# Subject Index 1985

### Errata

- Bellamy F.D. and Ou K., 25,839-842(1984): corrected procedure for reduction of p-nitrobenzoic
- Berkowitz W.F., Perumattam J., and Amarasekara A., 26, 3665(1985): acknowledgement of similar work published in Chem. Letters, 4562
- Brinkman K. and Helquist P., 26, 2845(1985): formula, last product in Table on p.2846 corrected and reference 8a replaced, 4154
- Danikiewicz W. and Makosza M., 26, 3599(1985): scheme 1 which was omitted from p 3599, 6523
- Davies J.W., Malpas J.R. and Moss R.E., 26, 4533(1985): last word of second paragraph on p.4533 diazabicycle; on p.4535 energy values corrected; revised Table
- 1 produced, 5226 Deshmukh M., Dunach E. and Kagan H.B., 25, 3467(1984): Line 15 after Abstract - correction, 402
- Gamboni R., Mohr P., Waespe-Sarcevic N. and Tamm C., 26,203(1985): structure 1d is
- corrected, 3058 Guo-qiang L., Hai-jian X., Bi-chi Guong-zhong G. and Weishan Z., 26, 1233(1985): reaction conditions for Schemes 1 and 2 corrected and arrangement of compds in Table 1 changed, 1690
- Jackson A.H., Rao K.R.N., Ooi N.S. and Adelakun E., 25, 6049(1984):
- p.6049 three corrections, 3522 Kondo N., Imai K., Isobe M., Coto T., Murasugi A., Wada C. Nakagawa C. and Harashi Y., 25, 3869(1984): corrected structures
- 3009(1904): corrected structures for formulae 1 and 2, 690 Mak C-P. and Fliri H., 26, 1433(1985): on p 1434 entry 8 corrected, 4010 Newcomb M., Williams W.G. and Crumpacker E.L., 26, 1183(1985): yield of isolated bromide 1 corrected, 2727
- Otera J., Yoshinaga Y., Hirakawa K. and Nakata T., 26,3219(1985): entries 6 and 7 in Table 1 should be deleted, 3746 Overman L.E. and Burk R.M.,

- line from bottom of p.5737, 4154 Shirahama H., Arora G.S., Osawa E.and Matsumoto T.: 24,2869-2872(1983): Formulas 10 and 11 on p.2871 corrected, 2
- Sunitha K., Balasbramanian K.K. and Rajagoplan K., 26, 4393: corrections in References and
- Notes, 5490 Swindell C.S., de Solms S.J., 25,3801(1984): data listed for conformation B of structure 3b
- corrected, 1474 Uma R., Swaminathan S. and Rajagopalan K.N., 25,5825-8 (1984): Sentence on lines 20-24,page 5827 rewritten, 1114
- Wakamiya T., Nakamoto H. and Shiba T., 25,4411(1984): chemical shift of 'H NMR corrected, 2138

# Acetalisation

selective-, of aldehydes catalysed by alumina, 4767

### Acetals

- 4-oxo-2-phenylthiobutanoic acid ethylene-, Li fianion, addirion to aldehydes and cyclisation to 2-alkyl-3-furoic acids, 3635
- g-amino silvl ketene-, aldol condensation, diastereoselective synthesis of α-amino-β-hydroxy acids, 3517
- addition to a-alkoxy aldehydes, Lewis acid promoted, diastereo-selective synthesis of syn-αmethylene-β-hydroxy-Y-alkoxy
- chiral acetylenic-, reaction with Grignard reagents in presence of Cu salt; prep of chiral alkoxy allenes, diastereoselective βelimination, 4197 dioxolane-, Pd(11)-catalysed
- hydrolysis or exchange
- reactions, 705 furanose and pyranose hemi-, stereoselective replacement of anomeric OH by F atom, 5
- Y-bromo α-acetylenic-, reduction with chromous ions to a-allenic acetals and hydrolysis to aldehydes, 51
- ketene-, diastereoselective coupling, asymmetric synthesis of R-(+)- $\alpha$ -lipoic acid, 2535

- aldehydes and ketones, cyclo-
- additions, 1889 ketene silyl-, Y-addition with quaternised methyl nicotinate, synthesis of (±)-sesbanine, 3267
- ketene silyl-, diastereoselective aldol reaction with 2,3-0-isopropylidene-D (and L)-glycer-aldehydes, 5777
- leukotriene-C., acetylenic acetals and dithioacetals as antagonists, synthesis of butanoic acid 4,4'-[(4E,6Z,9Z-pentadecatriene-2-ynylidene)]bis with LT-like activity, 6427
- mixed vinylketene-, prep and participation in cyclohexenone
- synthesis, 381 S-silyl-, S,N-acetals(cf with lithium enethiolates) reaction with Schiff bases to afford βaminothioamides, 1015
- silylketene-, reaction with 3,3dimethylacryloyl chloride to give Υ,δ-ethylenic β-keto esters, application to synthesis of (±)-turmerone, 4195
- silylketene-, reaction with acryloyl, methacryloyl and crononyl chlorides and addition of MeOH to form glutaric esters,
- thioester silyl ketene-, reaction with aldehydes in presence of BF,OEt, high diastereo selectivity in anti aldol condensation, 797

### Acetoxylation

2,3-dihydro-4-pyrones, route to glucal stereochemical series, 3411

- of nitrated phenoxide, anilide and benzyl anions, effect of solvation, 2435
- selective reduction of imonium salts by sodium hydrogen telluride, effect of pH, 3693

### Acvlation

and 5,6-dialky1-2-methoxy-4(3H)-pyrimidones transformation into 1-N-acylated pyrimidine derivs, 3345

- prep of optically active monobenzoate of cis-2-cyclopenten-1,4-diol, 3099
  Friedel-Crafts-, group direction
- and protection, 39
  Friedel-Crafts-, of 1-t-butyldimethylsilylpyrrole, route to
  3-substituted pyrroles, 5035
  intramplecular rearrangement of
- 4(R)-hydroxymethyl-3-methyl-1 thiazolidine to oxazolidine, 869
- Mander carbomethoxylation of ketone enolates with methyl cyano formate, mechanism, 2291 N-substituted 3-(4-fluorophenyl)-
- indoles, prep and direct formylation at 2-position, 2155 new synthesis of α,β-unsaturated Y-and δ-lactones via
- intramolecular acylation of asulphinyl carbanions, 2253 of extended enolate ions from  $\alpha$ -
- phenylthio(PhS-)crotonate esters, 2895 reaction of 2-chloro-4,6-
- disubstituted 1,3,5-triazines with carboxylic acids, useful acylating reagents in prep of esters, amides, acid anhydrides and peptides, 2901 reaction, 2-benzenesulphonyltetra-
- hydropyran anion with
- electrophiles, 535
  regioselective-, polyalcohols via
  3-acyl-2-oxazolone-zirconium complexes, 1977
- stable 3-acylindolenine deriv via intramolecular electrophilic acylation, 5593

### Adamantanes

- 1,3,4-thiadiazoline-2-spiro-2'adamantane by acid-base reactions of 1,3,4-thiadia-zolines and thiocarbonyl ylides, 1053
- 1-bromo-,.alkylayion of βdiketones to  $\alpha$ -(1-adamantyl)- $\beta$ -
- diketones via Co(11), Co(111) and Zn(11) complexes, 3735
  2-substituted-, unequivocal assignment of syn and anti 6 13C resonances, 2841
- 3-hydroxy-4-homoadamantyl 1adamantanecarboxylate, prep and conversion to 3,4-homo-adamantanediol, 661 adamantanethione and diazomethane.
- re-examination, 1049

### Addition reactions see also Cycloadditions

- 1,2-dicarbonyl compds with methyl 3-ketoglutarate, 2163
- 1,2-dihydrophosphorin with dimethylacetylenedicarboxylate, formation of novel phosphonium ylide, 531
- 2-lithio-1,3-dithiane to unprotected Co function of partially blocked carbohydrate derivs, diastereoselection, 3685
  3-methyl-2-cyclohexenone, photo-
- addition to optically active enoates, asymmetric induction and absolute configuration of adduct, 6163
- α, β-dehydro-β-amino acid derivs via tin-promoted addition of
- malonates to nitriles, 2603 α-alkylidene-β-lactams via addition of chlorosulphonyl isocyanate to allenyl sulphides, 5001
- acetals to a-alkoxy aldehydes, Lewis acid-promoted, diastereoselective synthesis of syn-amethylene-β-hydroxy-Y-alkoxy esters, 6509 acrylate of D-pantolactone to

cyclopentadiene, isoprene andbutadiene, diastereofacial selectivity, 3095

- active olefins, alkyl additions by tributylgermanium hydride reduction of alkyl halides, 6289
- adamantanethione and diazomethane,
- re-examination, 1049 adducts from peroxidase-catalysed reaction of N-2-methyl-9hydroxyellipticinium acetate with aliphatic amino acids, reinvestigation, 4933 adducts of 2,3-bis(bromomethyl)-
- 1,3-butadiene with propiolic acids, precursors of 2H-isoindoles and 2,3-dithia-naphthalenes, 4093 aldol dimerisation of 1-indanone
- and 1-tetralones, prep of α,βand B, Y-unsaturated ketones, 571
- alkyl radicals, to vinyl sulphones and vinyl phosphonium salts, 6349
- alkynyl-Cr compds, selective addition to aldehyde moiety without affecting CO group of
- without arrecting to group of substrate, 5585 all-trans-1-acyloxy-1,3-penta-diene-5-ols as dienes in new intramolecular Diels-Alder reactions, 1249
- amine-and phosphine-boryl radicals to 2-methy1-2-nitrosopropane, 2,4,6-tri-t-butylnitrosobenzene and phenyl-N-t-butylnitrone to nitroxide spin-adducts, 1349
- aminomalonates, to difluoro-and chlorofluoro-carbene, conversion of adducts to  $\beta$ -fluorinated alanine derivs, 2445
- and aromatic substitution reactions of 1H-cyclobuta[de]naphthalenes, 1157
- asymmetric Diels-Alder reaction between chiral isoprene units 2-(R)- and (S)-benzyloxy-2,5-dihydro-4-furancarboxyaldehyde
- and cyclopentadiene, 5605 carbonyl compds and 3-bromo-1,1,1-trichloroalkanes to give 3,3-dichlorotetrahydrofurans, 4899
- chiral 1,3-oxathianes via stereoselective additioncyclisation of hydroxythicls to
- electron poor acetylenes, 4539 chiral 4-hydroxymethylbutenolide use as template for stereocontrolled conjugate addition of C-Me group, 2-Carbon extenion and reformation of lactone with new side-chain, replicated butenolide subjected to second
- conjugate addition, 5627 conjugate-, (trimethylstannyl)copper reagents to α,β-acetylenic N,N-dimethylamides, trapping of intermediates with
- trapping of intermediates with electrophiles, 6265 conjugate-, allylsilanes with α,β-unsaturated acylsilanes, 2509 conjugate-, RCu.BF, to encotes of sulphonamide-shielded alcohols, synthesis of \$-substituted
- alcanoic acids, 6051 conjugate-, to thiin-4-ones, route to thiathromboxane analogs, 2821
- to thathromboxane analogs, #02 Cu-catalysed silplzincation of acetyelenes, 4629 cuprates to vinyltriphenyl-phosphonium bromide, synthesis of 1,5-disubstituted Z\_Z-penta-1\_B\_diages\_1700
- of 1,5-disubstituted 2 a-penda 1,4-dienes, 1799 daunosoamine, new route based on stereospecific electrophilic addition to 2-acetoxy-7-oxabioyolo[2,2,1]hept-5-ene-2-abiotylla [3]
- carbonitrile, 5127 diastereofaceselection, condensation of aldehydes with

Zr enolate of chiral Npropionyl-trans-2,5-disubstituted pyrrolidine, 5807

- diastereoselective-, benzene sulphonylchloride to optically active bornyl propenoates, 4335
- diastereoselective-, nucleophiles to α-methyl-β, Y-unsaturated carbonyl compds, 3707 diastereoselectivity in intra-
- molecular additions of allylsilanes to enones, 4711
- diastereoselectivity, silylazoles to chiral aldehydes, stereo-controlled homolgation of αhydroxyaldehydes, 5477
- diastereospecific aprotic conjugate-, of carban- ions to cyclic enones to form vinylic sulphoxides and vinylic phosphine oxides, 1565
- diastereospecific aprotic conjugate-, of allylic anions bearing polar, charge-
- stabilising groups, 1569 Diels-Alder reactions of 5.5-dicyanocyclopentadiene with dienophiles to 5-substituted 7,7dicyanobicyclo[2.2.1]hept-2-enes
- with endo-selectivity, 1691
  Diels-Alder reactions of o-quino-dimethanes, stereocontrol exerted by phenyl substituents, asymmetric induction effects,
- 3413
  dienolate-, of 2-butenoic acid to
  1,3-diphenylpropenone, almost
  exclusive formation of R,R(S,R)3,5-diphenyl-2-ethenyl-5-oxopentanoic acid, 2485
  electrophilic carbene to electron
- deficient olefin, kinetics of benzylchlorocarbene-diethyl-
- fumarate reaction, 3071 enol borates to aldehydes and subsequent metal addition to form 1,3-diols, 1643 ethylmetallic reagents to acrolein
- dimer and conversion of threo alcohol to exo-brevicomin and Mus musculus pheromone, 2619 fluorides as bases in cyclopropane
- ring formation from nucleophiles and Michael acceptors, 3001 fluorosulphuric acid to an
- epoxide, 6405 free radical-, iodoacetamide to α,β-unsaturated esters,
- synthesis of 4-substituted glutarimides, 5643 Y-, of ketene silyl acetal with quaternised methyl nicotinate,
- synthesis of (±)-sesbanine, 3267 Grignard reagent to terminal tri-
- methylsilylethene moiety, stereoselective, 2101 haloacetonitriles to aldehydes via cyanomethylnickel halides to βhydroxymitriles, 155
- intramolecular trapping by epoxides of intermediates generated by organocuprate additions to propiolate esters, 3437
- intramolecular, allylsilanes to
- conjugated dienones, regio-selectivity, 2747 intramolecular-, radicals and electrophiles to allylstannanes as methods for ring closure,
- intramolecular-, to chiral vinylic sulphoxides, enantioselective synthesis of (R)-and (S)-1,7-dioxaspiro[5.5]undecane, 2221 intramolecular-, unsymmetrical allylsilanes to enones,
- geometrical limitations, 2755 isocyanide and β-allenic aldehydes or ketones to give 5-

