#### GRAPHICAL ABSTRACTS

## VARIATION AMONG KNOWN KALIHINOL AND NEW KALIHINENE DITERPENES FROM THE SPONGE ACANTHELLA CAVERNOSA.

Jaime Rodríguez, Rosa M. Nieto, Lisa M. Hunter, M. Cristina Diaz, and Phillip Crews\* Department of Chemistry and Biochemistry and Institute for Marine Sciences, University of California, Santa Cruz, CA 95064. Emil Lobkovsky and Jon Clardy\* Department of Chemistry, Cornell University, Ithaca, NY, 14853-1301.

Seven new diterpene constituents in the kalihinane family were isolated from the sponge *Acanthella cavernosa*. These diterpenes are expected to be a chemotaxonomic marker for this sponge.

Tetrahedron, 1994, 50, 11079

3 R<sub>1</sub>=R<sub>2</sub>=NC 4 R<sub>1</sub>=NHCOH, R<sub>2</sub>=NC 5 R<sub>1</sub>=NC, R<sub>2</sub>=NHCOH 7 R<sub>1</sub>=R<sub>2</sub>=NC 8 R<sub>1</sub>=NC, R<sub>2</sub>=NHCOH

5 R<sub>1</sub>=NC, R<sub>2</sub>=NHCOH : 6 R<sub>1</sub>=R<sub>2</sub>=NHCOH :

Tetrahedron, 1994, 50, 11091

9 R<sub>1</sub>=NHCOH R<sub>2</sub>=NC 10 R<sub>1</sub>=NHCOH, R<sub>2</sub>=NCS

#### Reactions of N-Chloro β-Lactams

Peter R. Guzzo and Marvin J. Miller\*

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Methanol addition to C3 of N-chloro β-lactam 4 with concomitant loss of chloride gave α-methoxy β-lactam 5 under mild alkaline conditions but was accompanied by methanolysis of the β-lactam ring as the major reaction pathway.

# SYNTHESIS OF PORPHYRINS TAILORED WITH EIGHT FACIALLY-ENCUMBERING GROUPS. AN APPROACH TO SOLID-STATE LIGHT-HARVESTING COMPLEXES

Richard W. Wagner and Jonathan S. Lindsey\*
Carnegie Mellon University, Pittsburgh, PA 15213 USA

Ilona Turowska-Tyrk and W. Robert Scheidt Notre Dame University, Notre Dame, IN 46556 USA

Routes are established to porphyrins bearing various Ar-groups. The crystal structure is determined for  $Ar = C_6F_5$ .

Tetrahedron, 1994, 50, 11097

## CONCISE, EFFICIENT NEW SYNTHESIS OF PIPERCIDE,

AN INSECTICIDAL UNSATURATED AMIDE FROM

Tetrahedron, 1994, 50, 11113

PIPER NIGRUM, AND RELATED COMPOUNDS. George M. Strunz and Heather Finlay, Canadian Forest Service, P.O.Box 4000, Fredericton, New Brunswick, Canada, E3B 5P7.

Pipercide and piperolein A, from *Piper nigrum*, were synthesized in overall yields of 21% and 35% respectively, by a short route using the Sakai aldol-Grob fragmentation sequence.

## 7-EPIZINGIBERENE, A NOVEL BISABOLANE SESQUITERPENE FROM WILD TOMATO LEAVES David C. Breeden and Robert M. Coates,

Tetrahedron, 1994, 50, 11123

Department of Chemistry, University of Illinois, 600 S. Matthews, Urbana, IL 61801, USA. The identification of 7-epizingiberene (2), as the 7R diastercomer of zingiberene (1) from oil of ginger implicates the probable occurrence of opposite sidechain rotations in their biosyntheses.

## SYNTHESIS AND ABSOLUTE CONFIGURATION OF AN ISOTACTIC NONAMETHOXY-1-PENTACOSENE FROM THE BLUE-GREEN ALGA Scytonema ocellatum

Tetrahedron, 1994, 50, 11133

Yuji Mori,\* Narumi Kawajiri, Hiroshi Furukawa, and Richard E. Moore Faculty of Pharmacy, Meijo University, Tempaku-ku, Nagoya 468, Japan Department of Chemistry, University of Hawaii, Honolulu, HI 96822, USA

The novel isotactic nonamethoxy-1-pentacosene was synthesized and the absolute configuration was established

### Synthesis of the Pentacyclic Intermediate for Dynemicin A and Unusual Formation of Spiro-oxindole Ring

Tetrahedron, 1994, 50, 11143

Takaaki Okita and Minoru Isobe\*

Laboratory of Organic Chemistry, School of Agriculture, Nagoya University, Chikusa, Nagoya 464-01, Japan

A pentacyclic compound was synthesized as an important intermediate for dynemicins via the intramolecular Heck reaction.

