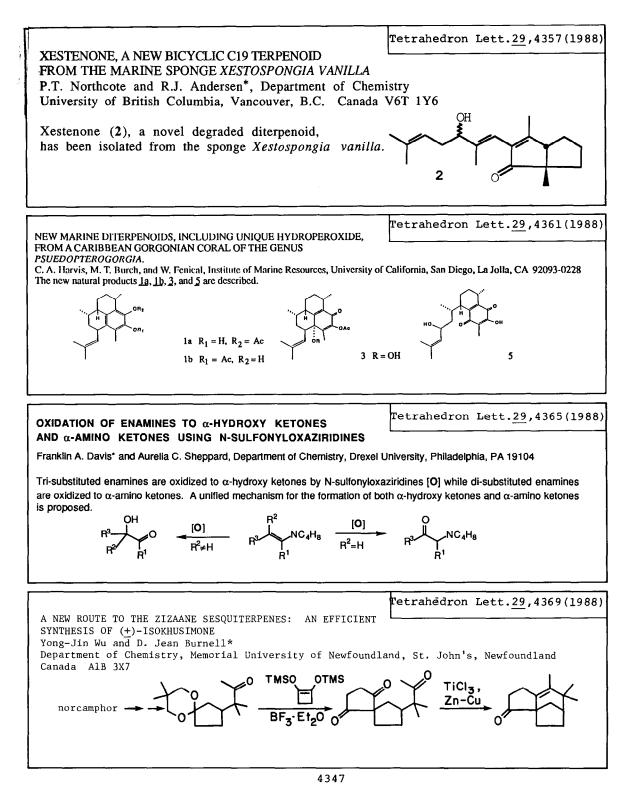
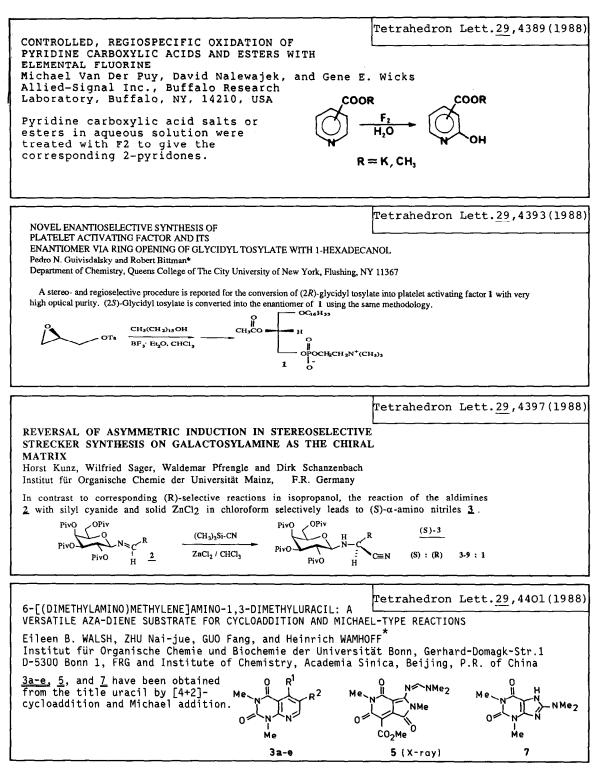
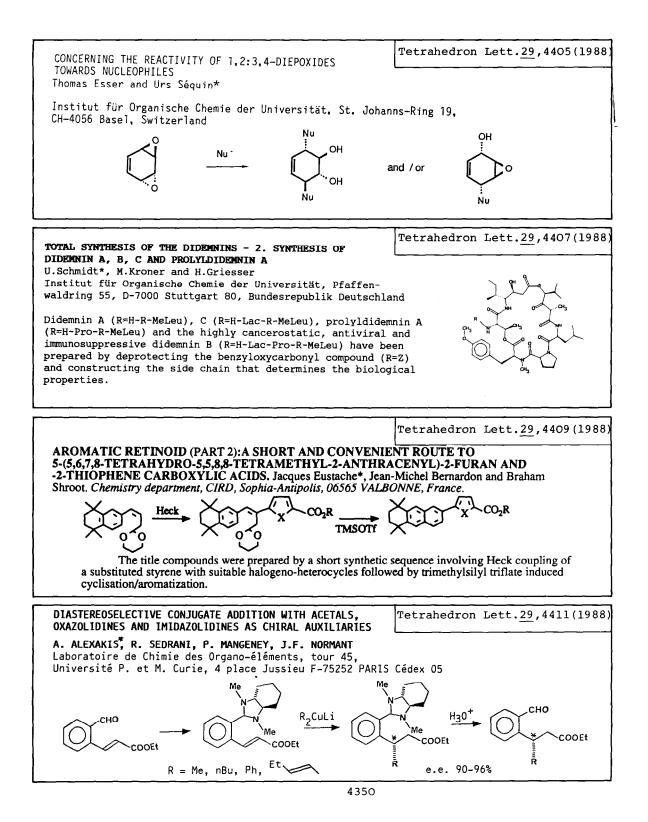
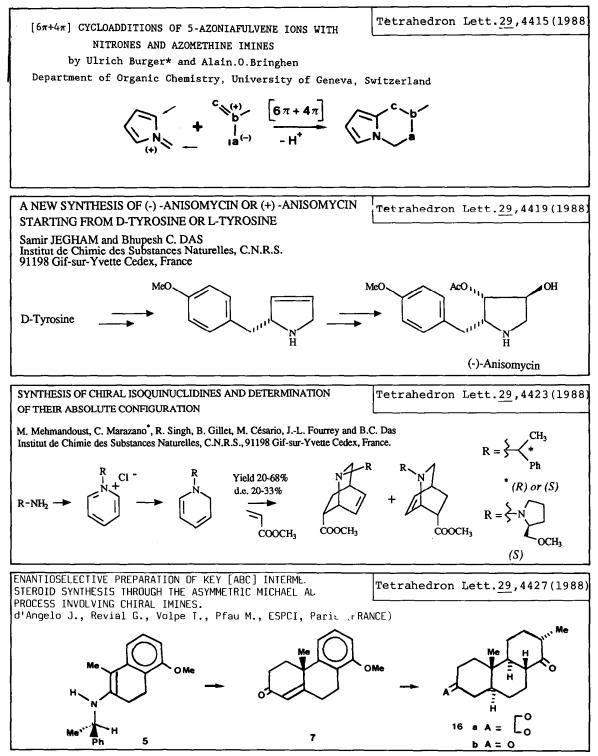
GRAPHICAL ABSTRACTS



CONFORMATIONAL ANALYSIS OF FLEXIBLE MOLECULES: Tetrahedron Lett.29,4373(1988) LOCATION OF THE GLOBAL MINIMUM ENERGY CONFORMATION BY THE SIMULATED ANNEALING METHOD Stephen R. Wilson*, Weili Cui, Jules W. Moskowitz and Kevin E. Schmidt An new approach for finding the lowest energy conformation of flexible molecules is reported. Examples are tetrahydroionone, arachidonic acid and enkephalin. COOH Tyr-Gly-Gly-Phe-Met 2 CH-1 Tetrahedron Lett.29,4377(1988) SPECTROSCOPIC PROPERTIES OF [n.1]-META-**CYCLOPHANEYLIDENES** David A. Shultz and Marye Anne Fox* Department of Chemistry, University of Texas, Austin, Texas 78712 The photophysical properties of the title compounds are discussed. These n = 4, 5, 6 compounds show large fluorescence quantum yields which are the result of relatively small torsional displacements of the phenyl rings upon relaxation in the first electronic excited state. Tetrahedron Lett.29,4381(1988) APHANORPHINE, A NOVEL TRICYCLIC ALKALOID FROM THE BLUE-GREEN ALGA APHANIZOMENON FLOS-AQUAE Nanda Gulavita, Akira Hori and Yuzuru Shimizu*, Department of Pharmacognosy and Environmental Health Sciences, The University of Rhode Island, Kingston, RI 02881, USA; Parkanyi Laszlo and Jon Clardy*, Department of Chemistry, Baker Laboratory, Cornell University, Ithaca, New York 14853-1301 USA An alkaloid, aphanorphine was isolated from blue-green alga, Aphanizomenon flos-aquae. Tetrahedron Lett.29,4385(1988) SYNTHESIS