- membered cyclic compds, 419 ketene silyl acetals, diastereoselective aldol reaction with 2,3-0-isopropylidene-D (and L)glyceraldehydes, 5777
- ketones and  $\beta$ -cyano or  $\beta$ -keto esters, addition to methacrylamide in presence of CsF/Si(OMe),, 1311 Li dianion of 4-oxo-2-phenylthio-
- butanoic acid ethylene acetal to aldehydes and cyclisation to 2alky1-3-furoic acids, 3635 Li reagents to dimethyl squarate
- at CO to give new 2-hydroxy-3,4dimethoxy-3-cyclobutenone, 1867
- lithiated cyclopropane ketals, prep and adducts with ketones, 2279
- lithium enolate of benzamidoacetone, regiospecific alkyl-ation at C-1 and stereospecific addition to cyclohexenone without protection of free NH, 3433
- Michael reactions promoted by solid-liquid phase transfer catalysis without solvent, application to addition of acetylacetone, methylacetoacetate and fluorene anions on hindered acceptors, 4601 N-tosylated α,β-unsaturated amides
- and lactams with R2CuLi or RMgX/CuI, 657
- new route to homoaporphines, addition of dichlorocarbene to aporphinoids, 1561
- nucleophile to cyclohexadienyliron cations, effect of ligand environment, 2399
- nucleophiles to alkyl  $\alpha$ -bromoacrylates, stereospecific formation of cyclopropanes, 1221
- nucleophilic-, to 2,3-0isopropylideneglyceraldehyde, high-pressure approach vs use of organometallic reagent, 4145 organocuprate-chlorotrimethyl-
- silane reagents to conjugated carbonyl compds, 6019
- organozine reagents to  $\alpha$ -enals in presence of nickel acetyl-acetonate, 829 paramagnetic heterocyclic aldo-
- nitrone, 2,2,5,5-tetramethy1-3imidazoline-3-oxide-1-oxyl, reaction with C=C and C=N dipolarophiles, formation of adducts with radical centers,
- phosphite to aldehyde, prep of chiral α-hydroxyphosphonates,
- photo-, trans-dichloroethylene with benzenoid compds, formation of tetracyclo[3.3.0.0², °0°, 6]-
- octanes and semibulivalenes, 949 pivalaldehyde and lithium enolate of pinacolone to a tetrameric
- aggregate, 3931 protected allylamines via Pdcatalysed amide addition to allylic substrates, 1749
- pyridine and Et<sub>2</sub>Mg plus EtLi resulting in 1,4-addition, 275 regioselective synthesis of 2-
- substituted pyridines via Grignard addition to 1-(alkoxy carboxy)-pyridinium salts, 3191 reversal of diastereofacial selectivity in intramolecular addition of δ-carbamoyloxy-α,β-
- unsaturated esters, synthesis of N-benzoyl-D,L-daunosamine, 4137 reversible  $d,\pi^*$ -complexation,  $\beta$ -
- cupration sequence in conjugate addition of Gilman reagents with  $\alpha,\beta$ -enones, 6015
- riboside analog of pyrimidine adduct from N-hydroxy-N-2acetylaminofluorene and DNA,

synthesis, 2659

- (+)(S)-3-trichloromethylbutyric acid via stereoselective conjugate addition of Cl\_CMgCl to erotonate of chiral auxiliary with sulphonylamino substituent, 6047
- selenothiocarboxylation, addition of S-benzoyl phenyl-
- selenosulphide to olefins, 3263 solvent controlled-, organo-titanium reagents to olefinic
- double bonds, 4423 stepwise-, of a C-electrophile and C-nucleophile to double bond of vinylacetylene, 1269
- stereoselective conjugate addition-alkylation of iron acyls, 3075
- stereoselective-, to a-alkoxy aldehydes, thioester silyl ketene acetal intermediates, 2373
- stereospecific vinyl substitution in free radical additionelimination, 4975
- t-butoxy radicals, reaction with norbornadiene, 5081 tertiary alcohols via CeCl,-
- promoted addition of Grignard reagents to ketones, 4763 tetraalkylalumin-
- ates, enantioselective addition to phenylgyloxalate, prep of  $\alpha$ substituted mandelic acid (-)-menthyl esters, 4181 thioester silyl ketene acetals and
- aldehydes in presence of BF, OEt, high diastereoselectivity in anti aldol condensation, 797 to symmetrical ketones via chiral
- propionate enolate equivs, stereoselective, 2129
- toxic marine polyethers, synthesis of segment A of okadaic acid via anti-selectivity by hetero
- conjugate addition, 5203 tricyclic diterpenes, daphnanes via high pressure intramolecular reaction of furan dienes, 2229
- uncatalysed asymmetric-, cyclopentadiene to  $(\underline{E})$ -and  $(\underline{Z})$ - $(\underline{R})$ -4,5-di-0-isopropylidene-pent-2enonates, 1631
- use of Yb(fod), in Diels-Alder reactions of acrolein with dienes, stereoselectivity, 2507 vinyl or aryl halide to enclate of β-allenyl malonate to give

# cyclopentenyl or vinylcyclopropyl derivs, 3795

- 1-vinyl-1-cyclobutanols, Pdrearrangement to 2-methyl-2cyclopentene-1-ones, 2503
- 3-buten-2-ol, model asymmetric olefin, conformational energy,
- acyclic allylic-, mercuration diastereoselectivity, 1197 africanol conversion to dactylol
- via 1,2-shift of Me group, 873 alkenols, prep, trans selectivity of ω-hydroxyalkyl-triphenyl-
- phosphonium bromides, 311 alkynyl-, conversion to β-iodoα, β-unsaturated ketones, 1967
- allyl-, benzyl and tertiary, conversion into halides with BF,-etherate/halide ion, 3863
- allylic- and benzylic-, conversion to iodides using NaI/BF,.Et<sub>2</sub>O, 2717
- allylic-, conversion into oxetane/tertiary acetate grouping of baccatin III, 3663 allylic-, hindered, rearrangement during vanadium epoxidation,

allylic-, new Claisen-nitrile oxide annulation of rings derivs

iii

- with rigid stereocontrol, 2031 allylic-, Pd-catalysed oxidative cyclisation and carbonylation to 5-membered ring lactones, 5639
- allylic-, stereospecific OHdirected ring closure, prep of tetrahydrofurans bearing 4 contiguous chiral centers and synthesis of rac-citreoviral.
- allylic-, t-butyl substituted, asymmetric epoxidation, 2543
- allylic-, transposition in stereocontrolled synthesis of  $\delta$ hydroxy allylic phosphine
- oxides, 1677 allylic-, [3.3] sigmatropic rearrangement with ethyl β,βdiethyloxyacrylate, regiospecific synthesis of
- substituted allylmalonates, 6261 and fatty acid, lipase-catalysed ester formation, preparative resolution of a-substituted
- β-acetylenic-, hydroboration followed by oxidation to hemi-acetals of Y-aldols and dehydration to 2,3-dihydrofurans, 5683
- β-substituted-, new synthesis, 657 β-trimethylsilyl-, Rh-catalysed dehydrogenation to α-trimethyl-silyl ketones, 4229
- bicyclic vinyl tertiary-, inversion of stereochemistry via [2.3] sulphoxide rearrangement, 607
- biomimetic oxidation by coenzyme PQQ-trimethyl ester, 4225
- bis-allylic-, regioselectivity in Claisen rearrangement, electronic and steric effects of
- C-4 substituents, 3655 chirality transfer in [2.3] sigmatropic rearrangement of anions derived from trialkylstannylmethyl allylic ethers, stereospecific synthesis of (2R)-3-benzyloxy-2-methyl-propanol, 5013
- cis-1,2-diacetoxycycloalkanedimethanols, enzymatic hydrolysis prep of chiral monoacetates with
- enantiomeric purity, 2073 conjugate addition of RCu.BF, to enoates of sulphonamide-shielded alcohols, synthesis of \$substituted alcanoic acids, 6051
- E-2-(phenylsulphonyl)vinyl ethers of 2,3-epoxyalcohols, stereo-specific synthesis of dihydro-furans and dihydroxy acids, 6301 E-homoallylic-, Y-hydroxyketones and cyclopropyl ketones,
- synthesis from 3-diphenylphosphinoyl propanols by acyl transfer, 5713
- epoxidation of 3(1'-hydroxyethyl)-5,8-dimethoxy-1,2-dihydronaphthalene-1-ol, allylic and homoallylic control of stereo-specificity, 6117 epoxy-, reactions, new results, 6277
- equatorial-, new stereocontrolled
- synthesis by ambiphilic reduct-ion of cyclohexanones, 3853 ion of cyclonexanones, 3653 exc-allylic, stereospecific synthesis, asymmetric synthesis of (R)-(-)-2-acety1-5,8-dimetrioxy-1,2,3,4-tetranydro-2-naphthol, 6497
- furan terminated cationic
- cyclisations with allylic alcohols, enones and N-acyl imminium salts as initiators,

- routes to spiro-cyclic systems, 5347
- gas-phase deoxygenation on iron catalyst, 3373
- HLAD-catalysed oxidation with acetaldehyde as coenzyme recycling substrate, 4527
- homoallylic-, OH-directed hydrogenation, effects of chiral and achiral Rh catalysts on 1,3stereocontrol, 6005
- homoallylic-, regio- and. stereoselective formation via anodic oxidation of polycyclic
- cyclopropanes followed by hydrolysis., 4513 homoallylic-, via Bi-mediated allylation of aldehydes, 4211
- improved pyridinium dichromate oxidation, 1699 inversion of configuration via
- nucleophilic displacement promoted by nitrate ions, 3369 new prep of alkyl iodides, 5445 novel deoxidation, use in
- synthesis of covalent perchlorates, 6243 optically active 1-phenylethanols via fungal hydroxylation of
- ethyl benzene and derivs, 6409 per-bicyclophosphoranylation , 2003
- poly-, regioselective acylation via 3-acyl-2-oxazolone-zirconium
- complexes, 1977 primary- and sec-, Pd-catalysed oxidations under solid-liquid
- phase conditions, 6257 primary-, p-anisyl as versatile protecting group, 6291 primary-, selective fluorination, 4207
- propargyl-, E-alkyl substituted, via fluoride ion catalysed reaction of F-akylacetylenes with silyl enol ethers, 79
- protected ester, nitrile, carbinol and carbinyl amine cyclopropanone hydrates, 2283
- protection, thexyldimethylsilyl chloride reagent, 5515 rate of 1,3-migration of silyl
- group from C to O using diastereomeric 7,8-epoxy-7-(trimethylsily1)-6-tridecanols, 4505
- reaction with benzosilacyclobutenes, rupture of benzyl-Si and aryl-Si, formation of (dialkyl)alkoxy-o-tolylsilanes and (dialkyl)alkoxybenzyl-
- silanes, 4761
  reaction with singlet and triplet
  diphenylcarbene, effect of temp
  on reversible ylide formation,
- regioselective ring opening of epoxides catalysed by organotin phosphate condensates, 3219 Ru-catalysed oxidation with sodium
- bromate, 2107
  sec-, new Cr(VI) reagent for
  catalytic oxidation to ketones,
- sec-, enantiomers, prep of precursor, 771
- selective desilylation of compds containing both phenolic and alcoholic t-butyldimethylsilyl ethers, 681
- sequential reactions of 2-halo-3 t-butyloxirane with p-chloro-phenolates and sodium triazolate to threo-triadimenol, 4341
- sugar cyclic sulphites stereoselective conversion to a-azido-alcohols, 6343
- tertiary-, deoxygenation via free radical method, formation of quaternary C-centres, 757 tertiary-, via CeCl,-promoted addition of Grignard reagents to

ketones. 4763

- tetrahydrofurfuryl-, regioselective opening to iodoacetonides, application to
- synthesis of brevicomins, 2315 trifluoromethylated propargylicand allylic-, Claisen rearrangement to trifluoromethylated diene derivs, 219
- via selective reduction of aldehydes by formic acid-trialkylamine-RuCl<sub>2</sub>(PPh<sub>3</sub>), 3365
- with adjacent PhS group, dehydration, 3133

