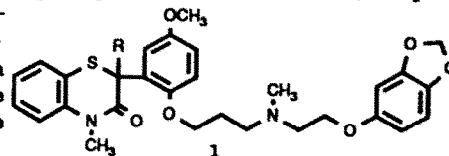


GRAPHICAL ABSTRACTS

Conformational Study of 2-Phenylbenzothiazine Part of SA2995, a Ca^{2+} Antagonist Having a Benzothiazine Skeleton, and Structure Activity Relationships

BioMed. Chem. 1994, 2, 235

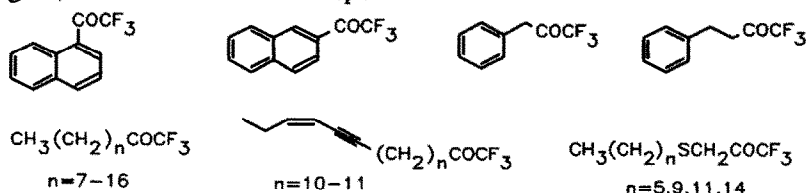
K.Fujimura*, M.Fujita, H.Suhara and Y.Kawashima, Developmental Research Lab., Santen Pharmaceutical Co., Ltd., 9-19, Shimoshinjo 3, Higashiyodogawa-ku, Osaka 533, Japan
Conformational analysis was done for the 2-phenylbenzothiazine part of 1 ($\text{R}=\text{H}, \text{OCH}_3, \text{SCH}_3, \text{CH}_3, i\text{-C}_3\text{H}_7$) by MO method. The molar fractional ratios within a particular range of 2-phenyl ring rotational angle that contained each global minimum conformation were found to correlate well with the activities.



SYNTHESIS OF TRIFLUOROMETHYL KETONES AS INHIBITORS OF ANTENNAL ESTERASES OF INSECTS

BioMed. Chem. 1994, 2, 243

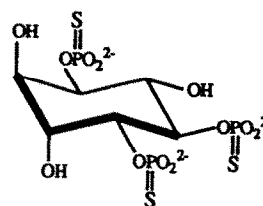
A. Parrilla, I. Villuendas and A. Guerrero*
Department of Biological Organic Chemistry, C.I.D. (CSIC)
Jordi Girona Salgado, 18-26. 08034 Barcelona. Spain



SYNTHESIS OF 1L-CHIRO-INOSITOL 2,3,5- TRISPHOSPHOROTHIOATE, THE FIRST PARTIAL AGONIST AT THE PLATELET MYO-INOSITOL 1,4,5- TRISPHOSPHATE RECEPTOR

BioMed. Chem. 1994, 2, 253

C Liu, J Al-Hafidh, J Westwick and B V L Potter*
School of Pharmacy & Pharmacology, University of Bath,
Claverton Down, Bath BA2 7AY, UK.

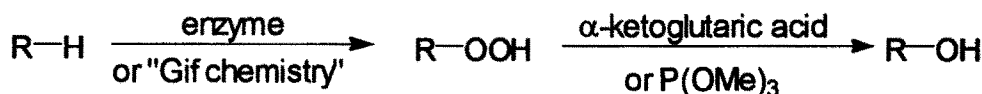


Chiro-inositol 2,3,5-trisphosphorothioate was synthesised from quebrachitol and established as a partial agonist for mobilisation of intracellular Ca^{2+} in permeabilised platelets.

The Functionalisation of Saturated Hydrocarbons. Part 30. Model Studies on the Mechanism of Some Oxygenases

BioMed. Chem. 1994, 2, 259

Derek H.R. Barton*, Christophe O. Bardin, Dario Doller.
Department of Chemistry Texas A&M University; College Station, TX 77843-3255



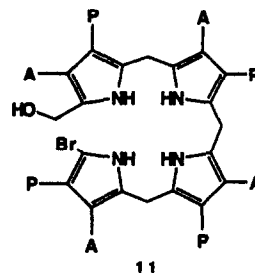
**19-BROMO-1-HYDROXYMETHYLBILANE,
A NOVEL INHIBITOR OF URO'GEN III SYNTHASE.**

Clotilde Pichon*, Barbara P. Atshaves, Richmond Danso-Danquah,
Neal J. Stolowich and A. Ian Scott*.

*Center for Biological NMR, Department of Chemistry, Texas A&M
University, College Station, Texas 77843-3255, USA.*

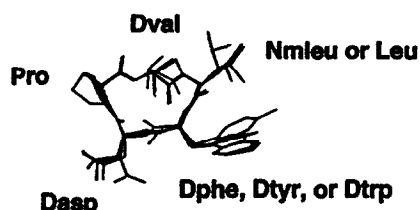
A novel hydroxymethylbilane analog, 19-Br-HMB (11),
has been synthesized. Its activity with the enzyme
Uro'gen III synthase shows competitive inhibition.

BioMed. Chem. 1994, 2, 267



**SYNTHESIS, MOLECULAR MODELLING, AND
NMR STRUCTURE DETERMINATION OF FOUR
CYCLIC PEPTIDE ANTAGONISTS OF ENDOTHELIN**

Erin K. Bradley*, Simon C. Ng,
Reyna J. Simon, & David C. Spellmeyer
Chiron Corporation, 4560 Horton Street, Emeryville, CA 94608

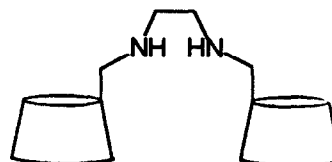


BioMed. Chem. 1994, 2, 279

**Synthesis and Transacylating Reactivity
of β -Cyclodextrin Ethylenediamines.**

John C. Beeson and Anthony W. Czarnik, Department of Chemistry, The Ohio State
University, Columbus, OH 43210, USA

The syntheses and transacylating activities of
an ethylenediamine-bridged cyclodextrin dimer
compound is reported, together with those of
reference cyclodextrinylamines.



BioMed. Chem. 1994, 2, 297