A Convenient Preparation of 3-Furoic Acid

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A three-step synthesis of 3-furoic acid from diethyl furan-3,4-dicarboxylate has recently been reported¹ to be better than older methods of preparation through decarboxylation of polycarboxyfurans².³. These decarboxylations were carried out in high-boiling basic oils on acids which themselves had to be synthesised. However, during a study of 3-furan derivatives, we found that the now commercially available furan-3,4-dicarboxylic acid (Aldrich Chemical Company) could be smoothly mono-decarboxylated in high yield in the absence of any solvent. This simple method thus represents the most convenient synthesis of 3-furoic acid.

3-Furoic Acid:

Furan-3,4-dicarboxylic acid (x g; m.p. $210-214^\circ$) and copper powder (x/2 g) were mixed well and heated gently with a Bunsen flame at atmospheric pressure. Decarboxylation occurred readily and 3-furoic acid, m.p. $117-119^\circ$ (Ref. 1, m.p. $122-123^\circ$) was collected as it sublimed from the reaction mixture. Yields of at least 86% were consistently obtained from 1 to 20 g (x g) lots of the diacid.

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M.R. BOYD, T.M. HARRIS, B. J. WILSON, Synthesis 1971, 545.
H. GILMAN, R.R. BURTNER, J. Amer. Chem. Soc. 55, 2903 (1933).