# JOURNAL OF THE CHEMICAL SOCIETY

# **Perkin Transactions 1**

Organic and Bio-organic Chemistry

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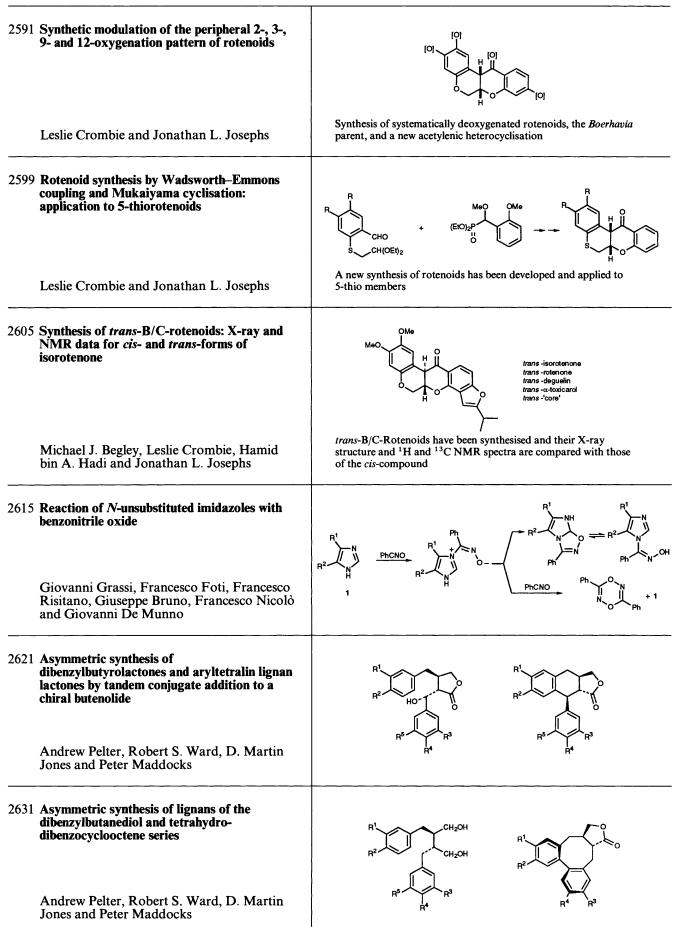
## **Perkin Communications**

	Syntheses of conduritol D derivatives from promatic compounds	$\begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \\ \end{array} \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array} \\ \end{array} \\ \end{array} \\ \end{array} \\$
H D	Ioward A. J. Carless, Kofi Busia, Yvonne Dove and Shahnaz S. Malik	X = H, Me, Cl, Br HO OH
re	<b>Jnusual chemo- and stereo-selectivities in the eactions of 1,2-dichlorocyclopropenes with itrile oxides</b>	$\begin{array}{c c} Me \\ \hline \\ Ci \\ Ci \\ Ci \\ \hline \\ Ci \\ Ci \\ \hline \\ Ci \\ \hline \\ Ci \\ \hline \\ \\ Ci \\ \hline \\ \\ Ci \\ \hline \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ $
D	Mark S. Baird, Xiaoming Li, Juma'a R. Al Dulayymi, Alexander I. Kurdjukov and /alery A. Pavlov	1a X = H b X = Cl Compound 1a reacts with a number of nitrile oxides to give 3, whereas 1b gives 6
et	Unprecedented route to enolates from silyl enol thers and enol acetates: reaction with hard and soft electrophiles	$ \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\$
	Pierre Duhamel, Dominique Cahard and ean-Marie Poirier	With a silyl dienol ether a catalytic amount of potassium <i>tert</i> - butoxide can be used