### Aldehydes see also also Oximes, Hydrazones, Halogenocarbonyl compds.

- 2-phenylpropanal, mutual kinetic resolution in aldol reaction with 4,5-dihydro-5-methyl-2-(trimethylsiloxy)-3-(trimethylsilyl)furan, 1061
- 3,4-epoxy-2-methylenealkanoic acid derivs synthesis using dianion of N-pheny1-2-(phenylsulphony1methyl)propenamide and aldehydes, 4751 5-stannyl-, cyclisation, formation
- of cyclopentanol ring with retention of configuration at C
- bearing stannyl group, 3857 6α(1R-hydroxyethyl) side chain of carbapenem and penem antibiotics, introduction via a stereocontrolled aldol reaction,
- α,α'-dihydroxy ketones, new prep from ketones and aldehydes, 4925 α,α-difluoroaldehydes via Claisen rearrangements, 2861
- a, B-dialkoxy-, pinacol allylboronate, stereochemistry of reactions pinacol allylboronate, comparison with reactions of crotylboronates,
- α,β-unsaturated-, cycloadditions of ketene acetals, 1889
- α,β-unsaturated-, prep from ketones via 1,3-carbonyl transposition through one C homologation, 1581
- α, β-unsaturated-, zinc chloride mediated reduction with silicon hydrides and Pd(0)-catalyst, 1353
- a-alkoxy-, stereoselective additions, thioester silyl ketene acetal intermediates,
- α-alkoxy-, Lewis acid -promoted additions of acetals, diastereoselective synthesis of  $syn-\alpha$ -methylene- $\beta$ -hydroxy-Y-alkoxy esters, 6509
- α-alkoxy-, Lewis acid mediated 1,2-asymmetric <u>lk</u>-induction, synthesis of a-nonsubstituted
- synthesis of α-nonsubstituted
   β, Υ, syn-dihydroxyketone and ester, 3467
   α-alkoxy-, stereoselective synthesis of vinyl ethers, 1419 a-costal-, structure and synthesis
- via S-t-butyl cyanothiolacetate incorporation of labile acrolein unit, 4843 a-enals, addition to organozine
- reagents in presence of nickel acetylacetonate, 829
- a-hydroxy-, stereoselective synthesis of 1,2-diols, 1189 a-hydroxy-, enantiomerically pure, prep as intermediates for
- synthesis of arachidonic acid metabolites, 319 a-methyl-, reaction with tri-methylsilylethylidenetriarylphosphoranes, introduction of vinyl group in Cram-selective

manner, 4471

- α-methylthio-, stereoselective reaction with allyltriphenylstannane, synthesis of anti-6methylthio alcohols, 6236 acrolein dimer, stereoselective addition to ethylmetallic reagents and conversion of threo alcohol to exo-brevicomin and Mus musculus pheromone, 2619 addition of haloacetonitriles
- mediated by Ni, 155
- addition of Li dianion of 4-oxo-2phenylthiobutanoic acid ethylene acetal and cyclisation to 2alkyl-3-furoic acids, 3635
- addition to enol borates, and subsequent metal addition to form 1,3-diols, 1643
- alkynyl-Cr compds, selective addition to aldehyde moiety without affecting CO group of substrate, 5585
- allylation in presence of ketones by Sn-or Zn-mediation, 1449
- and 1-bromo-3-iodopropane, stereoselective route to terminal conjugated (E)-dienes and trienes in presence of
- SnCl<sub>2</sub>, 753 and a single C-3 unit, antidiastereoselective homoaldol reaction, stereoselective synthesis of all 4 stereoisomeric 1-carbamoyloxy-1,3alkadienes, 411
- aromatic-, Darsens condensation with phenacyl halides, bovine serum albumin catalyst, 2471
- aromatic-, direct conversion to nitriles using S,S-dimethyl-sulphurdiimide as iminating agent, dithiatetrazocines as intermediates, 2739
- aromatic-, reaction with dimethyl N-ethoxycarbonyl methyliminodithio carbonate, synthesis of 5-aryl-2-ethoxycarbonyl-4methylthio-2,3-dihydrooxazoles, 243
- aromatic-, Wittig reaction of nonstabilised P-ylides, 4587
- aryl-, Wittig reaction with carbanions stabilised with adjacent dimesitylboron group, oxidation of intermediates to erythro-1,2-diols, 5093
- asymmetric Diels-Alder reaction between chiral isoprene units 2-(R)- and (S)-benzyloxy-2,5-dihydro-4-furancarboxyaldehyde and cyclopentadiene, 5605
- β-alkyl substituted α,βunsaturated, prep using anion of 1-[(2-methoxyethoxy)methoxy]-2-phenylsulphonylcyclopropane as new d3-reagent, 3613
- Rew d -reagent, 5015 R-allenic aldehydes, addition of isocyanide to give 5-membered cyclic compds, 419 β-substituted, new synthesis, 657 Bi-mediated allylation to homo-
- allylic alcohols, 4211
- bis(seleno) and bis(thio) alkylation, 6513
- branched amino sugars via hetero Diels-Alder cycloaddition of enaminecarbaldehydes with enol
- ethers, 5273 chiral α-siloxyketone, effect of cation on diastereofacial selectivity in aldol reactions, 6009
- chiral water-soluble butadienylether with glucose hydrophilic moiety, aqueous cycloaddition with meth-acrolein to two stereoisomers, 2653
- chiral-, condensation with ethyl 1,3-dithiolane and 1,3-dithiane-2-carboxylates, diastereoface

- selectivity, 2977 chiral-, diastereoselectivity in 1,2-addition to silylazoles, stereocontrolled homolgation of α-hydroxyaldehydes, 5477
- chiral-, reaction with preformed lithium enolate of methyl 2-
- methylpropanoate, 973 chiral-, stereoselective aldol condensation with protected glycine and 3-butyryl thiazolidine-2-thiones to give  $\beta$ hydroxyesters, intermediates for β-lactam antibiotics, 977
- g-lactam antiblotics, 9// cocondensation of C with o-,m and p-tolualdehyde at 77K, deoxygenation to o-,m and p-tolylmethylenes which rearrange to benzocyclobutenes and
- styrene, 2959 condensation with Zr enolate of chiral N-propionyl-trans-2,5disubstituted pyrrolidine, diastereofaceselection, 5807
- conjugated-, exclusive 1,4reduction by sodium dithionate. 831
- conversion to t-butyldimethylsilyl protected cyanohydrins, 4275 cross aldolisation with a-bromo-ketones in presence of chromium(11)chloride, 4371
- diastereoselective aldol reaction of 2,3-0-isopropylidene-D (and L)-glyceraldehydes leading to 2-deoxy-D (and L)-riboses, 5777
- (E)-cinnamyl chloride, diastereoselective reaction with aldehydes, mediated by Sn and
- Al, 6121
  facile synthesis of vinyl
  sulphides, 1795
  formylolefination by means of
- arsonium salts, 6447 fragile and unprotected, Wittig-
- Horner reaction, 53
- from alkyl and allyl sulphones,
- hexadeca-11Z,13Z-dienal, synthesis from Z,Z-dienes prepared via acetylene carbocupration, 3285
- hindered-, stereospecific reaction with ylides of 3-dimethylamino-propyltriphenylphosphonium salts to form Z-olefins which convert to 1,3-dienes, 5747
- HLAD-catalysed oxidation of alcohols with acetaldehyde as coenzyme recycling substrate,
- ketal-, from D-threonine, synthesis of cis-\$-lactams, 33 Knoevenagel condensation,
- catalysis, 4453 Knoevenagel-type condensation with 1,3-bis(phenylsulphonyl)propene to produce electron deficient1,3 bis(phenylsulphonyl)butadienes,
- nitroarenes, reaction with nitromethane anion to form nitromethyl derivs, conversion to nitroaromatic aldehydes, 3599
- nucleophilic additions to 2,3-0isopropylideneglyceraldehyde, high-pressure approach vs use of organometallic reagent, 4145
- olefination, by a-triethylsilyl t-butylimine of propionaldehyde,
- pivalaldehyde and lithium enolate of pinacolone addition to a
- tetrameric aggregate, 3931 prep by reduction of Y-bromo α-acetylenic acetals and hydrolysis of a-allenic acetals formed,
- prep of a-hydroxymethyl ketones via alcohols from reaction of 1,4-dioxen-2-yl lithium with aldehydes or ketones, 3453

- propargylic-, cycloaddition-retroreaction sequence to 14-membered
- ansa steroid, 1705 prostaglandins from Corey lactone involving BF,-mediated reactions of a sulphone and aldehydes. 5597
- reaction with chiral 2-[(tributylstannyl)methyl]propenamides via 1,6-asymmetric induction to give a-methylene-Y-butyrolactones, 1337
- reaction with P-chloroylides to give vinylphosphine oxides, 439 reaction with thioester silyl ketene acetals in presence of BF<sub>3</sub>OEt<sub>2</sub>, high diastereo-selectivity in anti aldol condensation, 797 reaction with trimethylsilyl
- trichloroacetate, formation of silylated trichloromethyl carbinols, 1175
- reinvestigation of reaction of glyoxal with 2-(N-alkylamino)ethanol, 1853
- route to 3-amino-2,3,6-trideoxy-Dor L-hexoses via nitroaldol reaction between O-benzyl D-or L-lactaldehyde and alkyl 3nitropropionates, 1261
- (S)-2-benzyloxypropanal, reactions with Li, Mg and Zr enclates of 2-methoxypropanoate, stereo-
- structures of products, 1001 selective acetalisation catalysed by alumina, 4767 selective Y-alkylation of triiso-
- propylsilylallyl anion, synthesis of a-triisopropylsilyl aldehydes, 5375
- selective methylenation with methylene dianion synthon, 5581 selective reduction by formic acid-trialkylamine-RuCl\_(PPh,),
- to alcohols, 3365 selectivity of alkyl metal complexes in reactions with heptanal and diethylketone, comparison with Hal<sub>2</sub>M-Me, 4063
- selectivity of Cl, Hf-Bu, in reaction with heptanal and diethylketone, comparison with alkyl-Zr and alkyl-Ti complexes, 4059
- stereoselective aldol reactions using TiCl. as stereochemical template, 4129
- stereospecific intramolecular Diels-Alder reaction to form
- lactones, 3307 syntheses using methylthiomethyl p-tolyl sulphone, 2455
- tandem aldol condensation-radical cyclisation sequence, new highly convergent method for annulation of carbocycles, 6431 tetrakis(2-formy1-2-propy1)-
- ethylene, reactions and structural analysis, 1639
- transient isopropylidene methylenemalonate, generation from either Meldrum's acid or diisopropylidenemethylenemalonate and formaldehyde and trapping by dienes, 3195
- unsaturated trifluoromethyl ketones via crossed aldol condensation of 1,1,1-trifluoro-acetone with aryl or  $\alpha,\beta$ -
- unsaturated aldehydes, 2873 unsaturated, enantioselective cyclisation, via chiral Lewis acid catalysis, 5535 use of thiazolium-2-carboxylates to induce benzoin condensations,
- use of Yb(fod), in Diels-Alder reactions of acrolein with dienes, stereoselectivity, 2507

### Alkaloids

- (±)-homochelidonine, stereoselective synthesis, 5163
- 1-deoxynojirimycin and 1deoxymannojirimycin, enantiomerically pure, total synthesis
- by aminomercuration, 1123
  3-(3,3-dimethylallyl)-4-(3,3-dimethylallyloxy)quinolin-2-one reversible Claisen rearrangement to buchapsine and loss of its 1,1-dimethylallyl group, 4253
- 3-methyl-6,7-dehydroaspidospermidine system,
- synthesis, 1963 3-w-epoxido faresyl indoles, intermediate in biogenesis of sesquiterpene indole alkaloids,
- 3R,5R,9R(-)-gephyrotoxin-223 AB, enantiospecific synthesis, 1515 (-)-elenolic acid and (-)-
- ajmalicine, enantioselective synthesis, 865 (±)-dihydrofumariline-1,
- stereoselective synthesis, 917 (±)-isoretronecanol total
- synthesis from pyrrole, 2831 (±)-obscurinervidine, total
- synthesis, 1769
  (+)-β-conhydrine enantiospecific synthesis from chiral 2-cyano-6-
- oxazolopiperidine synthon, 3803 (+)-castoanspermine, total
- synthesis from D-mannose, 4617 (+)-codonopsinine, and absolute
  configuration of (-)codonopsinine, 3255
- α-acyl iminium ion inyramolecular reaction with allyl silane, intermediate for synthesis of (±)-mesembrine, 4083
- α-mannosidase inhibitor related to swainsonine, stereospecific synthesis, 3747
- α-yohimbine, total synthesis, 5227 aaptamine, synthesis from 6-methoxy-3,4-dihydroisoquinoline, 1259
- Aizoaceae and Amaryllidaceae, synthons for construction, use of C-arylation by aryllead triacetates, 5331
- amauromine, total synthesis, 847 antitumour, enantiomeric pair of unprotected AB-ring systems of sesbanimide A, synthesis from D-and L-xylose, 4639
- aporphine, intermolecular benzyne cyclo-addition to dehydronor aporphines and oxoaporphines, total synthesis of PO-3, 4559
- ary1[2.3-a]quinolizin-2-ones, synthesis, 1277
- berbin-8-ones from 2'-halogeno-1benzylisoquinolines and metal carbonyls, 3159
- biomimetic route to clavicipitic acids, intermediate in ergot biosynthesis, 4043 buddamin from <u>Buddleya davidii</u>, reappraisal <u>of structure</u>, 5251
- canthin derivs, synthesis using dialkyl oxalates, new regioselective N-alklylation,
- 385 d,1-histrionicotoxin, total
- synthesis, 5887 dihydrodioscorine and new dumetorine from Diozcorea dumetorum, isolation and stereostructural elucidation, 1615
- ergot, total synthesis via dehydrogenation of indolines with phenylselenic anhydride,
- Fe++ containing enzyme forming methylenedioxy group in berberine acts on columbamine

and not (S)-tetrahydrocolumbamine, 201

- formamidines as a-amino carbanion precursors, synthesis of 2-aryl-piperidines, -pyrrolidines and nicotine analogs, 5863
- fused pyrrolines via intramolecular cycloadditions of azides, application to pyrrolizidine alkaloids, 3527
- gluco indole, ophiorines A and B, structure and isolation from Ophiorrhiza japonica, 5299 heliotridine and related compds,
- synthesis, 2857
- histrionicotoxin ring system, stereocontrolled synthesis, 3883 homoacridone, containing tropone pina ovetom icolatio

