## LETTERS TO THE EDITOR

UNUSUAL CONVERSION OF 3-(N',N'-DIALKYLHYDRAZINO)1,1,2,2,-TETRACYANOCYCLOPENTANES TO
1-DIALKYLAMINO-2-DICYANOMETHYLENE-5-CYANOPYRROLIDINES

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We have established that 1-dialkylamino-2-dicyanomethylene-5-cyanopyrrolidines IIa-d are formed when 3-(N',N'-dialkylhydrazino)-1,1,2,2-tetracyanocyclopentanes 1a-d [1] are refluxed in aqueous isopropyl alcohol with a twofold molar excess of triethyl-amine for 2-5 min with subsequent dilution with water.

 $I, \ II \ \textbf{a} \ R^1 = R^2 = CH_3; \ \textbf{b} \ R^1 = C_3H_7, \ R^2 = CH_3; \textbf{c} \ R^1 = C_6H_5, \ R^2 = CH_3; \textbf{d} \ R^1 = CH_3, \ R^2 = C_2H_5$ 

Compound IIa. This compound had mp 169-170°C. IR spectrum,  $\nu$  (in mineral oil): 2228, 2220 (C=C); 1584 cm<sup>-1</sup> (C=C). The yield was 44%.

Compound IIb. This compound had mp 118-119°C. IR spectrum,  $\nu$  (in mineral oil): 2229, 2221 (C=C); 1600 cm<sup>-1</sup> (C=C). The yield was 40%.

Compound IIc. This compound had mp 165°C. IR spectrum,  $\nu$  (in mineral oil): 2230, 2220 (C=N); 1585 cm<sup>-1</sup> (C=C). The yield was 30%.

Compound IId. This compound had mp 129-130°C. IR spectrum,  $\nu$  (in mineral oil): 2227, 2216 (C=N); 1581 cm<sup>-1</sup> (C=C). The yield was 39%.

The structure of IIa was established by x-ray diffraction analysis. Single crystals of IIa were investigated with an SAD-4 diffractometer using Mo Ka emission, a graphite monochromator, and  $\omega$  scanning. The principal crystallographic data were as follows: a = 10.553(2), b = 6.980(2), c = 15.884(3) Å,  $\beta = 100.38(2)$ ; V = 1150.9 Å<sup>3</sup>, space group P<sup>21</sup>/C, Z = 4, R factor 4.3%. The structures of IIb-d were established by comparing the IR and <sup>1</sup>H and <sup>13</sup>C NMR spectra of IIa with the spectra of IIb-d.

## LITERATURE CITED

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