### JOURNAL OF THE CHEMICAL SOCIETY

# **Perkin Transactions 1**

Organic and Bio-organic Chemistry

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#### **Perkin Communications**

1091 Application of an intramolecular Michaelis—Arbusov reaction to the synthesis of nucleoside 3'-S,5'-O-cyclic phosphorothiolate triesters  Xiang Li and Richard Cosstick	MMTO SH MeO O S
1093 Enzymatic resolution of oxalate esters of a tertiary alcohol using porcine pancreatic lipase  Ian Brackenridge, Raymond McCague, Stanley M. Roberts and Nicholas J. Turner	(±)-6 (30)% ee (88)% ee
1095 Total synthesis of (+)-kifunensine, a potent glycosidase inhibitor  Jacques Rouden and Tomas Hudlicky	From chlorobenzene using Pseudomonas putida 39D (+)-Kifunensine

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## Articles

1099 Investigation of the regioselectivity of some esterification reactions involving methyl 4,6-O-benzylidene D-pyranosides and Pseudomonas fluorescens lipase	Ph $\longrightarrow$ O $\longrightarrow$ OMe ( $\alpha$ or $\beta$ ) $\beta \cdot OMe \longrightarrow HO \qquad OH \longrightarrow \alpha \cdot OMe$
Gilles Iacazio and Stanley M. Roberts	The regioselectivity of lipase-catalysed acetylations is dependent on the orientation of the anomeric substituent. The positions for the formation of esters from $\alpha$ - and $\beta$ -anomers of the title compounds are indicated
103 Studies on quinones. Part 27. Diels-Alder reaction of 8,8-dimethylnaphthalene-1,4,5(8H)-trione	
Jaime A. Valderrama, Ramiro Araya- Maturana and Fernandio Zuloaga	The title compound I reacts with 1,3-dienes to afford adducts of type II. The regiochemistry of the cycloadditions was established by chemical correlation and/or FMO theory
109 Preparation of a novel potent inhibitor of methylthioadenosine nucleosidase	NH <sub>2</sub>
David Hendry, Edward J. Hutchinson, Stanley M. Roberts, Steven M. Dunn and John A. Bryant	The purine derivative 1 is a potent inhibitor of methylthioadenosine nucleosidase
113 Direct conversion of alcohols into thiols	
	$R^1R^2R^3COH \xrightarrow{LR} R^1R^2R^3CSH$
Takehiko Nishio	Treatment of alcohols with Lawesson's reagent (LR) affords the corresponding thiols accompanied by dehydration products, alkenes
119 Reactivity of 1,1-disubstituted indazol-3-ylio oxides: synthesis of some substituted indazolols and indazolinones	Z = NO <sub>2</sub> , H  A R <sup>1</sup> , R <sup>2</sup> = alkyl groups
Vicente J. Arán, Juan L. Asensio, José R. Ruiz and Manfred Stud	Some reactions of the represented indazol-3-ylio oxides, leading to differently substituted indazoles and/or benzohydrazides, have been studied