- annulation approach, 3523 racemic anhydrocannabisativine, via tetrahydropyridine moiety, 2237
- racemic cannabisativine, synthesis, 2241
- regio-and stereo-selective alkylation at 3'-terminal end of ribonucleotides by N-2-methyl-9hydroxyellipticinium acetate, 2891
- silica supported alkaloid for HPLC resolution of racemic arylalkyl-carbinols and binaphtol derivs,
- synthesis, 3361 stable 3-acylindolenine deriv via intramolecular electrophilic acylation, 5593

benzene, role of Lewis acid catalysis, 4699 conversion of 3-hydroxylvinyl

- selenides into 1,3-bis(seleno)-propenes, 1885
- dialkylketones, synthesis from alkenes via hydroboration-
- carbenoidation, 6361 electron deficient alkenes, cycloaddition of methylenecyclopropane, regioselective formation of 2,3-or 3,4disubstituted methylenecyclopentanes, 1045
- electron-rich, acid senstive olefins, stereospecific synthesis via α,β-epoxysilanes,

indoie, asymmetric synthesis, 3291 indolizidines, synthesis by intra-molecular ene cyclisations, prep of (E)-alkylidene analogs of pumiliotoxin, 4167

isoindolobenzazocine, magallanesine, isolation from Berberis darwinii, 993 isoindoloisoquinoline alkaloids, (±)-nuevamine structure

revision, 2925 janussines A and B from Strychnos johnsonii, cross point in biogenesis of quasidimeric alkaloids, 2441

mode of inhibition of glycosidase activity by polyhydroxylated

piperidines, pyrrolidines and indolizidines, mechanism, 5073 N-2-methyl-9-hydroxyellipticinium acetate peroxidase-cayalysed reaction with aliphatic amino

acids, reinvestigation, 4933 N-methyltetrahydroellipticine, intramolecular cycloaddition of carbazoles to acetylenic vinylketenimines, 1647

necatorone, synthesis, 5975 new 2-acylindole type from Vinca

difformia, structure, 1513
new class of glycosides of polyhydroxylated piperidine alkaloids from seed of

Xanthocercis zambesiaca, 1465 new pyrrolidizine N-oxide and revised structure of

sceleratine, 5721 new pyrrolizidine, from Senecio

latifolius, 929 new route to 1-oxygenated carbazoles, synthesis of murrayafoline-A, 5841 new route to homoaporphines,

addition of dichlorocarbene to aporphinoids, 1561

new, ekeberginine from Ekebergia senegalensis, structure by use of 2D long-range δ<sub>C</sub>/δ<sub>H</sub> correlation with 1D proton-coupled ''C NMR, 4249

nigellicine, structure and isolation from Nigella sativa,

phenanthrine, from Crinum bulbispermum identical with pratorine and pratorimine,

pratorine and pratorimine, revised structures, 4301 piperidine, synthesis of (±)-pinidine, 729 protoberberine and spirobenzyl isoquinoline alkaloids, synthesis 41-414 synthesis, diradical cyclisation processes probing photoinduced

electron transfer, 5867
pyrrolizidine, (±)-isoretronecanol
synthetic study, 3311

pyrrolizidine, total synthesis of (±)-supinidine via intramolecular [4+1]pyrroline

cyclisation of verticilianes, 3397

taxane, putative biogenetic precursor verticillene, total synthesis, 3393

thebaine, via attachment of Fe(CO), access to northebaine, 14a-substit thebainone deriv and codeine analog lacking oxide ring and dihydrophenanthrene nucleus replaced by dihydro-fluorene, 501 venoterpine and cantleyine,

absolute configuration, 837

vepridimerines A-D, biogentic-type total synthesis, 4529b0185 yohimbine congeners, use of iminediene cyclocondensation in synthesis, 5983

- 2,5-dihypochloro-2,5-dimethylhexane, thermal decomposition, 4027
- hybridisation-dependence of Dinduced 15C NMR istope shifts over one bond, data for ethane, ethylene and acetylene, 1491 n-hexane conversion into 1-hexane,
- 1999 regioselectiveity changes in hexane hydroxylation with iodobenzene catalysed by tetraarylporphyrinatoiron complexes, 4247

# Alkenes

- 1,1-difuryl olefins and unsymmetrical difurylmethane derivs, synthesis, 6399
- 1-alkenes, reaction with ethyl  $\alpha$ chloro-α-phenylthioacetate, to afford butyrolactones via ene reaction products, 2463
- 2,2-dimethyl-3-buten-1-yl, radical rearrangements to 2-methy1-4penten-2-yl, kinetics at 37-74°, 1179
- (±)-trans-bergamotene synthesis from geranylacetone using intramolecular ketene-olefin cyclo-addition, 3535 activated carboxylic acids,
- intramolecular lactone annulation onto olefins by Mn(111) yields bicyclo[3.3.0] and [4.3.0]lactones, 3761 active olefins, alkyl additions by
- tributylgermanium hydride
- reduction of alkyl halides, 6289 alkenes, electro rich, and N-methylformanilide, synthesis of quinolines, 1901
- alkenes, electron poor, trapping of tribromomethyl anion, 6353 alkenes, ionic reactions with Nhalo electrophiles promoted by BF, etherate, 1811
- and a, B-unsaturated ketones, oxidation using bleomycin-Zn(11) and bleomycin-Fe(111)-iodosyl-

exo-tetrasubstituted olerins, stereospecific construction, synthesis of cyano-carbacyclins,

formylolefination of aldehydes by means of arsonium salts, 6447 fragmentation of {2-arylethyl} phosphorchloridates to 1-chloro-2-arylethanes and/or arylethylenes, 945

hybridisation-dependence of D-induced <sup>13</sup>C NMR istope shifts over one bond, data for ethane, ethylene and acetylene, 1491

hydroxyalkenes, oxidative cyclisation, prep of Y- and δlactones, 127

new reagents for [2+2] and [2+4] cycloadditions to olefins, 1,3,2,4,5-dioxadithiazin-2,2,4,4-tetraoxides, 5689

olefin addition, dipole effect of diastereoselectivity, 6313 olefin, model asymmetric,

conformational energy of 3buten-2-o1, 3647 olefination by a-triethylsilyl t-

butylimine of propionaldehyde, olefinic double bonds, solvent

controlled additions of organotitanium reagents, 4423 olefinic enol ethers, Pd-mediated

cycloalkenyklation, interception of tetrahedral intermediates,

olefins, conversion to Y-lactones by intermolecular carbolactonisation using cyanoacetic acid or malonic acid derivs in presence of Mn(111), 4291

olefins, cyclopropanation with 2-lithio-2-phenylsulphonyl propane, 6115

olefins, hydroboration with acetoxyborohydride in presence of reducible functional groups,

olefins, photoaddition of 1,4diphenylbutadiyne to cross-linked products, 765

olefins, selenothiocarboxylation, addition of S-benzoyl phenyl-

selenosulphide, 3263 olefins, terminal and internal, conversion to ketones using polyethylene glycols, 3363

olefins, tetrasubstituted, hydrosilylation, new silicon based protective group reagent,

olefins, via conversion of alkylmercuric halides, 559 Pummerer rearrangemnet product,

reaction with 1-alkenes, synthesis of 1,3-dienes and sex pheromone of red bollworm moth,

tetraarylethylenes, by coupling of activated thiotetraarylbenzo-

phenone, 5135 tetrakis(2-formyl-2-propyl)-

- ethylene, 1639 vicinal dichloroethenes, Pdcatatalysed bisethynylation, stereospecific synthesis of cisand trans-1,6-bistrimethylsilyl-hex-3-ene-1,5-diyne, 709
- Z-olefins via stereospecific reaction of ylides of 3dimethylaminopropyltriphenylphosphonium salts with hindered aldehydes, 5747Z-olefins with  $\beta$ -nitroolefinic
- moiety and ethylthio group, regio-and stereoselective formation from 1,3-dienes, 779
- [Z]-1,2-d1halo-1-alkenes, stereoand regio-selective synthesis from 1-alkynes, 1065

- allyl(chloromethyl)dimethylsilane and (chloromethyl)dimethylvinylsilane, alkoxide attack on Si to give 1-(3-butenyl)methoxydimethylsilane and allylmethoxy dimethylsilane, 1119
- C-H bond-, regioselective intra-molecular insertion by (phenyl-thio)carbene, 4483
- Li enolate opening of cis and trans-but-2-ene oxides in
- presence of BF, OEt<sub>2</sub> by S<sub>N</sub>2 mechanism, 4815 methoxides, reactions with (halomethyl)silanes at Si, facilitation by aprotic solvents and cation separation, 1115
- Si-guided intramolecular 9,10:1',2' anthracene photo-dimerisation in bis(9-anthry1)-dimethylsilane, 1505

# Alkylation

- a-methylglucoside, protection by DHP and alkylation of resulting tetrahydropyranyl ethers, 5769
- and epimerisation of lithium norcaranylidenoids, 5371
- and silylation of 2H-azirines, 2637
- and sulphenylation of chiral 0silylated imide enolates, 5339 aromatic compds with Pummerer
- rearrangement intermediates, prep of methyl-aryl compds, 2787
- aromatic compds, silica-alumina supported transition metal oxide catalyst, 3281 arylalkynes by alkyl iodides, 1843
- asymmetric synthesis of  $\alpha$ -hydroxy acids by alkylation of chiral N-(benzyloxyacetyl)-trans 2,5-bis-(methoxymethoxymethyl)-pyrrolidine, 1343
- asymmetric tandem-, of αalkylidene-Y-butyrolactone derivs, 3027
- asymmetric-, of chiral dipole stabilised anions, mechanistic
- Horder, 3293 B-diketones with 1-bromoadamantane to α-(1-adamanty1)-β-diketones via Co(11), Co(11) and Zn(11) complexes, 3735 bis(seleno)- and bis(thio)-, of
- ketones and aldehydes, 6513 bisethynylation of vicinal
- dichloroethenes, Pd-catalysed, stereospecific synthesis of cis
- stereospectic symmetris of cir-and trans-1,6-bistrimethylsiTyl-hex-3-ene-1,5-diyne, 709 competitive reactions of RMgX with nitrobenzenes, ketone, ester, cyano- and iodo-derivs and some functionalised nitrobenzenes,

diastereoselective tandem-, of acyclic α,β-unsaturated estersbased on use of dithioacetalunit, 3031

- enantiomerically pure aminocyclo-hexanes via Birch reductionalkylation of chiral anthranilic acid deriv, 1619
- enolisable ketones, carbonyl
- methylenation, 5579 exclusive Y-, of diene allylic t-amines with Grignard reagents,
- Y-, of  $\alpha,\beta$ -unsaturated ketones, α'-phenylsulphonyl group as regioselective control, 169
- intramolecular, stereoselective-of 3-,5-,6-and 7-membered carbocyclic compds with extracyclic chiral center 1723
- lithium enolate of benzamidoacetone, regiospecific alkyl-ation at C-1 and stereospecific addition to cyclohexenone with-out protection of free NH. 3433
- methoxy(phenylthio)trialkylsilylmethane in new synthesis of αketosilanes, 2675
- N-, of weak N-H-acids by PT catalysis, mechanism, 297
- new regioselective N-, synthesis of canthin alkaloid derivs using
- dialkyl oxalates, 385 nitrones by N-, of aldoxime and ketoxime-O-trimethylsilyl ethers, 4331
- polyalkoxy-2-aza-propenylium salts, synthesis via oxygenalkylation of alkoxy substituted N-methylene amides,
- stereochemistry, 3547 quadruple bio-, of sterol side chain, 4031 quinone-imonium cation, synthesis
- using DMSO as methylating agent, 3145
- reaction, 2-benzenesulphonyltetra-hydropyran anion with electrophiles, 535
- regio-and stereo-selective-, at 3'-terminal end of ribo-nucleotides by N-2-methyl-9-hydroxyellipticinium acetate,
- regioselective-, of lithium enolates derived from 2heptanone, 1599
- regioselective-, of piperazine-dione derivs, 2955 selective-, Y-alkylation of tri-
- isopropylsilylallyl anion, synthesis of α-triisopropylsilyl aldehydes, 5375
- selective methylenation of aldehydes with methylene dianion synthon, 5581 spiro-, of indene, prep of spiro
- hydrindandione ring system of Fredericamycin A, 4725
- stereoselective conjugate addition-alkylation of iron
- acyls, 3075 stereoselective ortho-, of phenoxyvinylphosphorane to spirophosphorane, 5443
- stereoselective-, of benzylic acetates and free alcohols of (II-arene)tricarbonyl-Cr
- complexes to exo-alkyl Cr-complexes, 767
  stereospecific-, at C-6 of pyranose sugar, conversion of glycosyl fluorides into Cglycosides, 1823 trimethylsiloxy mercaptides to
- vinyl sulphides, 1795
- use of chiral butyrolactone as template to control stereo chemistry of C-methylation on

