Chem. Soc., **95,** 4248 (1973); A. Streitwieser, Jr. and F. Guibé, *ibid.*, **100,** 4532 (1978).

(15) R. Breslow and J. L. Grant, *J. Am. Chem. Soc.*, **99,** 7745 (1977).

(16) The "p*K*s" obtained by the electrochemical method are *relative* numbers. Although these p*K*s (in acetonitrile) are based on an arbitrary reference standard [the ion pair p*K* of triphenylmethane in cyclohexylamine (CHA) solution] a rough comparison with p*K*s in Me_2SO, which are based on an absolute standard (direct measurements in the low p*K* region), can be made because the p*K* of triphenylmethane in Me_2SO (30.6[17]) is close to its ion pair p*K* in CHA (31.5[14]) and because relative p*K*s *for acids forming delocalized anions do not differ much with medium.*[17]

(17) W. S. Matthews, J. E. Bares, J. E. Bartmess, F. G. Bordwell, F. J. Cornforth, G. E. Drucker, Z. Margolin, R. J. McCallum, G. J. McCollum, and N. R. Vanier, *J. Am. Chem. Soc.*, **97,** 7006 (1975).

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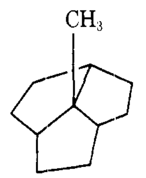
Additions and Corrections

Vol. 40, 1975

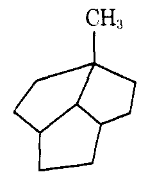
Naotake Takaishi, Yoshiaki Inamoto,* Kiyoshi Tsuchihashi, Kazuaki Yashima, and Koji Aigami: Identification of Intermediates in the Trifluoromethanesulfonic Acid Catalyzed Adamantane Rearrangement of 2,3-*endo*- and -*exo*-Tetramethylenenorbornane.

Page 2930. Chart I. Structures **10** (A_1) and **11** (A_3) should be:



10

(A_1)



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(A_3)

Page 2932. Table II. ^{13}C NMR signals should be read as:

compd notation as unknown	compd structure	^{13}C NMR signals, ppm (multiplicity, rel intensity)
A_2	4	24.8 (t, 3), 40.5 (t, 6), 60.4 (s, 0.9)
A_1	10	28.4 (q, 1), 31.8 (t, 6), 52.8 (d, 3), 62.8 (s, 0.3)
A_3	11	27.8 (q, 1), 31.7 (t, 4), 39.4 (t, 2), 45.5 (d, 2), 51.4 (s, 0.3), 61.7 (d, 1)

Slayton A. Evans* and Andrew L. Ternay, Jr.*: Preferred Conformer Assignments of Diaryl Sulfoxides Employing Aromatic Solvent Induced Shifts.

Page 2294. Experimental Section. Syntheses labeled *cis*-9-ethylthioxanthene *S*-oxide (*cis*-**5**) and *cis*-9-isopropylthioxanthene *S*-oxide (*cis*-**6**) should be relabeled trans. Syntheses labeled *trans*-9-ethylthioxanthene *S*-oxide (*trans*-**5**) and *trans*-9-isopropylthioxanthene *S*-oxide (*trans*-**6**) should be relabeled cis. Table I is correct. (We thank Professor Y. Tamura for noting this exchange.)

Werner Herz* and Ram P. Sharma: A *trans*-1,2-*cis*-4,5-Germacradienolide and Other New Germacradienolides from *Tithonia* Species.

Page 3122. Column 1, lines 24–25 should read: The CD curve of **1a** exhibits a positive Cotton effect while that of **4a** is negative, although no change has occurred . . .

Vol. 41, 1976

Courtland Symmes, Jr., and Louis D. Quin*: 1-Vinylcycloalkenes in the McCormack Cycloaddition with Phosphonous Dihalides. Stereochemistry of Some Resulting Bicyclic Phospholene Oxides.

Page 242. The ^{31}P NMR shifts on line 26 are reversed; the entry should read: δ +28.6 (**13a**, 56%) and +26.2 (**13b**, 44%).

K. Grant Taylor* and Melvin S. Clark, Jr.: Aliphatic Azoxy Compounds. 5. Functionalization of (*Z*)-Phenylmethyldiazene 1-Oxide.

Page 1145. Column 1, line 7 from bottom: "(ϵ 1900)" should be "(ϵ 11 900)".

D. C. Baker, J. Defaye, A. Gadelle, and D. Horton*: Reduction of Ketones with Incorporation of Deuterium at the α Position: Anomalous Reduction of Keto Sugar Derivatives.

Page 3836. Table I. The data for compound **9** should read: **9**, $CDCl_3{}^e$, 4.95 d (H-1), 4.79 dd (H-2), 4.15 t (H-3), 3.52 t (H-4), 3.84 sx (H-5), 4.29 q (H-6), 3.74 t (H-6′), 5.53 s (PhCH), 2.12 s (OAc), 3.39 s (OMe), 7.40 m (aryl).

Page 3837. Table II. The data for compound **9** should read: **9,** $CDCl_3{}^e$, 3.7 ($J_{1,2}$), 9.5 ($J_{2,3}$), 9.5 ($J_{3,4}$), 9.5 ($J_{4,5}$), 4.0 ($J_{5,6}$), 10 ($J_{5,6'}$), 9.5 ($J_{6,6'}$).

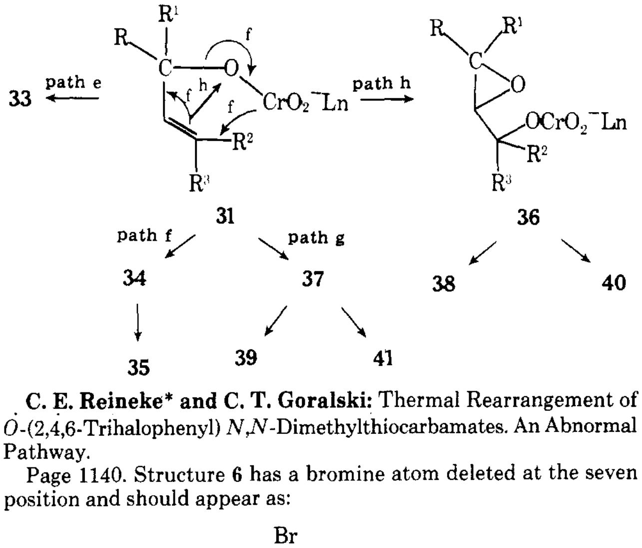
Charles A. Kingsbury,* Dan Draney, Alan Sopchik, William Rissler, and Dana Durham: Survey of Carbon-13–Hydrogen Splittings in Alkenes.

Page 3863. Reference to the elegant earlier work of Vogeli and von Philipsborn, *Org. Magn. Reson.*, **7**, 617 (1975), should have been given.

Vol. 42, 1977

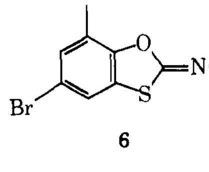
Padmanabhan Sundararaman and Werner Herz*: Oxidative Rearrangements of Tertiary and Some Secondary Allylic Alcohols with Chromium(VI) Reagents. A New Method for 1,3-Functional Group Transposition and Forming Mixed Aldol.

Page 817. The lower part of Scheme II was reproduced incorrectly. The correct formulation is as follows:



C. E. Reineke* and C. T. Goralski: Thermal Rearrangement of *O*-(2,4,6-Trihalophenyl) *N*,*N*-Dimethylthiocarbamates. An Abnormal Pathway.

Page 1140. Structure **6** has a bromine atom deleted at the seven position and should appear as:



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