OF N-SUCCINIMIDYL-2,4-DIMETHOXY-3-(TRI-N-BUTYLSTANNYL)BENZOATE VIA REGIO-SPECIFICALLY GENERATED LITHIUM 2,4-DIMETHOXY-3-LITHIOBENZOATE Acharan S. Narula and Michael R. Zalutsky Department of Radiology, Duke University Medical Center, Durham, NC 27710 Lithium 2,4-dimethoxy-3-lithiobenzoate can be efficiently generated from 2,4-dimethoxy-3bromobenzoic acid in THF at -100°C. HOOC . OMe (O)🖕 LiOOC Ó .OMe Br 0Me OMe 4348



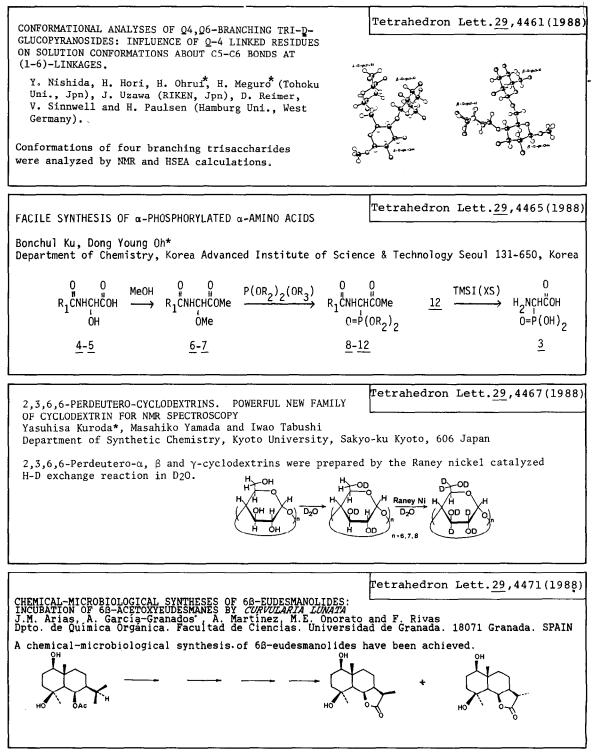




Tetrahedron Lett.29,4431(1988) A Multiple Hammett Study of the Nucleophilic Substitution Reaction of &-Carbonyl Derivatives Soo-Dong Yoh" and Oh-Seuk Lee Department of Chemistry, Kyungpook National University, Daegu 635, Korea The mechanism for the reaction of phenacyl benzenesulfonates with pyridines have been studied by means of multiple Hammett correlations. Tetrahedron Lett.29,4435(1988) AEROBIC OXIDATIVE-CARBONYLATION OF OLEFINS. A CONVENIENT PREPARATION OF 8-HYDROXYALKANOIC ACID DERIVATIVES FROM OLEFINS Hisao URATA, Atsuko FUJITA, and Takamasa FUCHIKAMI* Sagami Chemical Research Center, Nishi-Ohnuma 4-4-1, Sagamihara, Kanagawa 229, Japan $RCH=CH_2 + CO + Ac_2O + 1/2O_2 \xrightarrow{PdCl_2-CuCl_2-NaCl}{A_{COH}-80 \circ C-24 \text{ hr}} RCHCH_2COOAC$ 0Ac Tetrahedron Lett.29,4437(1988) REACTIONS OF NITROSONIUM HEXAFLUOROPHOSPHATE WITH SOME OLEFINIC COMPOUNDS IN ACETONITRILE: ONE STEP SYNTHESIS OF 4H-5,6-DIHYDRO-1,2-OXAZINES. Gae Hyung Lee, Jae Mok Lee, Won Bok Jeong, and Kyongtae Kim* Department of Chemistry, Seoul National University, Seoul 151-742, Korea $\frac{\text{NOPF}_6, \text{N}_2}{-23^\circ \text{C}, \text{CH}_3 \text{CN}}$ $X = S_0$ Y = H.MeTetrahedron Lett.29,4441(1988) SENSITIVE NATURE OF LIGAND COUPLING AND PSEUDOROTATION TO ELECTRONIC EFFECT OF SUBSTITUENT---LIGAND COUPLING IN THE REACTIONS OF BENZYLIC ARYL SULFOXIDES WITH BENZYLIC GRIGNARD REAGENTS. S. Wakabayashi, M. Ishida, T. Takeda and S. Oae, Okayama University of Science, Ridai-cho 1-1, Okayama 700, Japan In the reaction of p-benzenesulfonylphenyl benzylic sulfoxides with benzylic Grignard reagents, benzylic groups bearing an electron-withdrawing group tends to couple with the aryl group preferentially. THF, N₂ -CH₂-{O}-X + PhCH₂MgCl --X + r.t. X: CH₃ or Cl -so -Сн,

Tetrahedron Lett.29,4445(1988) LIGAND COUPLING THROUGH G-SULFURANE --- COMPLETE RETENTION OF GEOMETRIC CONFIGURATION OF ALLYLIC AND VINYLIC GROUPS IN THE REACTIONS OF ALLYLIC, AND VINYLIC SULFOXIDES WITH GRIGNARD REAGENTS. S. One^{*}, T. Takeda and S. Wakabayashi^{*}, Okayama University of Science, Ridai-cho 1-1, Okayama 700, Japan $S^{\text{THF, N}_2} \rightarrow S^{\text{CH}_2\text{-CH}_2\text{-CH}_2\text{-CH}_3} + C_2H_5MgBr \xrightarrow{\text{THF, N}_2} O SO_2 O CH_2\text{-CH}_2\text{-CH$ (cis/trans:26/74) 48.7% (cis/trans:26/74) + C_2H_5MgBr THF, N_2 r.t., 1h trans only trans-44.0% (cis-) (only cis-33.3%) Tetrahedron Lett.29,4449(1988) THERMAL REARRANGEMENT OF N-ACETYL-N-NITROSONEURAMINIC ACID DERIVATIVE: SYNTHESIS OF 3-DEOXY-D-NONULOSONIC ACID (KDN) Ryuichi Shirai, Mitsunobu Nakamura, Sumiko Hara, and Haruo Ogura* School of Pharmaceutical Sciences, Kitasato University, Shirokane, Minato-ku, Tokyo 108, Japan Thermal rearrangement of <u>1</u> followed by deprotection gave 3-deoxy-<u>D</u>-nonulosonic acd (KDN) (2). 