corresponding enolate, two-C homolgation and second Cmethylation, 5623

### Alkynes

- 1-alkynes, stereo-and regioselective synthesis of [Z]-1,2dihalo-1-alkenes, 1065
- 1-ethoxy-3-trimethylsilyl-1propyne, regio- and stereo-selective reaction with saturated and ethylenic carbonyl derivs giving new a-functional allylic, conjugated dienic or tienic silanes, 1861
- 1,4-diarylbutadiynes and mixtures of 1,4-dialkyl-1,3-butadiynes and 3-alkyl-4-(1-alkynyl)-hexa-1,5-diyn-3-enes, Pd-promoted prep, 523
- 1.4-diphenylbutadiyne and olefins photoaddition to cross-linked
- products, 765 7,8-acetylenic analogs of hexahydro leukotriene-E, with agonist and antagonist activities, synthesis, stereoselective route to E-and
- Z-enynes, 6423 acetylenes, Cu-catalysed silyl-
- zincation, 4629 alkoxalkyne, reaction with pyrrole-and phenyl-carbene Crcomplexes, dimer formation via o-quinonemethide and subsequent
- dimerisation, 1159
  alkynes, iodination, phenyl shifts
  to vinyl cations formed, 1967
  analogs of leukotriene via
- acetylene carbocupration, 2731
- arylalkynes, alkylation with alkyl iodides, 1843 cycloaddition with pyrrole-carbene chromium complex to give acetylated hydroindologuinone deriv, 2969
- electron poor acetylenes, stereoselective addition-cyclisation of hydroxythiols, prep of chiral 1,3-oxathianes, 4539
- enyne triflates, reaction with nucleophiles, 2301
- F-alkylacetylenes, generated in situ, treatment with silyl enol ethers, new route to F-alkyl substitutd propargyl alcohols or 4-(1H,1F-alkylidene)1,3dioxolane derivs, 79
- formation via coupling alkyl and allyl halides with propargyl dianion equivalent with high positional selectivity, 271
- hybridisation-dependence of D-induced <sup>13</sup>C NMR istope shifts over one bond, data for ethane,
- ethylene and acetylene, 1491 keto alkyne cyclisations to heterocyclic analogs of prostacyclins, 1171
- phenols from metal-carbynes and diynes, 2159
- phenyl acetylene oligomerisation over titanium oxide on silica gel-alumina, 6245
- photocycloaddition of N-methyl-thiophthalimide, 2249 terminal acetylenes, conversion to Mannich bases via gem-aminoethers, prep of cyclic aminoacetylenes from ω-acetylenic sec amines, 3203 terminal acetylenes, Pd-complex
- catalysed reactions with diphenylketene giving disub-
- stituted acetylenes, 3697 trimethylsilylethyne, double
- coupling with cis-and trans-1,2-dichloroethene, 709
  vinyl acetylene, stepwise addition of a C-electrophile and a C-nucleophile to double bond, 1269

### Allenes

- 1-chloro-1-aryloy1-3,3-diary1--cnioro-laryloyi-3,3-diary1-allene, Cu-assisted cross coupling to 3,7-dioxa-2,6-diary1-4,8-bis(diarylmethylene)-bicyclo[3.3.0]octa-1,5-diene, and 1,1,4,4-tetraary1-3,6-diaryloylhexa-1,2-diene-5-yne,
- allylic functional groups, reductive removal by nickel boride, 2581
- as acceptor-donor reagents in 3+2 annelations, synthesis of
- carbapenam ring system, 4315 bromo-, convesrion to alkylallene by t-Bu(CN)CuLi, stereochemistry, 65
- chiral alkoxy-, via reaction of chiral acetylenic acetals with Grignard reagents in presence of Cu salt; diastereoselective βelimination, 4197
- controlled lithiation, formation of efficient propargylic anion equivalent in coupling with alkyl and allyl halides,271
- Ni-promoted route to substituted allenes by reaction of 1-brome

- aldehydes, mediated by Sn and Al, 6121
- alkylation of aromatic compds, silica-alumina supported transition metal oxide catalyst,
- BF<sub>3</sub>-etherate on alumina catalysis in improved synthesis of cannabidiol, 1083 chiral bisazetidines via hydro-
- alane reduction of bis-6lactams, 2035
- organo-compds, ketone synthesis via Pd-catalysed carbonylation, 4819
- organo-, promoted pinacol-type rearrangement, conformational effects, 4781
- selective acetalisation of aldehydes catalysed by alumina, 4767
- tetraalkylaluminates enantioselective addition to phenylgyloxalate, prep of αsubstituted mandelic acid (-)-
- menthyl esters, 4181 trialkyl-, reagents, allylation of allylic cyclohexenyl esters, egio-and stereo-chemical study
- trialkyl-, stereoselective alkylation of benzylic acetates and free alcohols of ( $\Pi^s$ -arene)tricarbonyl-Cr complexes, 767

Ami des

- 2-alky1-3-oxoamides, potassium triethylborohydride reduction to
- triethylborohydride reduction to 3-hydroxyamides with antidiastereoselectivity, 4643
  3,4-epoxy-2-methylenealkanoic acid derivs synthesis using dianion of N-phenyl-2-(phenylsulphonyl-methyl)propenamide and aldehydes, 4751
  2-bydroxympotyl-anylamides
- 3-hydroxypent-4-enylamides stereoselective intramolecular aminocarbonylationcatalysed by Pd, 4479
- (+)-pederin, total synthesis, stereocontrolled synthesis of (+)-benzoylpedamide, 6461
- alkylamides via dialkylcuprolithium reagents and N-methyl 2-(2,4,6-trimethy1)cyclo-11,3-oxoimminium salts, 1013
- and esters from aziridine 2 carboxylic acid salts, 4439 anilides, synthesis of 2-phenyl-indoles via thallation, 5963
- carboxylic amides, a, B-unsaturated by homolgation of ketonesvia Pd-catalysed enol triflates, 1109 cathodic degradation of di-imines
- in presence of oxygen, synthesis
- in presence of oxygen, synthesis of amides, 3005 chiral 2-[(tributy1-stanny1)methy1]propenamides, reaction with aldehydes via 1,6-asymmetric induction to give amethylene-Y-butyrolactones, 1337
- methylether reductions (1) diamion of N-phenyl-2-(phenyl-sulphonylmethyl)propenamide, reagent for synthesis of (Ε)-α,β-unsaturated amides and 5,6dinydro-2H-pyrans, 4747
- methacryamide, addition toketones and β-cyano or β-keto esters in presence of CsF/Si(OMe), 1311
- N,N-diethylthiophene-2-carboxamide, 3,5-dillthiated, regioselective synthesis of 2.3-and 2,3,5-substituted thiophenes, 1149
- N-methylformanilide and electronrich alkenes, synthesis of quinolines, 1901 N-tosylated α,β-unsaturated
- amides, conjugated addition

- with R2CuLi or RMgX/CuI, 657 natural insecticidal isobutylamides, synthons for general routes, 2477 natural polyene isobutylamides,
- stereochemistry of Wittig reactions, 2481
- oxaziridine-amide isomerisation. use in creation of peptide bonds, 6065
- polyalkoxy-2-aza-propenylium salts, synthesis via oxygenalkylation of alkoxy substituted N-methylene amides, stereochemistry, 3547
- precursor of thienamycin and related carbapenems, cyclisation of N-propargyl epoxyamide to acetylenic 2-azetidinone, 4521
- protected allylamines via Pd-catalysed amide addition to allylic substrates, 1749
- resolution of β-hydroxycarboxylic acid derivs on chiral stationary phase derived from (R,R)tetramide, 4217 sec-,.carboxamides,R2NHCOR1, new
- synthesis using 2-methyl-2-ovazoline as huilding block
- stereoselective liver-esterase hydrolysis of rigid bicyclic meso-diesters, prep of optically pure 4.7-epoxytetra- and hexahydrophthalides, 4087
- synthesis, using new coupling reagent, benzotriazol-1-yl diethyl phosphate, 1341 tertiary benzamides, phthalamides
- and 0,0'-aryl dicarbamates, dilithiated synthons, 1145
- (trimethylstannyl)copper reagents conjugate addition to α,β-acetylenic N,N-dimethylamides, trapping of intermediates with electrophiles, 6265
- unsaturated-, synthesis of iodolactam, 1803

# Amidines

- chiral formamidines derived from  $\beta$ -carbolines, route to indole alkaloids via metallation and alkylation, 3291 chiral formamidines, stereo-
- selective deprotonation, deuteriation and alkylation, 3295
- convulsant pyrrole-3-carboxamidine from Brunfelsia grandiflora, 2623
- dipole measurements of 7 polar amidine containing compds of biological interest, 1897
- formamidines as a-amino carbanion precursors, synthesis of 2-aryl-piperidines,-pyrrolidines and nicotine analogs, 5863

### Amination

- azides using tetracarbonylhydrido-ferrate a selective reducung agent, 3277
- sodium azide, reaction with 9-bromo-6,7,8,9-tetrahydro-4H-pyrido[1.2a]pyrimidin-4-ones to give 9-amino-6,7-dihydro analogs, 3521

- 1,4-epimine compds, synthesis, reaction of IBDA-iodine with N-nitroamines, Ncianamines and N-phosphoroamidates, 2493
- 2-aminopyridine, reaction with benzil, correction, 247 α,α-dimethylarylalkylamines, new synthesis from 1-diphenyl-
- phosphiny1-2,2-dimethy1-

- photooxygenation of tetramethylallene competing with (2+2)cycloaddition and ene reactions with singlet oxygen, 5029
- tetraphenyl-, electron transfer photooxygenation, formation of 1,3-dihydroperoxy-1,1,3,3-tetrapheny1-2-propanone and decomposition under chemiluminescence, 173
- tricyclic thromboxane Az analogs, prep using oxyallyl method, 4411

### Allomones

5101

- $(\pm)-\delta^{9}(^{12})$ -capnellene, total synthesis, 4847
- aglajne-1 polypropionate metabolite, C-C connectivity via long range 'H-1'C couplings,
- defence against rice blast, stereoselective synthesis of coriolic and dimorphecolic acids, 465
- new sesquiterpenes in proposed Russulaceae chemical defence system, 3163

### Allylation

- aldehyde, in presence of ketones by Sn-or Zn-mediation, 1449 aldehydes, Bi-mediated allylation
- to homoallylic alcohols, 4211 carbonucleophiles and protection-deprotection of amines, 2449
- cyclopropene and cyclopropane derivs via selective
- derivs via selective cyclialiylation of alkenyl-lithiums, 5671 direct-, of aryl bromides with allyl acetate, 6457 Pd-catalysed-, of lithium 3alkenyl-1-cyclopentenolates-triethylborane and application
- (Z)-jasmonate, 2177 (2)-jasminate, 2177 regioselective photo-, of dicyano-polycyclic aromatic compds with allyltrimethylsilane, 5819 stereospecific-, of "ribo" and "galacto" aldosulose derivs, 823 with trialkylalumiunium reagents

to selective synthesis of methyl

of allylic cyclohexenyl esters, regio-and stereo-chemical study,

# Aluminium compounds

(E)-cinnamyl chloride, diastereoselective reaction with

- aziridine, 1245 activation of P-chiral dialkyl phosphates with triphenylphosphine-CCl, reaction of intermediate phosphonium salt with aniline or pyridine with inversion of configuration at
- aminobiphenyls, via reaction of arylhydrazines benzene in presence of TFSA, 5811
- aminocyclohexanes, enantiomerically pure, via Birch reduction-alkylation of chiral anthranilic acid deriv, 1619
- aromatic-, alkylation in presence
- of inorganic bases by PT catalyst, 297 aromatic-, and heteroaromatics, radical cations generated via action of nitroxides, 4201
- benzenediazonium chlorides, 4-substituted, reaction with methylamine-formaldehyde mixture to give 3,7-bis(4-X-aryl)-,5,3,7-dioxadiazocines, 1043
- CD of  $\beta$ -cyclodextrin with amino moiety, binding ability for ferrocenecarboxylic acid in DMSO, 899
- cyclodiene insecticides, amineinduced photodehalogenation,
- diene allylic t-amines, exclusive Y-alkylation with Grignard
- reagents, 1009 formamidines as α-amino carbanion precursors, synthesis of 2-arylpiperidines, -pyrrolidines and nicotine analogs, 5863 histamine liberating thapsigargin
- and trilobolide, absolute configurations, 107
- intramolecular reactions of acyclic N-acyliminium ions with propargyl silanes leading to derivs of 3-vinylidene-pyrrolidine, 3-vinylidene-piperidine or 1-amino-2-
- vinylidenecyclopentame, 3151 macrocycles with sulphide and amine binding sites as chiral ligands for Ni-catalysed cross coupling of Grignard reagent with vinyl bromide, 3499
- methylamine, photoreactions with 4-nitrocatechol ethers occurs at meta position in respect of
- nitro group, 2489 methylamine, reaction with cycloheximide carrying 4-nitrocatechol ether moiety, model for photoaffinity labelling, 2489
- N,N-disubstituted hydroxylamines Ti(111) induced transformations to imines and sec amines, 4633
- N-acetoxy-N-acetylaminobiphenyl, solvolysis, no nitrenium ion involved in production of hydroxamic acid, 147
- N-methanesulphonylalkylamines, regiospecific cyclisation to Nmethanesulphonylpyrrolidines, 1877
- neutral aminyl radicals, prep and H-abstraction to epimine compds,
- open-chain aliphatic aminium radicals, intramolecular H-abstraction with 8-membered cyclic transition state, 3407
- primary and sec, via BF3-etherate ring opening of substituted aziridines by organocopper reagents, 1153
- protected allylamines via Pdcatalysed amide addition to allylic substrates, 1749 reaction with di-2-pyridyl thionocarbonate to form isothio-

cyanates, 1661 sec, dehydrogenation with phenylselenic anhydride or acid,

- synthesis of α-cyanoamines, 1229 sec-, reductive carboxylation, synthesis of tertiary Nmethylated amines, 5367
- selective reduction of C=C in conjugated enones by benzyl-amine, variant of Sommelet reaction, 5217
- stereochemistry of products of reaction with 1,3-dimethylthymine epoxide, 3587
- tertiary aromatic-, electron transfer of organic ligands from iron carbonyl complexes, 737 tertiary-, axially disymmetric,
- reductive cleavage by LAH, synthesis of new 1,1'-binaphthyl
- substituted amines, 421 tryptophan, oxidative transformation to 3-(2-aminophenyl)-2-pyrrolidine or kynurenine, 5871
- vanillylamine, direct lithiation and metal-halogen exchange of bromoisovanillylamines, 2151
- vanillylamine, direct litiation and metal-halogen exchange of bromoisovanillylamines, 2151 via rapid reduction of aliphatic
- nitro compds, 6413

