.

Pyrido[4,3-b]acridine
There are no reported examples of this ring system, either natural or synthetic.

Pyrido[3,2-b]acridine
Strong acid-catalysed cyclisation of symmetrical diacid-diarylamine (34) affords the linear benzopyrido[3,2-b]acridine (35) (Scheme 21).38

Scheme 21  Reagent : i, PPA, 120-160°C.

The use of a decahydroquinolone-amide in a Friedländer synthesis provided 36.39

Scheme 22  Reagent : i, NaOH, EtOH (85%).

Benzo[b][1,7]phenanthroline (Pyrido[2,3-c]acridine)
The syntheses of compound (17)25 and of isoascididemine (9)19 have already been discussed.

The route described in Scheme 11 has been utilised, but with 5-aminoquinoline instead of 7-aminoquinoline, to prepare 37.40
β-Naphthol condenses (Scheme 23) with 5-aminoquinolines and paraformaldehyde generating naphthaleno[1,7]phenanthrolines (38, R = H and Me).\(^{41}\)

\[
\text{R} = \text{H, Me}
\]

**Scheme 23**  **Reagent**: i, paraformaldehyde, 250°C, (15%).

In a double Skraup synthesis (Scheme 24), 3,6-diaminoacridine produced the dipyridoacridine (39); the direction of ring closures, both adjacent to the acridine nitrogen, is notable.\(^{42}\)

**Scheme 24**  **Reagent**: i, HO\text{-}\text{CH}_2\text{-}\text{CH(OH)}\text{-}\text{CH}_2\text{-}\text{OH}, \text{H}_3\text{AsO}_4, \text{H}_2\text{SO}_4, reflux.

Scheme 25 shows how the condensation of 2-aminoacetophenone and 2-aminobenzophenone with 1,3-diiodobenzene produces 40, double ring closures of which give the angular systems (41 R = Me or Ph).\(^{33}\)

A large number of benzo[\(b\)]\{1,7\}phenanthrolines have been synthesised by the route shown in Scheme 26, in which an aminoacridine is converted into a aminomethylenemalonate, (42, \(R^1, R^2,\) and \(R^3=\text{H, and alkyl (C}_{1-10}\), Ar, alkylamino, alkylthio, nitro, cyano, alkylsulphonyl, etc.) and this cyclised in acid; again, the regioselectivity of cyclisation is interesting.\(^{43}\)
Scheme 25  Reagents: i, K$_2$CO$_3$, Cu, n-Bu$_2$O, reflux (75%); ii, H$_2$SO$_4$, AcOH (66%).

Scheme 26  Reagents: i, Ph$_2$O, 260°C.

**Benzo[b][1,8]phenanthroline (Pyrido[3,4-c]acridine)**
The synthesis of amphimedine, (6) has already been discussed (Schemes 4, 5, and 6).

A sequence of the type encountered before (Schemes 11 and 14) allowed the assembly of 43 (X = H or Cl) from 5-aminoisoquinoline (Scheme 27).

Scheme 27  Reagents: i, K$_2$CO$_3$, Cu, EtC(Me)$_2$OH, reflux (X=H (22%) and X=Cl (15%)); ii, POCl$_3$, reflux, (X=H (39%) and X=Cl (85%)).
In a manner similar to that described above for 5-aminoquinoline, β-naphthol combines (Scheme 28) with paraformaldehyde and 5-aminoisoquinolines giving naphthaleno[1,8]phenanthrolines (44, R=H or Me).41

Scheme 28  Reagent: i, paraformaldehyde, 250°C (40%), ii, Ph₂O, reflux (68%).

In a variation of this approach, the Mannich base derived from β-naphthol was used.48

In work (Scheme 29) relevant to the development of pyridone-fused acridines of the type found in amphimedine, coupling of 2-aminoacetophenone with 45 gave 46, then double cyclisation was achieved in one pot, in which one of the closures involved a Pummerer process.49

Scheme 29  Reagents: i, Na₂CO₃, Cu (cat.), PhNO₂, reflux (68%); ii, MCPBA, CH₂Cl₂, 0°C → room temperature (87%); iii, c. H₂SO₄, AcOH, 130°C (79%).
Benzo[bl,b]phenanthroline (Pyrido[4,3-c]acridine)
There are no reported examples of this ring system, either natural or synthetic.

Benzo[bl,b]phenanthroline (Pyrido[3,2-c]acridine)
The chloro compounds (47, R1=H or Cl and R2=H or F or OMe) were prepared by routes totally analogous to those shown in Scheme 11, but starting with an 8-aminoquinoline as the amine component.30,40,44,47,50-54

Skraup cyclisations (Scheme 30) were successful with 4-aminoacridine and with 4-aminoacridin-9-one.54

The amino-aldehyde (48) was prepared by condensation of 4-aminopyrimidine-5-carboxaldehyde with ketone (49) followed by release of the required functional groups by hydrolysis of the pyrimidine unit; the ketone and the ortho-amino-aldehyde groups were then utilised to add two more pyridine rings.55
Scheme 31  Reagents : i, 4-aminopyrimidine-5-carboxaldehyde, KOH, MeOH, reflux (93%);
ii, 2N HCl, reflux (97%); ii, O₃, MeOH, CH₂Cl₂, -78°C then Me₂S (100%, 2 steps)

2-Aminobenzophenone and 1,2-diiodobenzene underwent a double Ullmann-amine
coupling and now, in contrast to the isomeric *para* situation, double ring closure *could*
be brought about (Scheme 32) with strong acid.³³

Scheme 32  Reagents : i, K₂CO₃, Cu, n-Bu₂O, reflux (67%); ii, H₂SO₄, AcOH, (48%).

In a search for dyestuffs, each of the phenylenediamines was reacted with 1-
nitroanthraquinone and 1-amino-4-chloroanthraquinone as Ullmann partners, and
the resulting bisquinone cyclised with acid.⁵⁶  No details of structure are given, so that
the direction of cyclisation of the *meta*- and *para*-phenylenediamines is not described;
the *ortho*-isomer can of course close in only one way, and it is the cyclisation of the
product of condensation of this isomer with the former quinone which is shown in
Scheme 33.
The final step in the synthesis of 50 involves high temperature decarboxylation with concomitant dehydrogenation (Scheme 34); the skeleton was built up using two quinoline-type syntheses, the ortho-aminocarbonyl component being hydrolysed isatin and the other component cyclohexane-1,2-dione dioxime.57

Scheme 33  Reagents: i, Na$_2$CO$_3$, Cu bronze, PhNO$_2$; ii, c. 70% H$_2$SO$_4$, 160°C (73%).

Scheme 34  Reagents: i, H$_2$O, 1080°C (22%); ii, paraffin oil, N$_2$, 320°C (93%).

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References and Footnotes


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