Reissert Compound Studies. XL. The First Monocyclic Reissert Compounds

Seshadri Veeraraghavan, Debkumar Bhattacharjee, and Frank D. Popp*

Department of Chemistry, University of Missouri-Kansas City, Kansas City, Missouri 64110, USA Received January 15, 1981

Pyridazine and pyrimidine have been reacted with trimethylsilyl cyanide and benzoyl chloride to give the first examples of Reissert compounds derived from monocyclic systems.

J. Heterocyclic Chem., 18, 443 (1981).

Sir:

While the formation of Reissert compounds (1-4) from quinolines, isoquinolines, phthalazines, and other diazasystems (4) has provided a very useful synthetic intermediate (5), the extension of this work to monocyclic systems has not been actively persued because of the observation that pyridine (1,2) does not form a Reissert compound.

We have now found that pyrimidine reacts at room temperature with trimethylsilyl cyanide, benzoyl chloride, and a catalytic amount of aluminum chloride in anhydrous methylene chloride to give a 59% yield of the di-Reissert compound 1 (6) m.p. 148-151. The formation of 1,3-dibenzoyl-2,4-dicyano-1,2,3,4-tetrahydropyrimidine (1) from pyrimidine is analogous to the formation of the Reissert compound 2 from quinazoline (7).

A similar reaction of 3-methylpyridazine with trimethylsilyl cyanide and freshly distilled benzoyl chloride gave a 41% yield of the pyridazine Reissert compound 3 (6), m.p. 122-124°. When either pyridazine or 3-methylpyridazine was reacted under the above conditions with undistilled benzoyl chloride the deep red Reissert salts 4 (6) and 5 (6), m.p. 171-173° and 179-181°, respectively, were obtained.

Confirmation of this structure, which is analogous to the salt obtained by acid treatment of the phthalazine Reissert compound (8), was had by reaction of 4 with dimethylacetylenedicarboxylate in anhydrous dimethyl formamide at 100° to give 6 (6) m.p. 179-180°.

The reactions of these new and novel Reissert compounds are being studied in detail.

REFERENCES AND NOTES

- (1) W. E. McEwen and R. L. Cobb, Chem. Rev., 55, 511 (1955).
- (2) F. D. Popp, Adv. Heterocyclic Chem., 9, 1 (1968).
- (3) F. D. Popp, ibid., 24, 187 (1979).
- (4) F. D. Popp, Heterocycles, 14, 1033 (1980).
- (5) F. D. Popp, ibid., 1, 165 (1973).
- (6) All new compounds exhibited satisfactory spectral properties and gave satisfactory C, H, and N analyses.
- (7) D. Bhattacharjee and F. D. Popp, J. Heterocyclic Chem., 17, 1211 (1980).
 - (8) D. Bhattacharjee and F. D. Popp, ibid., 17, 1035 (1980).