Retraction

Biogenetic hypothesis and first steps towards a biomimetic synthesis of haouamines

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We, the named authors, hereby wholly retract this Chemical Communication. Signed: Edmond Gravel, Erwan Poupon and Reynald Hocquemiller, Châtenay-Malabry, France, November 2007.

Retraction endorsed by Sarah Thomas, Managing Editor.
A detailed hypothesis for the biogenesis of haouamines is reported herein, supported by experiments headed towards biomimetic synthesis of these compounds.

Marine organisms belonging to the Ascidiacea class are a rich source of structurally unique secondary metabolites. Among the most remarkable of these, Haouamines A and B were isolated by E. Zubia and co-workers in 2002 from Aplidium haouarianum collected off Tarifa Island (Spain). Haouamines A and B belong to a new family of complex polycyclic alkaloids (Fig. 1).

The molecular complexity of 1 constitutes an intellectual challenge for organic chemists. In fact, several groups have disclosed their own approaches to the construction of haouamines. These efforts culminated in a total synthesis by the Baran team in early 2006, soon followed by a formal total synthesis by Weinreb and co-workers. We wish to report in this communication the first biomimetically inspired approach to haouamine A featuring an expeditious assembly of an advanced intermediate towards the total synthesis of I.

The combination of ring systems into a single and small architectural unit like haouamine A leads one to wonder how such structures can be assembled in nature. The tetrahydropyridine center ring suggests that the biogenetic origin is likely to be an assembly of simple L-phenylalanine derived units which can be traced four times in the final structure of 1. Baran and colleagues suggested a condensation of four meta-hydroxylated phenylacetaldehyde molecules with ammonia as the nitrogen source followed by oxidative events. They also reported they had failed to take advantage of that route. We suggest a modified hypothesis. Central in our proposals is the formation of the tetrahydropyridine core through a natural Chichibabin-like pyridine synthesis reaction involving three aldehydes and a primary amine as the nitrogen source, all units being presumably derived from the aforementioned amino-acid.

Based on the fact that, in nature, amines can be transformed into aldehydes through an oxidative deamination process, we can assume that from L-phenylalanine, through hydroxylation to L-m-tyrosine and decarboxylation, amine 2 can be obtained and then undergo oxidative deamination to give rise to meta-hydroxylated phenylacetaldehyde 3 (Scheme 1). The condensation of 2 and three molecules of 3 will lead to the formation of dihydropyridinium 4 (that already contains all atoms of the final molecule) which can then be reduced to compound 5. The formation of the C8–C9 bond can be seen as the result of an ortho–para coupling that occurs under oxidative conditions to yield intermediate 6. Such conditions might as well lead to the oxidation of C26 and generate an allylic hydroxyle group (7). It has been reported by Rawal and co-workers that under acidic conditions, the remaining bond (C24–C26) is then easily formed, via carbocation formation, completing the biosynthesis of haouamine A (Scheme 2).

Biomimetic investigations were performed with this possible biosynthetic route taken into consideration, and it was decided to study the outcome of the condensation reaction of 3-methoxyphenethylamine 8 with 3-methoxyphenylacetaldehyde 9. The two species were reacted together (in a 1 : 3 ratio) along with catalytic quantities of ytterbium triflate in different solvents, at room temperature. After 20 hours, pyridinium 11§ was found as the major product, whatever the solvent was, with yields of 60 to 75%
The reaction involves a cascade of reactions similar to the ones depicted on Scheme 2 (up to product 4). Results highlight the great interest of rare earth-metal triflate catalyzed reactions in organic synthesis: the reaction takes place in very mild conditions instead of high temperatures and/or pressures historically required for the Chichibabin pyridine synthesis reaction. 

Ytterbium triflate plays the role of a Lewis acid and probably activates both aldehyde and imine groups of the reactants and intermediates. It should be noted that no traces of dihydropyridinium salt 10 (the logical outcome of the reaction) were detected in the crude reaction mixture. Spontaneous oxidations of dihydropyridinium salts into pyridinium salts have previously been observed with Chichibabin-like reactions, and appear to be especially prominent when condensations are carried out with phenylacetaldehydes.

Compound 11 was then easily transformed into 12 by reaction with tribromoborane (4.1 equiv.) in methylene chloride. The addition was performed at \(-78\, ^\circ\text{C}\) and the mixture was allowed to slowly warm up to room temperature within the next three hours. Compound 12 was obtained in very good yield without any column purification needed.

Our initial plan was to perform reduction of 11 with sodium borohydride but the reaction only gave very limited quantities of the desired product 14 as a mixture of cis/trans isomers

Scheme 3 Biomimetic self-assembly of pyridinium 11 followed by phenol deprotection to yield compound 12.

Scheme 4 Reduction of 11 into tetrahydropyridine 14.
(Scheme 4). The reason for such low yields is yet under investigation in our laboratory but it is likely to be an issue of pseudo-dimerisations through Diels–Alder type cycloadditions involving 14 and its direct precursor 13.13

With compound 14 obtained in too little amounts, it was decided to try out different types of oxidative conditions applied to product 12 in order to see if the desired C8–C9 coupling can take place despite high constraint resulting from the pyridinium’s planarity. The outcome of such reactions is still under investigation in our laboratory.

In conclusion, we have proposed a new detailed biogenetic hypothesis for haouamines that has led us to successfully achieve the first steps of the first biomimetic total synthesis of haouamine A. An advanced biomimetic precursor of that compound has been obtained through a convenient and efficient multicomponent reaction and the final two bonds that are needed to complete the synthesis are currently being investigated in our laboratory. The described biosynthetic hypothesis can serve as a useful framework for which to develop a coherent and straightforward synthetic plan towards haouamines (as well as close analogs such as compounds 11, 12, 14) that competes with other synthetic approaches. This example also contributes to demonstrate the power of biomimetically inspired strategies in total synthesis.14

Notes and references

† It is interesting to note that oxidative conditions under which radicals can be generated on phenols to make phenolic couplings possible may also generate free hydroxyl radicals responsible for the formation of 1-meta-tyrosine involved in our biogenetic proposal.

§ Compound 11: orange oil; Rf = 0.40 (silica gel, CH2Cl2-MeOH 9 : 1); IR (film, CHCl3): v max = 2937, 1585, 1260, 1030 cm−1; 1H NMR (400 MHz, CDCl3), δ 1.25 (2 H, s), 3.27 (2 H, t, J = 6.5 Hz), 3.67 (6 H, s), 3.86 (6 H, s), 5.15 (2 H, t, J = 6.5 Hz), 6.55–7.45 (16 H, m), 8.47 (1 H, s), 8.75 (1 H, s); 13C NMR (100 MHz, CDCl3), δ 29.6, 37.8, 55.1, 55.7, 63.5, 112.5, 113.5, 114.2, 116.6, 118.2, 119.6, 121.1, 122.6, 130.2, 130.8, 134.0, 136.9, 139.8, 140.5, 141.3, 160.2, 160.6; MS (ES) m/z 546 (M+), 440 (10), 426 (100); HRMS (ES) calcd for C36H36NO4+ 550.2919, found: 550.2921.