2 HO-H NO ACO Tetrahedron Lett.29,4453(1988) ASYMMETRIC BORANE REDUCTION OF KETONES CATALYZED BY OXAZABOROLIDINE In Kwon Youn, Sang Who Lee, and Chwang Siek Pak* Korea Research Institute of Chemical Technology, Chungnam, Korea + 2 BH3 · S(CH3)2 THE Tetrahedron Lett.29,4457(1988) CONFORMATIONAL ANALYSIS OF α AND β (1-6) MANNODI-SACCHARIDES BY DEUTERIUM SUBSTITUTION EFFECT ON RELAXATION RATE AND NOE Hiroshi Hori, Yoshihiro Nishida, Hiroshi Ohrui, Hiroshi Meguro and Jun Uzawa^a Department of Food Chemistry, Tohoku University, Sendai, Japan, RIKEN(Institute of Physical and Chemical Reserch)HO-The study of H-differential relaxation rate HOand NOE using chirally deuterated sugars 1 and 2, revealed the preferred conformation of each α and OMo β (1-6) oligomannose in solution. HO 1

4353



	Tetrahedron Lett.29,4473(1988)
METAL-AMMONIA REDUCTION AND REDUCTIVE ALKYLATION OF NAPHTHALENE SULPHONAMIDES. A NEW ROUTE TO SUBSTITUTED NAPHTHALENES. L. Gottlieb and H.J.E. Loewenthal, Dept. of Chemistry, Technion-Israel Institute of Technology, Haifa, Israel. Naphthalene-1- and -2-sulphonamides can be conver= ted into dihydro compounds which can be alkylated in situ or subsequently. On heating the products rearomatise, giving specifically substituted naphthalenes.	$R \longleftrightarrow SO_2NHR' \qquad SO_2NR'R'' \qquad X$ $R \longleftrightarrow SO_2NHR' \qquad SO_2NR'R'' \qquad X$ $R \longleftrightarrow SO_2NHR' \qquad SO_2NR'R'' \qquad X$ $R \longleftrightarrow SO_2NR'R'' \qquad X$
	Tetrahedron Lett.29,4477(1988)
