

Revised Structures for Cytochalasins E and F

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Summary The structures of cytochalasins E and, by analogy, F are revised to **(3)** and **(4)** respectively.

CYTOCHALASINS E and F have been assigned structures **(1)** and **(2)** respectively.¹ We now report evidence which requires modification of their structures to **(3)** and **(4)**.

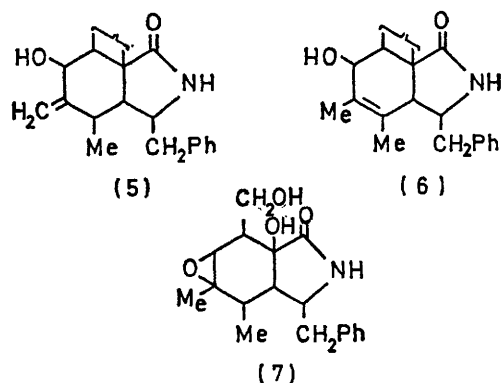
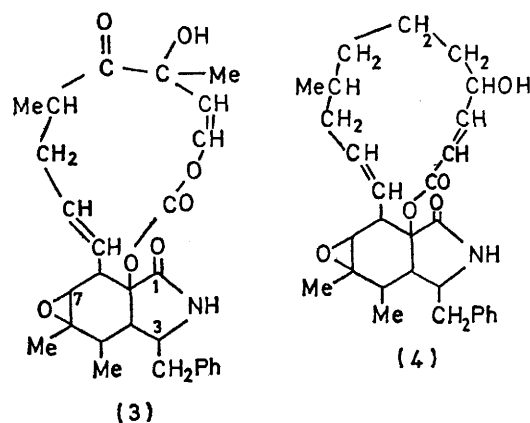
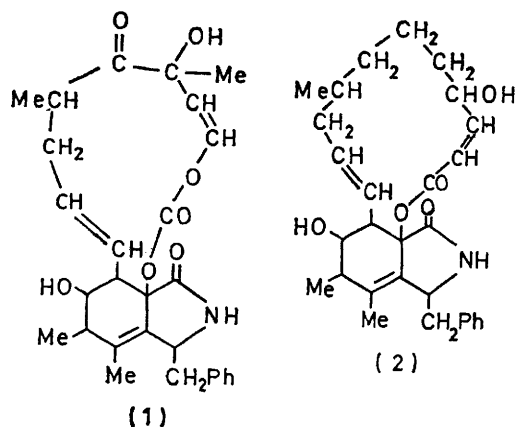
The ¹³C-n.m.r. spectrum of cytochalasin E shows signals due to only thirteen *sp*² carbon atoms in the range δ (Me₄Si) 122—213 whereas structure **(1)** possesses fifteen such atoms. On the other hand the spectra of the isomeric compounds **(5)** and **(6)** formed by mild acid treatment of cytochalasin E¹ show the expected fifteen *sp*² carbon atoms, so that a new

double bond has been formed in the acid catalysed rearrangement and cytochalasin E must possess one more ring than structure (1).

Ozonolysis of cytochalasin E and reduction of the ozonide with sodium borohydride² gave compound (7), whose structure was clear from its ¹H n.m.r. spectrum. In particular it shows the presence of only two hydroxy-protons and a doublet (*J* 5 Hz) at δ 2.68, a chemical shift characteristic of a proton on an epoxide ring.

Structure (3) for cytochalasin E readily accounts for the formation of the isomers (5) and (6), and is supported by the fact that cytochalasin E fails to form an acetate under conditions which readily acetylate the 7-hydroxy-group of cytochalasins A, B, C, and D. Structure (4) for cytochalasin F follows from the similarity of its ¹H n.m.r. spectrum to that of cytochalasin E,¹ its similar behaviour towards acid,¹ and the fact that it only forms a mono-acetate.

The earlier structures of cytochalasins E and F were based on the incorrect assignment of a signal at δ ca. 3.7 to



H-7, which gives a signal of 3.85 in cytochalasins A and B.³ This signal is in fact due to H-3, which appears at lower field in the spectra of cytochalasins E and F than in those of cytochalasins A, B, C, and D.

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¹ D. C. Aldridge, B. F. Burrows, and W. B. Turner, *J.C.S. Chem. Comm.*, 1972, 148.

² W. Rothweiler and Ch. Tamm, *Helv. Chim. Acta*, 1970, **53**, 696.

³ D. C. Aldridge, J. J. Armstrong, R. N. Speake, and W. B. Turner, *J. Chem. Soc. (C)*, 1967, 1667.