Aphidicolin Synthesis (II)--An Expeditious and Efficient Tetrahedron, 1994, 50, 11153

Formal Synthesis of (±)-Aphidicolin

Masahiro Toyota, Youichi Nishikawa, and Keiichiro Fukumoto\* Pharmaceutical Institute, Tohoku University, Aobayama, Sendai 980-77, Japan

An expeditious and efficient formal total synthesis of an antiviral and antitumor tetracyclic diterpene aphidicolin (5) has been achieved. An intramolecular Heck reaction (1→2) and an intramolecular Diels-Alder reaction (3->4) were utilized for the key steps of the sequence.

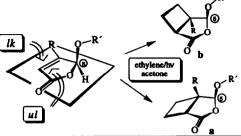
CHIRAL INDUCTION IN PHOTOCHEMICAL REACTIONS 15.<sup>1</sup>
DETECTION OF STEREOELECTRONIC EFFECTS BY TEMPERATURE
DEPENDENT MEASUREMENTS OF THE DIASTEREOSELECTIVITY
IN THE PHOTOSENSITIZED [2 + 2)-CYCLOADDITION

Tetrahedron, 1994, 50, 11167

Norbert Hoffmann, Helmut Buschmann, Gerhard Raabe and Hans-Dieter Scharf\*

Institut für Organische Chemie der RWTH Aachen Prof.-Pirlet-Str. 1, D-52056 Aachen

The mechanism of the diasteroselection in the corresponding reaction is investigated by determination of the dependence of the stereoselectivity on the temperature and on R and R'.



BIS(2,3-DIMETHYLBUTENYL)SULFANES, THE PRODUCTS OF SULFANE ADDITION TO 2,3-DIMETHYL-1,3-BUTADIENE AND OF THE REACTION OF SULFUR WITH 2,3-DIMETHYL-2-BUTENE

Tetrahedron, 1994, 50, 11187

D. Jungk, N. Schmidt and J. Hahn $^{a,*}$ , P. Versloot, J. G. Haasnoot and J. Reedijk $^b$ ; aUniversität Köln (FRG),  $^b$ Leiden University (NL)

$$H_2S_n$$
 +  $CS_2 \over RT, 1 h$  1, 2 and 3  
 $S_8$  +  $TMTD$ ,  $Z_nO$  1 and 3  
 $TMTD$  = tetramethylthiuram disulfide

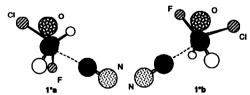
 $\searrow$   $S_n$  3

Conformational Analysis of 2-Chloro-2-fluoroecstaldehyde and Calculated Transition State Structures of Nucleophilic Addition Reactions.

Tetrahedron, 1994, 50, 11197

G. Frenking\*, K.F. Köhler, and M.T. Reetz\*\*

Fachbereich Chemie, Philippe-Universität Marburg, Hans-Meerwein-Straße, D-35032 Marburg, Germany, and \*\*Max-Planck-Institut für Kohlenforschung, Kaiser-Wilhelm-Platz 1, D-45470 Mülheim/Fluhr, Germany.



Quantum mechanical calculations of the transition states for CN° addition to 2-chloro-2-fluoroacetaldehyde predict anti-Felkin-Anh diastereoselectivity for the addition reaction. Transition state 1°a is 0.7 kcal mol<sup>-1</sup> (MP2/6-31G(d)/HF/6-31G(d) + ZPE) lower in energy than 1°b, because the conformation of the substrate is more favorable in the former transition state than in the latter. The importance of the conformation of the substrate for the diastersoeelectivity is emphasized.

SHORT INTERMOLECULAR S...S CONTACTS IN A REACTION PRODUCT FROM THE CYCLIC SULFATE ESTER OF DIMETHYL L-TARTRATE AND 2-THIOXO-1,3-DITHIOLE-4,5-DITHIOLATE.