### Amino acids

- 2-alkylated-3-deuteriated-1-aminocyclopropane-1-carboxylic acids, regiospecific synthesis, 481
- 2-amino-3-hydroxy-4-pentenoic acid derivs, lactonisation, synthesis of natural 3,4-dihydroxyprolines, 5307
- 4-amino-2(5H)-furanones, new
- synthetic method, 2459 α,β-dehydro-β-amino acid derivs via tin-promoted addition of malonates to nitriles, 2603 α-amino silyl ketene acetals, aldo
- condensation, diastereoselective synthesis of α-amino-β-hydroxy
- acids, 3517

  α-dehydroxyglutamic acid derivs,
  synthesis and reactions, 85
  αamino acid derivs, prep ofpyrrolidines, 2775
- alanine derivs, aminomalonates reaction with difluoro-and chlorofluoro-carbene, conversion of adducts to  $\beta$ -fluorinated alanine derivs, 2445
- aliphatic-, peroxidase-catalysed reaction with N-2-methyl-9hydroxyellipticinium acetate, reinvestigation, 4933
- analog structure of host-specific toxin from Helminthosporium carbonum, 969
- bicyclic dipeptide with shape of β-turn central part, synthesis from L-glutamic acid and L-
- cysteine,, 647 bioactive (+)-3-methylenecyclopropyl)glycine and  $(\pm)-3$ -(carboxycyclopropyl)glycine,
- synthesis, 83 carzinophilin degradation product, enantioselective synthesis of (2S,3S) 4-amino-2,3-dihydroxy-3-
- methylbutyric acid, 3299 chiral synthesis of statine, 2973 condensation with meso-tetra(2-aminophenyl) and meso-α, Y-bis-(2,6-diaminophenyl)porphyrins, 1019
- cycloaddition of 2-chloroacyrlo-nitrile and N-(2-pyridinecarbonyl)proline and thio analog afford bridged compds, 5447 depsipeptides with D-α-hydroxy

carboxylic acids, synthesis via

- $L\text{-}\alpha\text{-}\text{halo-carboxylic}$  esters and cesium salts of N-protected
- amino acids, 5257 erythro- and three-3-fluorophenyl-alanine from protected glycine synthon, 3067
- glycine cation equiv, electrophilic, prep and reaction with heteroatom nucleophiles, 695
- glycine, protected, and 3-butyryl thiazolidine-2-thiones stereoselective aldol condensation with chiral aldehydes to give β-hydroxyesters, intermediates for β-lactam antibiotics, 977
- helminthosporic acid analog, prep using oxyallyls, 6133
- inhibitor of Y-aminobutyric acid transaminase, (Ε)-β-fluoro-methyleneglutamic acid prep, 4091
- isoxzolin-5-one-2-yl ring system, synthesis via 5-endo-dig cyclisation, enantio-synthesis of L-β-(isoxazolin-5-one-2-yl)-
- alanine, 5931 L-proline, transformation to Namino-2-pyrrolidones via mesoionic system, 5739
- N,O-bisacryloyl-L-phenylalaninol, synthesis, application in prep of L-phenylalanine ethyl ester selective acrylic polymer, 3623 new bulgecinine, amino acid in
- bulgecins, synthesis, 4759
- new route to chiral β-hydroxy-αamino acids, cleavage of oxirane with hydrazoic acid, 5309
- optically active aminoketones from (S)-serine, 4055
- protected racemic β, Y-unsaturated  $\alpha$ -amino acids via Y-phenyl-seleno- $\alpha$ ,  $\beta$ -unsaturated esters, 1959
- protected vinylic amino acids via intermolecular Lewis acid
- catalysed ene reactions, 3115 rac 2-alkyl-1-aminocyclopropane-1carboxylic acids, resolution, 485 S-(5'-deoxy-5'-adenosy1)-(±)-2-
- methylhomocysteine from (±)-2methylmethionine, 1135
- S-nitroso and S-phenylsulphonyl derivs of L-cysteine and glutathione, 2013 specific inhibition of β-D-
- glucuronidase by 3R,4R-
- dihydroxy-L-proline, 5379 stacking interaction between tryptophan and uracil in synthetic model, 4467
- synthesis, reaction of glycine cation equiv with carbon nucleophiles, 699
- thiosulphonate derivs, synthesis,
- tryptophan, oxidative transformation to 3-(2-aminophenyl)-2-pyrrolidine or kynurenine, 5871

### Amino alcohola

- 1,2-amino-, and carboxylic acids, stereochemistry of Ph,P-CCl, mediated cyclisation to oxazolines with inversion of carbinol center, 4687 β-amino-, from epoxides and halo-
- magnesium alkylamides, 3107 β-amino-, enantiospecific prep
- from N-cyanomethyl-4-phenyl-1,3-oxazolidine synthon, 6345 regiospecific opening of oxetanes with trimethylsilyl cyanide-zinc iodide, approach to Y-amino alcohols, 4971
- reinvestigation of reaction of glyoxal with 2-(N-alkylamino)-

ethanol, 1853 Ru-catalysed oxidation of cyanohydrins to acyl cyanides, reagents for selective Nbenzoylation of aminoalcohols, 925

### Aminocarbonylation

3-hydroxypent-4-enylamides catalysed by Pd, 4479

### Amino esters

- α-amino esters, .via reaction Grignard reagents with methyl 2-acetamidoacrylate in presence of CuI, 4387
- active-, of amino acid derivs, reagents for synthesis, 1721
- amino norbornenecarboxylates, synthesis via methyl acrylate and amino butadienecarboxylates,
- aromatic amino acid ethyl-, via  $\alpha$ -chymotrypsin in solns of high EtOH concentration, 6081
- methyl-, of N(3,5-dinitro-benzoyl)amino acids, chiral solvating agents for NMR determination of enantiomeric purity, 2989
- N,O-bisacryloyl-L-phenylalaninol, synthesis, application in prep of L-phenylalanine ethyl ester selective acrylic polymer, 3623 t-butyl 4-alkoxy-3-amino-2alkenoates, lactonisation, 2459

### Amino ethers

gem-amino-, conversion of terminal acetylenes to Mannich bases. prep of cyclic aminoacetylenes

unsaturated esters, synthesis of N-benzoyl-D,L-daunosamine, 4137 route to 3-amino-2,3,6-trideoxy-Dor L-hexoses via nitroaldol reaction between O-benzyl D-or L-lactaldehyde and alkyl 3nitropropionates, 1261

stereocontrol in isoxazoline reductions, synthesis of aminodeoxy-DL-xylo-and arabinopentose derivs from furo-isoxazolines, 2997

### Ammonium salts

- allylic ammonium salt inter mediates in subsitution of allylic t-amines with Grignard reagents, 1009 anti-[2.2](1,6)azulenophane via
- fluoride induced 1,8-elimination from trimethylsilyl-tetraalkyl-
- ammonium salts, 4835 cationic dihexadecyldimethylammonium bromide, permeation of Ellman's anion from surface to interior, 603
- dehalogenation of mono-or polychlorinated aromatic compds via ammonium formate catalytic transfer hydrogenolysis, 1381
- quaterary-, reductive cleavage by LAH, synthesis of new 1,1'-binaphthyl substit amines, 421 quaternary-, cleavage by sodium hydrogentelluride, 6197

### Anilinium salts

N-adamantylidylcarbinyl-N,Ndimethyl-4-cyanoanilinium bromide, synthesis and aqueous reactivity, 2183

- dehydroaza[18]annulene, synthesis from annulenones, 639 4:7,10:13-diepoxy[15]annulenone, transformation by carbonyl transposition reactions, 5567 dihydro[10]annulen-11-one, skeletal rearrangement, effect of temp and counter ion, 3511 dimeric[14]annulenes, electrochemical behaviour, 4879 doubly charged ions of bridged [4n]annulenes, evaluation of diatropic and current effects,
- hydrazino bridged [14]annulenes, frontier orbital control, 5425 new bridged [4n]annulene, 1,6:9,14-bismethano[6]annulene,
  - molecular structure and dynamic
- behaviour, 3087 Ni-catalysed coupling of bromides of 1,6-methano[10]annulene and azulene, synthesis of biannulene and bi-, ter-, quater- and poly-azulenes, 3829

# Anthocyanins see also Pigments

bis-malonylated, from Monardia didyma, structure of monardaein,

### Anthracene and derivatives

- 1,3-bis[2-(2-naphthyl)vinyl]benzene photocyclisation leads regioselectively to dinaphth-[1,2-a:2',1'-j]anthracene, 3773
- 9,10-anthracenediones, functionalised, synthesis via tosylate and triflate phenolic activation, 157

### Amino ketones

azabicyclic enaminones by pyrolysis of aminomethylene Meldrum's acid involving -1,4 H shift from aminomethyleneketene intermediate, 833 L-proline, transformation to N-

amino-2-pyrrolidones via mesoionic system, 5739 optically active aminoketones from

(S)-serine, 4055

# Aminolysis

chiral α-aminonitriles from new Ncyanomethyl-1,3-oxazolidine synthon, 3567

# Amino sugars

- branched, via hetero Diels-Alder cycloaddition of enaminecarbaldehydes with enol ethers,
- chiral nitronate dianion from (+)-D-gylceraldehyde, generation and use in synthesis of 3-amino-2,3-dideoxy sugars, 6269 daunosoamine, new route based on stereospecific electrophilic addition to 2-account.
- addition to 2-acetoxy-7-oxabicyclo[2.2.1]hept-5-ene-2carbonitrile, 5127
- deoxygenated aminoglycosides, prep and new pathway to L-daunosamine, 1863 glycosidase inhibitors,
- enantiomerically pure 1-deoxynojirimycin and 1deoxymannojirimycin, total synthesis by aminomercuration,
- N-acetyl-D,L-acosamine and Nbenzoyl-D,L-ristosamine, diastereoselective synthesis, 4133
- reversal of diastereofacial selectivity in intramolecular addition of δ-carbamoyloxy-α,β-

- benzamides, use in tandem directed lithiations, 6213 10-methyl-4-ketohydroazulenes synthesis using organotin, (3+2)-cyclopentane annulations, 3531
- carbocycles, new highly convergent method, via tandem aldol condensation-radical cyclisation sequence, 6431
- fused cyclopentanes with bridgehead OH group, synthesis using Lewis activated bisfunctional annulation, 3629
- intramolecular lactone annulation of activated carboxylic acids onto olefins by Mn(111) yields bicyclo[3.3.0] and [4.3.0]lactones, 3761 new approaches to 5- and 6-
- membered carbocyclics, 2513
- new Claisen-nitrile oxide annulation of rings onto allylic alcohol derivs with rigid stereocontrol, 2031 new cyclohexenone, reaction of a-
- n-butylthiomethylene ketones with 1-lithio-1-methoxycyclopropane to afford α-n-butylthiomethylene spirobutanones and
- their ring expansion, 713 pyrrolizidine alkaloids, total synthesis of (±)-supinidine via intramolecular [4+1]pyrroline annulation approach, 3523 ring transfer of furans to fused
- furans by tandem intramolecular Diels-Alder and ring opening of adducts, 2689

# Annulenes

2-ethoxy-6,11-dimethyl-7,9-bisdehydroaza[14]annulene and benzannelated derivs, synthesis, 635 2-ethoxy-6,11-dimethyl-7,9-bis-dehydroaza[16]annulene and 2ethoxy-8,13-dimethyl-9,11-bisnew tetrahydroanthraquinones from genus Cortinarius, 2593

