

CORRECTION OF STRUCTURE FOR SEVERAL SUPPOSED 2-SUBSTITUTED 1-NAPHTHALENE DERIVATIVES

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The work of Leonard and Hyson (1) indicates that the Friedel-Crafts acetylation of 2-acetamidonaphthalene (2) and 2-bromonaphthalene (2, 3) yields 8-acetyl derivatives, the 2-substituents [in contrast with the 1-substituents (4)] orienting the entering group to the unsubstituted ring. Thus it is necessary for us to correct the structures of several supposed 2-substituted 1-naphthalene derivatives.

In confirmation, we have found that the hypochlorite oxidation of the supposed 2-chloro-1-bromoacetanaphthone (2) yields an acid, m.p. 235-237° (uncorr.), undepressed on admixture with a sample of 7-chloro-1-naphthoic acid, m.p. 235-239° (uncorr.) available from the separation of the mixture of acids from the reaction of chlorobenzene with methyl furoate (5). Further, samples of the supposed α -di-*n*-butylaminomethyl-2-chloro-1-naphthalenemethanol hydrochloride, m.p. 124-124.5° [previously reported (2) 124.5-125°, 136.5-137°; two modifications], and the 7-chloro isomer, m.p. 123.5-125° [previously reported (6) 123-125°], melted at nearly identical temperatures. The mixed melting point showed no depression.

The necessary corrections involve change in structure from I to II



in the following cases (2): X = NHCOCH_3 , R = COCH_3 ; X = NH_2 , R = COCH_3 ; X = Cl, R = COCH_3 or COCH_2Br (7) or $\text{CH}(\text{OH})\text{CH}_2\text{Br}$ or $\text{CH}(\text{OH})\text{CH}_2\text{N}(\text{C}_2\text{H}_5)_2$ or $\text{CH}(\text{OH})\text{CH}_2\text{N}(\text{C}_4\text{H}_9)_2 \cdot \text{HCl}$ or $\text{CH}(\text{OH})\text{CH}_2\text{N}(\text{C}_6\text{H}_{11})_2 \cdot \text{HCl}$ or $\text{CH}(\text{OH})\text{CH}_2\text{N}(\text{C}_6\text{H}_{13})_2 \cdot \text{HCl}$; X = Br, R = COCH_2Br (7) or $\text{CH}(\text{OH})\text{CH}_2\text{Br}$ or $\text{CH}(\text{OH})\text{CH}_2\text{N}(\text{C}_4\text{H}_9)_2 \cdot \text{HCl}$.

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