R. L. GILES, J. D. SULLIVAN, A. M. STEINER, R. E. LOOPER* (UNIVERSITY OF UTAH, SALT LAKE CITY, USA)

Addition–Hydroamination Reactions of Propargyl Cyanamides: Rapid Access to Highly Substituted 2-Aminoimidazoles

**Synthesis of Substituted 2-Aminoimidazoles**

**Significance:** Looper and co-workers described a three-step synthesis of highly substituted 2-aminoimidazoles. The first step consists of the preparation of propargyl cyanamides by copper(I)-catalyzed addition of an iminium generated from condensation of aldehydes ($R^2 = \text{alkyl or aryl}$) with secondary amines. Without further purification, the resulting tertiary amines are subjected to a von Braun reaction. The use of 4-methoxybenzyl-, 2,4- and 3,4-dimethoxybenzyl-substituted propargyl amines lead to the propargyl cyanamides in good yields whereas symmetric $N,N$-dialkylpropargyl amines give exclusively propargyl bromide. Screening of catalysts led to the use of $\text{La(OTf)}_3$ for the final addition–hydroamination step. Both acyclic and cyclic secondary amines generated guanidine intermediates which underwent cyclization to the alkyne to afford 2-aminoimidazoles in good yields.

**Comment:** Several alkaloids containing the 2-aminoimidazole ring with interesting structures and biological properties have been isolated from marine sponges (S. M. Weinreb Nat. Prod. Rep. 2007, 24, 931). This scaffold improves physicochemical properties like lipophilicity, blood-brain barrier passage, cell permeability and bioavailability. Polysubstituted 2-aminoimidazole synthesis remains challenging as the current methods present some disadvantages: i) use of unstable precursors for the condensation of $\alpha$-amino/$\alpha$-haloketone with a cyanamide or a guanidine derivative respectively; ii) multi-step synthesis for the decoration of the imidazole scaffold. This three-step sequence represents a useful tool for the efficient synthesis of diversely substituted 2-aminoimidazoles from readily available starting materials.

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**Category**

- Synthesis of Heterocycles

**Key words**

- 2-aminoimidazoles
cyanamides
hydroamination
lanthanum(III)

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**SYNFACTS Contributors:** Victor Snieckus, Emilie David

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