$\begin{array}{c} \textbf{CARBON-SULPHUR BOND FORMATION CATALYSED BY}\\ \textbf{BIS(DIPHENYLPHOSPHINO)-METHANE COMPLEXES OF PLATINPhilip C. Bulman Page*, Sukhbinder S. Klair, Michael P. Brown, MarieStephen J. Maginn, and Suzanne MulleySchool of Chemistry, University of Liverpool, P.O.Box 147, Liverpool,RSH + R'I \begin{array}{c} PtCl_2(dppm) \\ \hline Na_2CO_3 \\ \end{array} \qquad up \\ RSH + CH_2I_2 \\ \hline Na_2CO_3 \\ \end{array} \qquad up \\ \hline RSCH_2SR \\ \end{array}$	orie M. Harding,Christopher S. Smith,
	Metrohedron Lott 29 4481 (1988)
	Tetrahedron Lett. <u>29</u> ,4481(1988)
A SYNTHESIS OF THE C(1)-C(15) SEGMENT OF TSUKUBAENOLIDE (FK 506). Philip Kocienski* and Michael Stocks Chemistry Department, The University, Southampton, SO9 5NH, U. K. and David Donald*, Martin Cooper, and Anthony Manners Fisons Pharmaceuticals PLC, Bakewell Road, Loughborough, Leicestershire, LE11 0RH, U.K. A synthesis of the C(1)-C(15) segment (4) of the immunosuppressant Tsukubaenolide from Tri-O- acetyl-D-glucal and (S)-Pipecolinic acid methyl ester is described.	
	INS Tetrahedron Lett.29,4485(1988)
BIO-ORGANIC SYNTHESIS OF OPTICALLY ACTIVE CYANOHYDRINS AND ACYLOINS J. BRUSSEE*, E.C. ROOS, and A. VAN DER GEN. Department of Chemistry, Gorlaeus Laboratories, Leiden University, P.O. Box 9502, 2300 RA, Leiden, The Netherlands. Chiral acyloins of high optical purity have been obtained in good yields by enzyme catalyzed formation of optically active cyanohydrins, followed by hydroxyl protection and reaction with a Grignard reagent. $R^{1} \longrightarrow R^{1} \bigoplus C^{0H} \longrightarrow R^{1} \bigoplus C^{0H} \bigoplus R^{1} \bigoplus C^{0} \bigoplus R^{1} \bigoplus C^{0} \bigoplus$	
4355	