

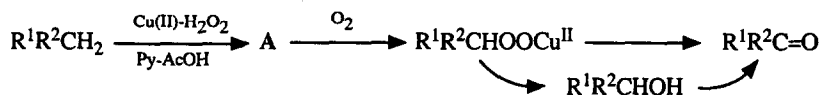
## GRAPHICAL ABSTRACTS

*Tetrahedron*, 1992, 48, 2895

### THE FUNCTIONALISATION OF SATURATED HYDROCARBONS. PART XXI. THE Fe(III)-CATALYZED AND THE Cu(II)-CATALYZED OXIDATION OF SATURATED HYDROCARBONS BY HYDROGEN PEROXIDE: A COMPARATIVE STUDY

Barton, D.H.R.\*; Beviere, S.D.; Chavasiri, W.; Csuhai, E. and Doller, D.\*

Department of Chemistry, Texas A&M University, College Station, Texas 77843-3255, USA.



Mechanistic studies show that the ketonisation of saturated hydrocarbons by Cu(II)-H<sub>2</sub>O<sub>2</sub> or Cu<sup>0</sup>/O<sub>2</sub> in pyridine-acetic acid follows the pathway *alkane* → *intermediate A* → *alkyl hydroperoxide* → *ketone and alcohol*.

*Tetrahedron*, 1992, 48, 2911

### N-HYDROXYAMINOMETHYLENEPHOSPHONATES:

#### THE REACTION OF HYDROXYLAMINES WITH ALDEHYDES AND SECONDARY PHOSPHITES

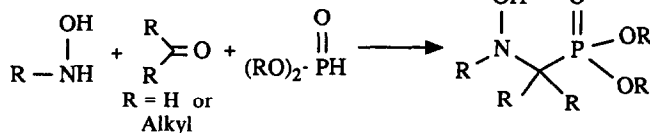
Stephen D. Pastor,\* Ramanathan Ravichandran and Roger Meuwly

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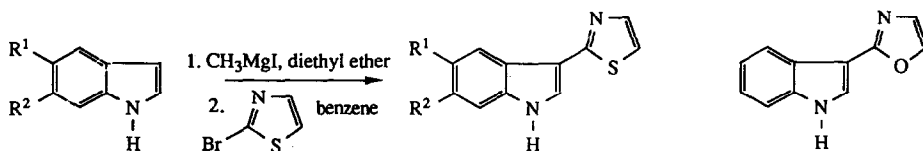
*Tetrahedron*, 1992, 48, 2919

### SYNTHESIS OF CAMALEXIN AND RELATED PHYTOALEXINS

William A. Ayer\*, Peter A. Craw, Yu-ting Ma, and Shichang Miao

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A number of 3-(2'-thiazolyl)indoles and 3-(2'-oxazolyl)indole have been prepared and the antifungal activity of these 3-alkylindoles has been examined.



*Tetrahedron*, 1992, 48, 2925

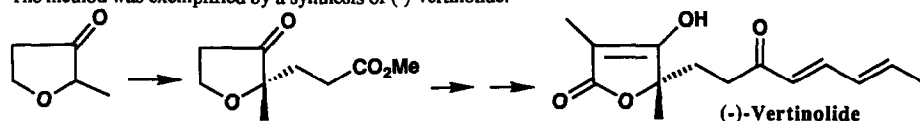
### New Approach to Chiral 5,5-Disubstituted Tetronic Acids.

#### Enantioselective Synthesis of (-)-Vertinolide

Didier Desmaële

Laboratoire de chimie Organique, Faculté de Pharmacie, 5, rue J.B. Clément, 92296 Châtenay-Malabry Cedex (France).

2,2-Disubstituted-4,5-dihydrofuranones obtained by an asymmetric Michael addition were converted into chiral tetronic acids. The method was exemplified by a synthesis of (-)-vertinolide.



*Tetrahedron*, 1992, 48, 2935

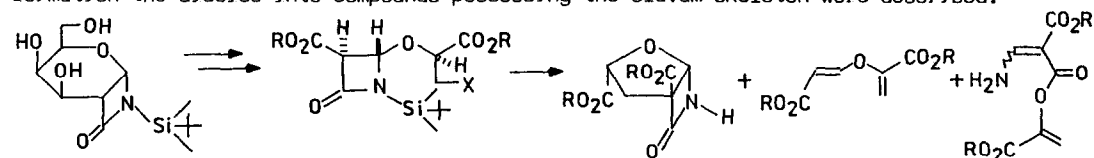
### AN ATTEMPT ON THE SYNTHESIS OF THE CLAVAM SKELETON FROM GLYCALS AND ISOCYANATES

M. Chmielewski\*, J. Grodner, Wang Fudong and Z. Urbańczyk-Lipkowska

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N-Protected 2-C:1-N-carbonyl-2-deoxy-glycopyranosylamines were subjected to the two-step oxidation to afford dicarboxylic acids. Unsuccessful attempts on transformation the diacids into compounds possessing the clavam skeleton were described.

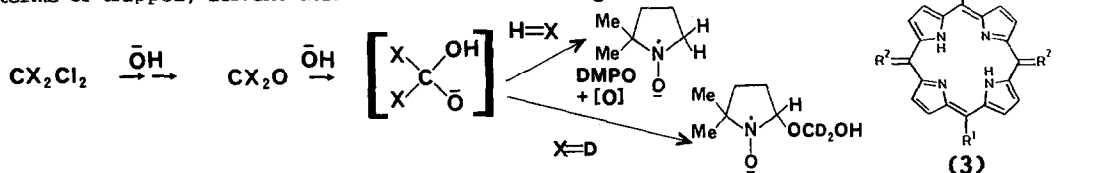


*Tetrahedron*, 1992, 48, 2951

### FACILE AERIAL OXIDATION OF A PORPHYRIN. PART 12. SPIN-TRAPPING OF SOLVENT-DERIVED INTERMEDIATES FROM THE ADDITION OF BASE TO AN OXIDISED PORPHYRIN.

L.R. Milgrom\* and W.D. Flitter<sup>b</sup>; \*Molecular Probes Unit, Department of Chemistry, and <sup>b</sup> Department of Biochemistry, Brunel University, Uxbridge, Middlesex, UB8 3PH., U.K.

Esr-active adducts are produced on addition of base and a spin-trap to dichloromethane solutions of the oxidised porphyrin (3): an explanation is offered in terms of trapped, solvent-derived intermediates being oxidised to radical species.



**INTRAMOLECULAR Pd<sup>0</sup>-CATALYSED CROSS COUPLING;  
A DIRECT ROUTE TO  $\gamma$ -OXO- $\alpha,\beta$ -UNSATURATED MACROCYCLES.**

Jack E. Baldwin, Robert M. Adlington and Steve H. Ramcharitar.  
*The Dyson Perrins Laboratory, University of Oxford, South Parks Road, Oxford, OX1 3QY.*

Intramolecular Pd<sup>0</sup>-catalysed cross coupling of acid chlorides and *Z*- and *E*- $\beta$ -stannylalkenoates has provided a new and an efficient route to 10-20 membered  $\gamma$ -oxo- $\alpha,\beta$ -unsaturated macrolides. The method has been used to synthesise the macrocyclic framework of the antibiotic A26771B and norpatulolide B.