Tetrahedron, 1994, 50, 11205

Turan Ozturk<sup>1</sup>, David C. Povey<sup>b</sup> and John D. Wallis<sup>1</sup>, \*Centre for Materials Research, University of Kent, Canterbury, CT2 7NH, b\*Chemistry Department, University of Surrey, Guildford, GU2 5XH, U.K.

Reaction of 4 and 5 give meso and dl products 8 and 9. Short S...S contacts in crystalline 9 arise from mutual polarisations and the asymmetric shape of the bonded S atom.

#### CONFORMATIONAL ANALYSIS OF 4-ARYL-2,2,6,6-TETRA-METHYLHEPTANE-3,5-DIONES. DIPOLE MOMENT DETER-MINATIONS AND MOLECULAR MECHANICS CALCULATIONS.

Tetrahedron, 1994, 50, 11213

M. Moreno-Mañas,\*,a M. Fathallah, I. Hernández-Fuentes, C. Jaime, M.E. Lloris, J. Marquet, M.F. Rey-Stolleb

<sup>a</sup> Department of Chemistry. Universitat Autònoma de Barcelona. Bellaterra. 08193-Barcelona. Spain. <sup>b</sup> Department of Physical Chemistry I. Faculty of Chemistry. Universidad Complutense. 28040-Madrid. Spain

Dipole moment determinations and MM calculations show that these diketones exist (74-94%) in pairs of enantiomeric conformations with angles between both carbonyl dipoles in the range 62-66°

# RESOLUTION OF PRENYL BROMOHYDRIN ESTERS AND DERIVATIVES:SYNTHESIS OF THE QUINOLINE ALKALOID (+)-(R)- AND (-)-(S)-LUNACRIDINE

Tetrahedron, 1994, 50, 11219

S.A. Barr, a D.R. Boyd, a N.D. Sharma, b T.A. Evans, J.F. Malonca and V.D. Mehta a School of Chemistry, The Queen's University of Belfast, Belfast, BT9 5AG, U.K. b Department of Chemistry, University of Delhi, Delhi 110007, India.

Chromatographic separation of the bromo MTPA ester derivatives of prenylated compounds provides a new route to enantiopure prenyl epoxides and derivatives e.g. (+)-(R)- and (-)-(S)-lunacridine and 8-methoxyplatydesmine.

## Cs<sub>2</sub>CO<sub>3</sub> OR CaO AS PROMOTERS OF

Tetrahedron, 1994, 50, 11235

ETHYL N-{[(4-METHYLPHENYL)SULPHONYL]OXY}CARBAMATE IN AMINATION REACTIONS

Marco Barani, Stefania Fioravanti, Lucio Pellacani, and Paolo A. Tardella Dipartimento di Chimica, Università "La Sapienza", P.le Aldo Moro 2, I-00185 Roma, Italy

A convenient functionalisation of benzene and substituted alkenes of different reactivity was achieved by the action of heterogeneous inorganic bases on the title reagent.

$$\begin{array}{c|c} Et_3N & \hline \\ \hline CH_2Cl_2, r.t. & R_2C=CR_2 + TsONHCQEt \\ \hline \hline CH_2Cl_2, r.t. & CQEt \\ \hline \end{array}$$

# AN EASY ACCESS TO 2-OXOHYDRAZONES VIA ELECTROPHILIC α-p-TOLYLHYDRAZONYLATION OF KETONE ENOLATES WITH tert-BUTYL p-TOLYLAZO SULFIDE

Tetrahedron, 1994, 50, 11239

Carlo Dell'Erba, Marino Novi, Giovanni Petrillo\* and Cinzia Tavani

Istituto di Chimica Organica dell'Università, C.N.R. Centro di Studio per la Chimica dei Composti Cicloalifatici ed Aromatici, Corso Europa 26, 16132 Italy

 $\alpha$ -(p-Tolylhydrazono)ketones 3 or their N-methylderivatives 7 are conveniently prepared through the title reaction from ketones 5.

Polyhydroxyl Oligothiophenes. I. Regioselective Synthesis of 3,4'- and 3,3'-dl(2-hydroxyethyl)-2,2'-bithiophene via Palladium Catalyzed Coupling of Thienylstannanes with Thienylbromides.