## Articles

	A REAL PROPERTY AND A REAL
<ul> <li>2513 Investigation of the potential of molybdenum(VI) hydrazido(2—) complexes as sources of nitrenium ions: cleavage of the N–N bond and incorporation of the β-nitrogen group into solvent molecules</li> <li>Marc M. Baum and Edward H. Smith</li> </ul>	$Ph(R)N - C - CHCl_2 \xrightarrow{Cl_2CHCHCl_2}_{hv \text{ or heat}} N^{-N(R)Ph}_{MoL_n} \xrightarrow{AgNO_3}_{R'OH} \xrightarrow{VNR}_{OR'} NO_2$
2521 Diamide analogues of phosphatidyl choline as potential anti-AIDS agents	$\begin{array}{c} CH_2NHCO(CH_2)_n Me \\ I \\ CHNHCO(CH_2)_n Me \\ CHNHCO(CH_2)_n Me \\ I \\ CH_2OP(C)OCH_2CH_2NMe_3 \\ I \\ CH_2OP(C)OCH_2CH_2NMe_3 \\ I \\ O \end{array}$
	Analogues of phosphatidyl choline, 5 and 6, containing amide
	groups instead of the usual ester functionality have been prepared in racemic form and 5 was found to possess moderate anti-HIV
Chunhua Jia and Alan H. Haines	activity
2525 Biosynthetic relationships in the desertomycin family Marion Mayer and Ralf Thiericke	A complete picture of the biosynthetic relationships of the desertomycin family, and the structures of oasomycin E and F are presented
2533 Synthesis of podophyllum lignans <i>via</i> an isolable <i>o</i> -quinonoid pyrone	$O = OBz$ $Ar = 3.4.5-trimethoxyphenyl; E = CO_2Me$ $OBz$
David W. Jones and Adrian M. Thompson	The adduct is converted into lignans a key step being direct lactonisation of, for example, methyl podophyllate to podophyllotoxin with ZnCl <sub>2</sub> /molecular sieves in THF
2541 Synthesis of (±)-4-deoxypodophyllotoxin, (±)-podophyllotoxin and (±)- epipodophyllotoxin	$ \begin{array}{c}                                     $
Devid W7 Tener and Advise M6 These	Podophyllum lignans
David W. Jones and Adrian M. Thompson	1

2549 Synthesis of 3-octyl-, 3-cyclohexylmet 3-carboxyl-3-(prop-2-ynyl)pyrrolidine diones required as potential aromatase inhibitors	<b>→2,5-</b>
Lawrence W. L. Woo, H. John Smith J. Barrell and Paul J. Nicholls	h, Kevin Synthesis of 3-substituted-3-(prop-2-ynyl)pyrrolidine-2,5-diones required as potential aromatase inhibitors
2555 Purines, pyrimidines and imidazoles. I Some N-substituted o-(2-hydroxyethy purines, -pyrimidines and -imidazoles aromatic acyclonucleoside analogues	<b>/I)benzyl-</b>
Demetrios C. Agathocleous and Gor Shaw	rdon Synthesis of imidazole, pyrimidine and purine nucleoside analogues
2561 Synthesis of the naturally occurring indolequinone BE 10988, an inhibitor topoisomerase II	of $H_2N \xrightarrow{V} N_{Me}$
Christopher J. Moody and Elizabeth	A total synthesis of the topoisomerase II inhibitor BE 10988 is described
2567 The synthesis of 5-ylidenepyrrol-2(5H from maleimides and pyrrol-2(5H)-on	$ \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \\ \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \\ \\ \end{array} $
G. Bryon Gill, Gwyn D. James, Kare Oates and Gerald Pattenden	en V. 5 6 7
2581 Total synthesis of pukeleimide A, a 5- ylidenepyrrol-2(5H)-one from blue gre	een algae
Gwyn D. James, Stuart D. Mills and Pattenden	Gerald
2585 Synthesis of deoxydinucleoside phospl containing 6-thio-substituted purine nucleobases	HN HO HO $R^3$ N N N N N N N N N N
Pascale Clivio, Jean-Louis Fourrey a Alain Favre	



2639	Convenient synthesis of alcohol <i>O</i> -hemiesters using isopropenyl esters as acylating reagents: synthesis of hydrophilic oxaunomycin 10- <i>O</i> - hemiester derivatives	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
	Yasuyuki Kita, Hiroshi Maeda, Fumie Takahashi and Seiji Fukui	Reaction of β-rhodomycinone derivative 8 with isopropenyl esters 4 in the presence of a catalytic amount of acid followed by selective deprotection of the terminal ester gave hydrophilic oxaunomycin 10-O-hemiester derivatives 14a, b and 19a-c
2651	Bicycloannulation of $\alpha$ -bromo $\alpha,\beta$ -unsaturated esters; synthesis of the tricyclo[4.4.0.0 <sup>1.5</sup> ]-decane framework and its congeners	$ \begin{array}{c} & & \\ & & $
	Hisahiro Hagiwara, Futoshi Abe and Hisashi Uda	CO <sub>2</sub> Me
2657	Synthesis of 5-(acylethynyl)uracils and their corresponding 2'-deoxyribonucleosides through palladium-catalysed reactions	$C \equiv C - C - R'$ $HN$ $C \equiv C - C - R'$ $HN$ $I = H, R' = Ar$ $H = 1 = 2'-deoxy-D-erythro-pentofurancse, R' = Ar$
	Nitya G. Kundu and Swapan K. Dasgupta	The synthesis of compounds I and II, through palladium-catalysed reactions, are reported

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