ph dii	ormation of chiral β-lactams by otocyclisation of achiral <i>N,N</i> -sopropylarylglyoxylamides in their chiral ystalline form	Ar—C CHMe2  N— Solid state  OPTICAL PLANT OF Me2  CHMe2  Chiral crystal  Optically active β-lactarn derivatives
Fu	ımio Toda and Hisakazu Miyamoto	
ary	elluroxide elimination by oxidation of alkyl yl tellurides: remarkable effect of added ethylamine	RCHCH <sub>3</sub> [O] RCH=CHCH <sub>3</sub> + RCH=CH <sub>2</sub> Ar: Ph.
Yo Ko	oshiaki Nishibayashi, Naoki Komatsu, ouichi Ohe and Sakae Uemura	Treatment of various alkyl phenyl tellurides with 1-2 mol equiv. of m-chloroperbenzoic acid (MCPBA) in diethyl ether in the presence of triethylamine (Et <sub>3</sub> N) at 25 °C for 2 h produced the corresponding alkenes in fair to good yields and highly selectively
rac	ziridines. Part 60. Electron transfer from dical anions to N-alkanoylaziridines. cocyclic cleavage of an aziridino ketyl	
Pe an	en-Yuan Lin, Jürgen Werry, Gunther Bentz d Helmut Stamm	R <sup>-</sup> (R = CMePh <sub>2</sub> ) products
1147 Pr 1,3	reparation of all- <i>trans-</i> (1,4-phenylenehexa- 3,5-trienylene) oligomers	H-{\}
Yo	oriko Sonoda and Yukimichi Nakao	n = 2,3 all-trans-(1,4-Phenylenehexa-1,3,5-trienylene) oligomers PHT2 ( $n = 2$ ) and PHT3 ( $n = 3$ ) have been prepared by a Wittig reaction
Pa ph	nenolic constituents of <i>Glycyrrhiza</i> species.  art 10. Glyasperin E, a new 3- enoxychromen-4-one derivative from the ots of <i>Glycyrrhiza aspera</i>	Me O O O O O O O O O O O O O O O O O O O
	a Zeng, Toshio Fukai, Taro Nomura, Ru-Yi nang and Zhi-Cen Lou	The structure of glyasperin E was established with spectroscopic and synthetic methods
	ovel 10β-aziridinyl steroids; inhibitors of omatase	17; 19R X = 0 16; 19S, X = 0 19; 19R X = HOH 20; 19S, X = HOH
Vi an	ncent C. O. Njar, Elam Safi, J. V. Silverton d Cecil H. Robinson	Synthesis, structure and aromatase inhibitory properties of four novel 10β-aziridinyl steroids 17–20

	Synthesis and first X-ray structure analysis of a stabilized chiral chlorobismuthine. Fixation of molecular geometry induced by the intramolecular coordination of a sulfonyl group	Bi
	Hitomi Suzuki, Toshihiro Murafuji and Nagao Azuma	[2-(tert-Butylsulfonyl)phenyl]chloro(4-methylphenyl)bismuthine
	Preparation and reaction of difluorinated malonaldehydic acid derivatives: a new route to functionalized α,α-difluorinated esters and amides	$(X = OEt, NEt_2) \xrightarrow{Zh, (EIO)_2 SO_2} EtO \xrightarrow{F} X$ $Me_2N O$ $H^{\bullet} EIOH$
	Takashi Tsukamoto and Tomoya Kitazume	Nu FF X Nu EXO FF X
	Synthesis of very long chain fatty acid methyl esters	
		CH <sub>3</sub> (CH <sub>2</sub> ) <sub>n</sub> (HC=CHCH <sub>2</sub> ) <sub>m</sub> (CH <sub>2</sub> ) <sub>p</sub> CO₂Me  cis 1
	Marcel R. Kling, Christopher J. Easton and Alf Poulos	Esters 1 were obtained from copper-mediated coupling reactions of $\omega$ -iodo esters with Grignard reagents and from reactions of $\omega$ -oxo esters with phosphoranes
1191	Estrogen biosynthesis: 2β-hydroxy-19- oxoandrost-4-ene-3,17-dione revisited	D D D D D D D D D D D D D D D D D D D
	Eliahu Caspi, H. Ranjith, W. Dharmaratne, Esther Roitman and Cedric Shackleton	Incubation of 1 in the presence and in the absence of aromatase gave estrone-D₂ and HC●OH; mechanistic implications of the results are reviewed
	Intramolecular Diels-Alder reaction with furans: effect of the substitution pattern reinvestigated	CH=N-R - NC R
	Dipak Prajapati, Dipak R. Borthakur and Jagir S. Sandhu	IMDA reaction of α-cyanofurfurylamine and N-furfurylarylamines affords 4 or 10 in good yields

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