### Antibiotics

- 1, neothramycin synthesis, application of 4-benzodiazepines, one-step synthesis, 5947
- 1α-and 1β-methyl thienamycin, total synthesis and and stereochemical assignments, 587
- 2-azidocarbapenems, prep and
- reactions, 5407 4-demethoxy-8-nordaunomycinone, total synthesis, 691
- 6-deoxy-daunomycinone and 1hydroxy-4-demethoxy-11-deoxydaunomycinone, regiospecific synthesis, 5693 11-deoxyanthracyclinones via
- cycloaddition of substituted tetrahydrohomophthalic anhydride, regiospecific synthesis, 1549
- (-)-carpetimycin A, synthesis,
- (±)-aphidicolin, total synthesis, 6147
- (±)-cycloeudesmol, new stereoselective total synthesis via stereoselective cyclopropane ring formation, 235
- (±)-pyrenophorin and (±)-
- colletallol, syntheses, 1989
  (+)-thienamycin from 3(R)-hydroxybutyric acid, stereocontrolled
  synthesis, 1523
- (+)-thienamycin, carbapenem and penems, saynthon for synthesis, 937
- acetoxyazetidinones, reaction with trimethylsilylacetyl thiol-esters, prep of azetidinone-thiolester precursors to carbapenems, 573 amphotericin B, synthesis of C-D fragment and C(21)-C(37) unit of

the aglycone, 5239 anthracycline precursor, synthesis of chiral tetralin from α-D-isosaccharino-1,4 lactone, 5295 anthracyclinones, optically active, asymmetric synthesis,

anthramycin analogs structure and

dioxabicyclononane unit of tirandmycin, synthesis, 2173 discodermins B,C and D, structure and isolation from Discodermia kilensis, revised structure of discodermis A, 855 enatioselective synthesis of

chlorothricolide intermediate,

polyene macrolide, comparative study of polyene syntheses, 4891 polyene macrolides, association with alkali metal salts in MeOH, 253 precursor of thienamycin and

triazolvl lphosphorothicate, prep

related carbapenems, cyclisation

antiherpetic agent, synthesis of (S)-9-(2,3-dihydroxy-1-propoxy-methyl)guanine, 1815 antitumour and antimicrobial

flavonoid, uvaretin, synthesis, 4807

antitumour geiparvarin, new nitrile oxide based synthesis,

antitumour halogenated clavulone analog, synthesis, 5053 antitumour hexamethylmelamine,

acylhydroperoxide oxidation to 2.4-bis(dimethylamino)-6-

2,4-bis(dimethylamino)-o-[(dimethylamino)oxy]1,3,5-triazine, 5247 antitumour, 6-ethyl-10-methoxy-7H-pyrido[4.3-c]carbazole, new synthesis, 4929 antiviral aphidicolin via reaction

of key dienoid intermediate with unsymmetrical dienophiles, 2833

avermectin B<sub>1</sub>a, chemical degradation, 4279 avermectin-milbemycin hybrid

synthesis, 4283 benz[a]anthraquinone X-14881 C and ochromycinone, synthesis by

directed ortho metallation, 9 biosynthesis of terrecyclic acid A, confirmation of cyclisation mechanism and H-shifts using [2-2H,] acetate and [2-13C2H,]-

acetate, 3845 B/C component of CC-1065, synthesis via regiospecific introduction of oxygen

functionality, 2985 carbapenem and penem, 6α(1R-hydroxyethyl) side chain, introduction via a stereocontrolled aldol reaction, 6285 carbapenem, 1-substituted,

synthesis, 4739 carbapenem, synthesis based on symmetrisation-asymmetrisation, 5831

carpetimycins, optically active, via stereoselective reduction of chiral 3-isocyanoazetidinone,

carzinophilin degradation product, enantioselective synthesis of (2S,3S) 4-amino-2,3-dihydroxy-3-

methylbutyric acid, 3299 construction of optically active tricyclic lactone sub unit for total synthesis of (+)-pleuromutilin, 1603,1607

construction of pentacyclic carbon skeleton and total synthesis of (±)-naphthyridinomycin, 1911

cyclobutanone analogs of β-lactam antibiotics, synthesis of Nacetyldeazathienamycin, 5363

cycloheximide with 4-nitrocatechol ether moiety, reaction with methylamine, model for photoaffinity labelling, 2489

arrinity labelling, 2489 DEF-ring system of nogalmycin, chiral synthesis, 2693 degradation of (+)-pleuromutilin or tiamulin direct to (-)-bicyclic ketone intermediate,

dihydromaesanin, prep from p-methoxyphenol via substitutions of new orthoquinones, 3993

fredericamycin, biomimetic model for construction of spirocyclic diketone center, 3063

galbonolides A and B, nonglycosidic macrolides from Streptomyces galbus, 6167 hexahydrobenzofuran fragment of

milbemycins and avermectins, model study for synthesis, 5759 key tricyclic lactam intermediate

and total synthesis of (±)-naphthyridinomycin, 1907 kinamycin D, C-assignments, long-

range heteronuclear COSY experiment, 4019

kinamycin, biosynthesis, 4023 lactam, intermediates via stereoselective aldol condensation of protected glycine and 3-butyryl thiazolidine-2-thiones with chiral aldehydes to give βhydroxyesters, 977

methylenomycin B, total synthesis, 6397

mitomycin C, new chemistry, 3923 N-formyl, new formylation reagent, 4-formyl-2-methyl-1,3,4thiadiazolin-5-thione, 3703

naphthacenequinone SS-228R, synthesis, 3887

nargenicin studies, dehydrative rerarrangement of a dihydrooxyoctahydronaphthalene, 5845

neocarzinostatin, with bicyclo-[7.3.0]dodecadiyne structure, 331

neooxazolomycin, structure, 1077 neplanocin and related cyclopententenyl-containing

nucleosides, synthesis, 3669 new 3'-deamino-3'-hydroxy-daunorubicin analogs, synthesis and rate of glycoside elimination under electro-reduction, 2641

new C, -unit substitution at C-4 of 4-acetoxyazetidinone deriv, synthesis of dethiathienamycin,

new halogenated marine prostanoids, antitumour chlorovulones from Clavularia viridis, 5787

new isotetracenone, capoamycin structure, 3471

new isotetracenone, kerriamycins A,B and C structures, 3475 new macrolide-polyether,

sorangicin A, structure and isolation from Sorangium cellulosum,6031

new, chromoxymycin, structure,

northern hemisphere of milbemycins, synthesis via formation of C-15 to C-16 bond, 5837

oxazolidone reaction with Nbenzylimines, asymmetric induction to evcloadducts and subsequent reduction to homochiral B-lactam derivs, 3783 oxazolomycin, β-lactone structure,

1073 patulolide A, new macrolide from Penicillium urticae. structure,

phosphorylating agent, S-4-methyl-phenyl-0,0-bis[1-benzo-

rebeccamycin, two syntheteic approaches, 4015

regiospecific and stereoselective conversion of ribonucleosides to 3'-deoxynucleosides; cordycepin from adenosine, 4295 right fragment of salinomycin,

stereocontrolled synthesis, chelation controlled Grignard reaction, 1541

route to 3-amino-2,3,6-trideoxy-D-or L-hexoses via nitroaldol reaction between 0-benzyl D-or L-lactaldehyde and alkyl 3nitropropionates, 1261 simple route to 4-demethoxy-7-

deoxydaunomycinone, 4703 stereo-and regio-controlled

synthesis of racemic and optically active (-)blastmycinone, 2815 stereoisomers of 7-hydroxyethyl-

2,2,5-trimethyl-3-oxa-1-azabicyclo[4.2.0]octan-8-one, structure/activity of 1-methyl carbapenems, 583 stereoselective construction of C-

ring fragment of trichothecenes from 4-cumyloxy-2-cyclopentenol, 3765

stereoselective synthesis of C(1)-C(9) and C(11)-C(17) fragments of protomycinolide 1V based on asymmetric pinacol-type

rearrangement, 3711 substituted 4-(bromomethyl)-Nhydroxy β-lactams, synthesis, 5385

swinholide-A, new macrolide from Theonella swinhoei, structure,

tetracyclic compds of potential value as anthracyclinone precursors, 111

tetrahydropyran "left wing" of ionophore X-14547A via stereo-selective 1,2-carbonyl addition and oxapyranone-dihydropyran rearrangement, 1163

thienamycin and epithienamycin C,

enantiomeric synthesis of intermediates, 5493 thienamycin synthesis, improved entry to key intermediate from methyl (R)-3-hydroxybutenoate via direct epimerisation at C-3 on 2-azetidinone rings, 4647

# Antimony and compounds

carbonyl compds, selective reduction with diphenylstibine,

cycloaddition of oxetane and CO2 catalysed by Ph.SbI, formation of trimethylene carbonate, 1323

debromination of phenacyl and benzylic bromides with tertiary stibine, 3211

### Aromaticity

some cyclic C.H. isomers and their potential trefoil aromaticity, 1585

# Aroylation

Ru-catalysed oxidation of cyanohydrins to acyl cyanides, reagents for selective N-

benzoylation of aminoalcohols.

### Arsenic and compounds

formylolefination of aldehydes by means of arsonium salts, 6447

Aizoaceae and Amaryllidaceae alkloids, synthons for construction, use of C-arylation

by aryllead triacetates, 5331 intramolecular-, C-, of 2-0-benzylated cyclic sugar derivs, useful 1,2-cis-C-gylcosylation, 2055

Pd-catalysed-, of vinyltrimethylsilane in presence of AgNO,, 3131

regioselective benzylation of lithium enolates derived from 2heptanone, 1599

### Association

n vs  $\pi$  competition in phenolsalkyl ketones association is a function of phenols acidity and branching of CO group, 3331

# Asymmetric induction and synthesis

- 3-methyl-2-cyclohexenone, photo-addition to optically active enoates, asymmetric induction and absolute configuration of
- adduct, 6163 α-tocopherol side chain, synthesis, 5153
- alkylation and sulphenylation of chiral O-silylated imide
- enolates, 5339 alkylation of chiral dipole stabilised anions, mechanistic
- model, 3295 allylic esters, asymmetric coupling with arylmagnesium-bromide, 3259 anthracyclinones, optically active, asymmetric synthesis,
- chiral 2-[(tributy1stannyl)methyl]propenamides, reaction with aldehydes via 1,6asymmetric induction to give  $\alpha$ -methylene-Y-butyrolactones, 1337

chiral a-aminonitriles from new Ncyanomethyl-1,3-oxazolidine synthon, 3567

chiral intermediates for chiral intermediates for prostanoids, asymmetric synthesis of cis-4-t-butyl-dimethylsiloxy-2-cyclopenten-1-ol and cis-4-t-tetrahydropyranyl-oxy-2-cyclopenten-1-ol, 5803 conjugate addition of RCu.BF, to encates of sulphonamide-shielded alcohols, synthesis of ß-substituted alcanoic acids, 6051 cycloaddition of diphlorycetene

cycloaddition of dichloroketene with chiral enol ethers, approach to optically active cyclopentenone derivs, 5525

diastereofaceselection, condensation of aldehydes with Zr enclate of chiral Npropionyl-trans-2,5-disub-stituted pyrrolidine, 5807

diastereoselection in Rh-mediated intramolecular C-H insertion, prep of a trans-3,4-dialkyl-cyclopentane, 3059 diastereoselective coupling

reactions of a ketene acetal, synthesis of  $R-(+)-\alpha$ -lipoic acid, 2535

dictyopterene B and its enantiomer, syntheses via

asymmetric S<sub>ON</sub> reactions, 3319
Diels-Alder reaction between
chiral isoprene units 2-(R)- and
(S)-benzyloxy-2,5-dihydro-4-

furancarboxyaldehyde and cyclopentadiene, 5605

Diels-Alder reactions of o-quino-dimethanes, stereocontrol exerted by phenyl substituents, asymmetric induction effects,

dihydroxylations via chiral

oxazolidines, 5459 effect of metal hydride and silica gel on Sharpless asymmetric epoxidation, 6221

enantio-and stereo-controlled synthesis of (+)-and (-)-eldanolide based on asymmetric pinacol-type rearrangement, 861 enantiospecific synthesis of (-)-

megaphone by controlled consecutive 1,4-and 1,3-asymmetric induction, 903

epoxidation of t-butyl substituted allylic alcohols, 2543

exo-allylic alcohol, stereospecific synthesis, and synthesis of (R)-(-)-2-acetyl-5,8-dimetrhoxy-1,2,3,4-tetra-hydro-2-naphthol, 6497

high induction during reduction of pyruvate and phenylglyoxylate esters of chromium complexed

carbinol, 429
higher induction by reduction of β-ketosulphoxides and transformation into optically active epoxides, prep of \$-hydroxysulphoxides of opposite stereochemistry, 435 indole alkaloids, synthesis, 3291 intramolecular Diels-Alder

reactions of N-acyl-camphorsultam trienes, 5437

Lewis acid mediated 1,2-asymmetric 1k-induction to α-alkoxy-aldehyde, synthesis of α-non-substituted β,Υ,syn-dihydroxy-ketone and ester, 3467 micobial metabolite, (+)-

citreoviral, enantioselective synthesis using hydroxylation of tiglate esters, 6485 new approach to 11-deoxy-prostanoids, 5009

optically active allylsilanes by hydrosilylation of 1-arylbutadienes, 3023

oxazolidone reaction with Nbenzylimines, asymmetric induction to cycloadducts and subsequent reduction to homochiral β-lactam derivs, 3783

oxidation of 3-phenylpropionates of chiral alcohols derived from (+)-camphor by MoO, Py.HMPT, 203 reduction of 2-ethoxycarbonyl-4,4-

thylenedioxycyclohexanone to ethyl (18,28)-4,4-ethylenedioxy-2-hydroxycyclohexane 451 (R)- and (S)-citramalate synthesis in high enantiomeric purity,

regio- and stereo-controlled synthesis of octahydro-naphthalenol portion of dihydrocompactin, 1131

