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A Novel Amination Dehydration of Nitromethane by Tris(dimethylamino)arsine

By HAROLD WEINGARTEN

(Central Research Department, Monsanto Company, St. Louis, Missouri 63166)

Tris(dimethylamino) arsine has been reported to interact with a number of dipolar molecules to give primarily exchange of substituents or insertion type reactions. However, its reaction with nitromethane was found to be quite different giving rise to both amination and dehydration.

shown to be dimethylcyanamide (80% yield; b.p. $42^{\circ}/6$ mm.; n.m.r.: τ 7.97, 7% in benzene, tetramethylsilane internal standard; infrared spectrum identical with that of an authentic sample*).

A plausible mechanism for this transformation is described below.

$$CH_{3}\cdot NO_{2} \longrightarrow CH_{2} = \overset{+}{N} \overset{OH}{O^{-}} \xrightarrow{As(NMe_{2})_{3}} \qquad CH_{2} = \overset{+}{N} \overset{OAs(NMe_{2})_{2}}{O^{-}} + HNMe_{2}$$

$$As(NMe_{2})_{3} \longrightarrow CH_{2} = \overset{+}{N} \overset{OAs(NMe_{2})_{2}}{OAs(NMe_{2})_{2}} \longrightarrow Me_{2}N - C = N + 2OAs(NMe_{2}) + 2HNMe_{2}$$

$$OAs(NMe_{2})_{2} \longrightarrow Me_{2}N - C = N + 2OAs(NMe_{2}) + 2HNMe_{2}$$

Tris(dimethylamino)arsine reacts with nitromethane exothermically evolving a large quantity of dimethylamine (identified by mass spectroscopy). A precipitate of arsenic trioxide (identified by X-ray diffraction) is formed, and an oil which was

OAsNMe₂ very likely has no discrete existence except as a unit in an -As-O-As-O- ring or chain molecule which can continue to react until the dimethylamino-groups are depleted.

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- * An authentic sample of dimethylcyanamide was generously supplied by the American Cyanamide Co.
- ¹ H. J. Vetter and H. Nöth, Z. anorg. Chem., 1964, 330, 233.