Tetrahedron, 1994, 50, 11249

Giovanna Barbarella\* and Massimo Zambianchi

I.Co.C.E.A., Area Ricerca C.N.R., Via Gobetti 101, 40129 Bologna, Italy

## SYNTHESIS OF 7-SUBSTITUTED DEHYDRONORAPORPHINES, WITH SOME BIOGENETIC CONSIDERATIONS

N. Atanes, S. Pérez, E. Guitián\*, L. Castedo, J.M. Saá. Universidad de Santiago, Santiago de Compostela and Universidad de les Illes Balears, Palma de Mallorca. SPAIN

N-protected 7-methyl-6a,7-dehydronoraporphines 2 were synthesized by the IBC approach. Oxidation of these compounds by oxygen led to 7-hydroxy-7-methyl-6,6a-dehydronoraporphines 13 in what may be a biomimetic process.

## Tetrahedron, 1994, 50, 11257

Tetrahedron, 1994, 50, 11267

Tetrahedron, 1994, 50, 11289

$$\begin{array}{c}
R_1O \\
R_2O \\
R_3 \\
OH
\end{array}$$

$$\begin{array}{c}
R_1O \\
CH_3
\end{array}$$

$$\begin{array}{c}
R_2O \\
R_3 \\
CH_3
\end{array}$$

$$\begin{array}{c}
CH_3 \\
3
\end{array}$$

### ALKYL RADICAL CYCLISATIONS OF

### METHYLENECYCLOPROPANE DERIVATIVES

Christine Destabel, a Jeremy D Kilburna and John Knightb

- a Department of Chemistry, University of Southampton, Southampton, SO9 5NH, UK
- b Glaxo Research and Development Limited, Park Road, Ware, Hertfordshire, SG12 0DP, UK

(Methylenecyclopropyl)propyl radicals underwent selective

5-exo cyclisation, followed by opening of the resulting

cyclopropylmethyl radical to give methylenecyclohexanes.

(Methylenecyclopropyl)butyl radicals gave a mixture of products.

## MALONATE RADICAL CYCLISATIONS OF METHYLENECYCLOPROPANE DERIVATIVES

Christine Destabel, a Jeremy D Kilburna and John Knight

a Department of Chemistry, University of Southampton, Southampton, SO9 5NH, UK

b Glaxo Research and Development Limited, Park Road, Ware, Hertfordshire, SG12 ODP, UK

Cyclisations of methylenecyclopropyl substituted malonate radicals have been investigated. Highly regioselective 5-exo, 7-endo and 8-endo cyclisations are reported.

(Bu<sub>3</sub>Sn)<sub>2</sub>, hv,  
toluene Me<sub>3</sub>Si 
$$n = 1$$
  $CO_2Et$  or  $CO_2ET$  or

ix

#### TOTAL SYNTHESIS OF MICROGININ, AN ANGIOTENSIN-CONVERTING ENZYME INHIBITORY PENTAPEPTIDE FROM THE BLUE-GREEN ALGA MICROCYSTIS AERUGINOSA

Tetrahedron, 1994, 50, 11303

Fumiyoshi Matsuura, Yasumasa Hamada, and Takayuki Shioiri\*

Faculty of Pharmaceutical Sciences, Nagoya City University, Tanabe-dori, Mizuho-ku, Nagoya 467, JAPAN

microginins	C-2	C-3
proposed structure 1	R	-
1a 1b	R R	R S
1c	s	s
revised structure 1d	s	R

An Enantioselective Total Synthesis of (+)-Altholactone from Diethyl L-Tartrate Peter Somfai, Organic Chemistry 2, Chemical Center, Lund Institute of Technology,

Tetrahedron, 1994, 50, 11315

University of Lund, P. O. B. 124, S-221 00 Lund, Sweden

An enantioselective total synthesis of (+)-altholactone (1) from diethyl L-tarrate via the L-threitol derivative 2 is described.

### HETEROGENEOUS CATALYTIC ASYMMETRIC DIHYDROXYLATION OF OLEFINS: A NEW POLYMERIC SUPPORT AND

Tetrahedron, 1994, 50, 11321

A PROCESS IMPROVEMENT

Dario Pini, Antonella Petri , Piero Salvadori\*

Centro CNR Macromolecole Stereordinate ed Otticamente Attive, Dipartimento di Chimica e Chimica Industriale, Università di Pisa, Via Risorgimento 35, 56126 Pisa, ITALY

OsO4 cat. secondary oxidant polymer bound quinine derivative

A new polymeric support containing a derivative of 4-chlorobenzoylquinine was found to give good enantioselectivity (up to 95%) in the catalytic asymmetric dihydroxylation of olefins.