ERRATA

A. Ortega and E. Maldonado (1982) Phytochemistry 21, 785. The Publishers regret that the following structures were omitted:

A. Banerji and S. C. Pal (1982) *Phytochemistry* 21, 1321. The Publishers regret that on p. 1321, first column, lines 23–25 of the Results and Discussion were misprinted. They should read: ... study of the 270 MHz spectrum including several decoupling experiments provided the following additional information regarding the structure:

A. M. Morfaux et al. (1982) Phytochemistry 21, 1767. The authors regret that on p. 1767, second column, the sentence beginning on line 28 should read: '16-Epi-isositsirikine, 12, was identified with the most polar reduction product of geissochine according to ref. [12].'

H. Suzuki et al. (1982) Phytochemistry 21, 1824. The authors regret that on p. 1824, the absolute configurations for (+)-arctigenin (1) and (-)-arctigenin (2) should be described as 8(S),8'(S) and 8(R),8'(R), respectively, instead of 8(R),8'(S) for 1 and 8(S),8'(R) for 2 according to CIP rule.

A. D. Kinghorn et al. (1982) Phytochemistry 21, 2269. The Publishers regret that in Table 1, Templetonia egena (plant J) should have appeared containing 11,12-dehydrosparteine and sparteine, and not tetrahydroleontichine.

F. Bohlmann et al. (1982) Phytochemistry 21, 2691. The authors regret the following errors, pp. 2691 and 2696: lavendulol-2-methylbutyrate should be replaced by santolinyl 2-methylbutyrate; p. 2696: The missing ¹H NMR data of santolinyl 2-methylbutyrate (8) are as follows: 1.74 d (J = 1.5 Hz), 1.63 d (J = 1.5 Hz), 5.08 br d (J = 8 Hz)(Me₂C=CH), 5.02 dt (J = 1.7, 1.5 Hz), 5.05 dt (J = 10, 1.5 Hz), 5.80 ddd (J = 17, 10, 8 Hz), 3.67 br t (J = 8 Hz)(-CHCH=CH₂), 5.20 and 5.18 br s, 4.71 br s (H_2 C=C-CH₂O), 2.46 ddq (J = 7, 7, 7 Hz), 1.61 ddq (J = 14, 7, 7 Hz), 0.92 t (J = 7 Hz), 1.17 d (J = 7 Hz)[OCOCH(Me)Et].