(S)-1-acetoxy-2-aryloxypropionitriles, synthesis by asymmetric hydrolysis of the racemates with an enzyme, synthesis of (S)-propanolol,

5533 (+)(S)-3-trichloromethylbutyric acid via stereoselective conjugate addition of Cl.CMgCl to crotonate of chiral auxiliary with sulphonylamino substituent,

sesquiterpenoid botrydial, synthesis of quaternary Ccentres, 5433 spatane diterpene, synthesis of stoechospermol via intramolecular(2+2)-photocycloaddition, 3035

stereoselective construction of Cring fragment of trichothecenes from 4-cumyloxy-2-cyclopentenol, 3765

stereospecific synthesis of Ybutyrolactones from acyclic vinyl sulphoxides, synthesis of optically pure oak lactones,

synthesis of (R) or (S)-4,4dialkyl-2-cyclopentenones, 2047 synthesis of (S)-methyl-3-hydroxy-alkanoates from ketene and 2,2dichloroaldehydes via 4-(1,1dichloroalkyl)-2-oxetanones,

synthesis of α-hydroxy acids by alkylation of chiral N-(benzyl-oxyacetyl)-trans 2,5-bis-(methoxymethoxymethyl)-pyrrolidine, 1343

synthesis of a-hydroxy carboxylic acids, direct oxidation of chiral amide enclates using 2 sulphonyloxaziridines, 3539 tandem alkylation of α-alkylidene-

Y-butyrolactone derivs, 3027

### Autoxidation

alkylbenzenes, catalysed by cerium(1V)ammonium nitrate, 3353

1,1-diphenyl-3-arylisoquinolin-4ones new synthesis via cyclisation of 2-azabuta-1.3-

dienes, 5213
2-aza-1,3-diene prep via N-(1-triethylsilylallyl) protodesilylation, mechanism, 47 2-azidocarbapenems, prep and

reactions, 5407 2-ethoxy-6,11-dimethyl-7,9-bisdehydroaza[14]annulene and benzannelated derivs, synthesis, 635

2-ethoxy-6,11-dimethyl-7,9-bisdehydroaza[16]annulene and 2ethoxy-8,13-dimethyl-9,11-bis-dehydroaza[18]annulene, synthesis from annulenones, 639 4-aza-1,6,8-trienes, intra-

molecular Diels-Alder cyclisation, ratio of cis vs trans isomers, effects of solvent, steric bulk and nature of electron demand, 3575 aza-β-lactams via rhodium

carbenoid mediated cyclisation, 3171

carcinogenic aza-arene dihydro diols and diol epoxides, novel synthesis of trans-tetrahydro diol diacetates as precursors, 6417

polyalkoxy-2-aza-propenylium salts, synthesis via oxygen-alkylation of alkoxy substituted N-methylene amides, stereochemistry, 3547 regioselectivity in photo ring contraction of 4-diazo-

pyrrazolidine-3,5-diones to azaβ-lactams, 3167

# Azetidines

(3S,4R)-3-[(R)--1-hydroxyethy1]-4acetyloxy-azetidin-2-one, stereocontrolled total synthesis of chiral building block, 937 4-iodomethylazetidin-2-one

reaction with t-butyl ester of propiolic acid, formation of β-lactam ynamide, 4141 acetoxyazetidinones, reaction with trimethylsilylacetyl thiol-

esters, prep of azetidinone-thiolester precursors to

carbapenems, 573 azetidinone, introduction of alkyl, aryl, allyl and alkenyl groups at 4-position, 547

chiral 3-isocyanoazetidinone. stereoselective reduction, prep of optically active

carpetimycins, 223 chiral bisazetidines via hydro-alane reduction of bis-βlactams, 2035

ethyl 3-hydroxybutyrates and Narylaldimines cycloaddition to series of 3-(1'-hydroxyethy1)-2azetidinones, stereochemistry,

N-hydroxyazetidine, precursor of β-lactam, synthesis, 2809 new C<sub>3</sub>-unit substitution at C-4 of

4-acetoxyazetidinone deriv, synthesis of dethiathienamycin,

precursor of thienamycin and related carbapenems, cyclisation of N-propargyl epoxyamide to acetylenic 2-azetidinone, 4521

thienamycin synthesis, improved entry to key intermediate from methyl (R)-3-hydroxybutenoate via direct epimerisation at C-3 on 2-azetidinone rings, 4647

trapping of 1-azetin-4-one with siloxydienes as cyclocondensation adducts, 6309

2,6-dimethylphenyl azide, photolysis in N2 at 12K gives 2,6-dimethylphenylnitrene; in CO, isocyanate or azacycloheptatetraene are formed, 359

amination, using tetracarbonyl-hydridoferrate a selective reducing agent, 3277

fused pyrrolines via intramolecular cycloadditions of

pyrrolizidine alkaloids, 3527 phenyl azide, temp-dependent photochemistry in diethylamine, 2147

phenyl azides, 1,3-dipolar cyclo-addition to methyl 3pyrrolidinoacrylate, kinetics and mechanism, 4661

sodium azide, reaction with 9-bromo-6,7,8,9-tetrahydro-4H-pyrido[1.2a]pyrimidin-4-ones to give 9-amino-6,7-dihydro analogs, 3521

Azido acids and esters conjugate bases of 1,2,3,4tetrahydro-9H-carbazole and other indole derivs, action of t-butyl azidoformate, 2017

### Aziridines

1-diphenylphosphinyl-2,2-dimethylaziridine, new synthesis of  $\alpha,\alpha$ dimethylarylalkylamines, 1245

BF, -etherate ring opening of substituted aziridines by organocopper reagents leading to primary and sec amines, 1153 esters and amides from aziridine

2-carboxylic acid salts, 4439 N-tosyl-2-(1,3-butadienyl)-aziridine Pd-promoted transformation into N-tosy1-2vinyl-3-pyrroline, 857

2H-azirines, alkylation and silylation, 2637 new bi-2H-azirin-3-yl compds from 2,3-diazido-1,3-butadienes, 5261 spiro- and ring-fused azirines, conformational analysis, 4399

### Azocines

aromatic aldehydes, direct conversion to nitriles via S.Sdimethylsulphurdiimide iminating agent, dithiatetrazocines as intermediates, 2739

Azo and Azoxy compds 2,2'-hydrazonaphthalene, enantioselective reduction, 119 2-methylphenylazo compds conversion to diareno-1,2-

diazepines, 1345 azocyclobutane and azocyclopropyl-

methane, generation of C.H. radicals, comparison of PE spectra with data from pyrolysis

of methylnitrites, 2427 formation of 1,2,4-triazoles from oxidative addition of 1,4-diphenylazomethane and aryl aldehyde phenylhydrazones to

nitriles, 5655
vicinal '3C-13C coupling constants
for series of bi(poly)cyclic
azo-,azoxy- and imino compds for N=N-, N=NO- and N=C, 4427

diastereoselectivity in 1,2-addition of silylazoles to chiral aldehydes, stereocontrolled homolgation of ahydroxyaldehydes, 5477

### Azomethines

pyrazolid-3-one azomethinimines and (E)-β-nitrostyrene, 1,3dipolar cycloaddition, generation of seemingly "non cisoid" adducts, 5123

10-methyl-4-ketohydroazulenes synthesis using organotin, (3+2)-cyclopentane annulations.

anti-[2.2](1,6)azulenophane via fluoride induced 1,8-elimination from trimethylsilyl-tetraalkylammonium salts, 4835 cycloaddition of tetrachloro-

thiophene-S,S-dioxide, 2607 Ni-catalysed coupling of bromides of 1,5-methano[10]annulene and azulene, synthesis of biannulene and bi-, ter-, quater- and poly-azulenes, 3829

stable diatropic 1,6-and 2,6-di-t-butyl-cyclopent[f]azulen-ides, dominated by polymethide conjugation, prep, 2567

# Benzene derivatives

alkylation of aromatic compds with Pummerer rearrangement intermediates, prep of methylaryl compds via methylthiomethylaryl products, 2787

alkylbenzenes, photo-autoxidation catalysed by cerium(1V)ammonium

nitrate, 3353 benzene, novel compexation with cyclophane, 1325

dilithiated species generated by metal-halogen exchange and ortho metallation, reaction with electrophiles to give polysubstituted aromatics, 1145 ethyl benzene and derivs, fungal

hydroxylation, 6409 phenylindenes, electro-oxidation

depends on electrolyte anion and non-nucleophilic solvents, 1265 spiro hydrindandione ring system

of Fredericamycin A via spiroalkylation of indene, 4725 stable 3-methylene-1,4-cyclo-

hexadiene, a non aromatic tautomer of benzene, 4715 t-butyldimethylsiloxy-3,5-dimethoxy-benzene, regioselective lithiation, 123

toluene-α-d and-α-t in aqueous NaOH, measurements of H-isotope exchange, 6317

### Benzodiazepines

1,4-benzodiazepines, one-step synthesis, application to synthesis of neothramycin, 5947

### Benzofurans

hexahydrobenzofuran fragment of milbemycins and avermectins,

model study for synthesis, 5759 precursor of 4,5-dimethylbenzo[b]-furan from Desmia hornemanni, isolation of 2-(1-enloro-2hydroxyethyl)-4,4-dimethylcyclohexa-2,5-dienone, 2335 tetrahydrodibenzofurans, new

route, 3449 triacetoxybenzofuran, conversion to khellinone, 1385

### Benzof urazans

4,6-dinitro-benzofuroxans and benzofurazans new 7-substituted, by H-displacement in 4,6-dinitro-benzofuroxan andbenzofurazan, 1307

### Benzofuroxans

4,6-dinitro-benzofuroxans and benzofurazans, new 7-substituted, by H-displacement in 4,6-dinitro-benzofuroxan and benzofuran, 1307

# Benzoisothiazoles

2,3-dihydro-2,3-disubstituted-1,4benzothiazines from 2-aryl-2,3-umydrobenzothiazotes, (45)

3-methyl-2,3-dihydrobenzothiazoles, use in dye enhanced photoreductions of halides, sulphonium salts and pyridinium salts, 1453

### Benzophenoxazines

benzo[a]phenoxazine derivs, formation by triethylphosphite deoxygenation of 1-(2-nitrophenoxy) and 2-(2-nitrophenoxy)naphthalenes, 393

# Benzopyrans

substituted derivs via tributyltin hydride induced heteroring annelation of o-iodoaryl allyl ethers derived via aromatic directed metallation, 6001

## Benzoquinones

benzoquinone/porphyrin/benzoquinone molecule, structure and fluorescence, 6055 new benzoquinone-bridged porphyrin, 6059

### Benzotriazines

4-methylene-1,2,3-benzotriazines, pyrolysis, 941

# Benzotriazoles

new coupling reagent, benzotriazol-1-yl diethyl phosphate for amide and peptide synthesis, 1341

### Biogenesis

Br-containing diterpenoids, (±)-concinndiol and (±)-aplysin 20, synthesis, 343 fredericamycin, biomimetic model

for construction of spirocyclic diketone center, 3063 indolosesquiterpenes isopolyalthenol and neopolyalthenol

from Polyalthia sauveolens, biogenesis, 4937 janussines A and B from Strychnos

johnsonii, cross point in biogenesis of quasidimeric alkaloids, 2441

taxane alkaloids putative biogenetic precursor verticillene, total synthesis,

### Biologically active compounds see also Drugs

- 2'-5' linked oligoribonucleotides, synthesis, 761
  3-deazooxanosines. C-nucleoside
- isosteres of oxanosine, synthesis, 5785
- (±)-aphidicolin, total synthesis, 6147
- (±)-indolactam fraction of teleocidin B, conformational isomerism, 1069
- (S)-1-acetoxy-2-aryloxypropionitrile, synthesis by asymmetric hydrolysis of the racemates with an enzyme, synthesis of (S)-propanolol, 5533
  (+)-pederin, total synthesis,
- stereocontrolled synthesis of
- (+)-benzoylpedamide, 6461 absolute stereochemistry of panaxynol by exciton chirality method, 5775
- alkaloid necatorone, synthesis, 5975
- ancovenin, new inhibitor of angiotensin 1 converting enzyme, structure, 665
- antitumour and antimicrobial flavonoid, uvaretin, synthesis,
- ary1[2.3-a]quinolizin-2-ones,
- synthesis, 1277 azadirachtin and 22,23-dihydro-238-methoxyazadirachtin, structures, 6435
- base catalysed formation of Nalkyltriazenes from N-hydroxy-methyl-N-alkyltriazenes, 1557 benzyl6-β-aminopenicillinate
- conversion into crystalline 6-aformido deriv without
- chromatography, 377 bioactive phenyltetrahydrofurano-2-pyrone, goniothalenol, characterisation and isolation from Goniothalamus giganteus,
- carcinogenic aza-arene dihydro diols and diol epoxides, novel synthesis of <u>trans</u>-tetrahydro diol diacetates as precursors, 6417
- cyclic pentapeptide analogs of thymopentin, NMR, 177 cyclopentenonyl fatty acids, synthesis of (±)-dicranenones,
- cyclopropyl analogs of precocene I, synthesis and activity against bean aphid, 4677
- dipole measurements of 7 polar amidine containing compds of biological interest, 1897 diterpenes, (+)-spatol with natural configuration, total
- synthesis, 6109 diterpenoids, identity of coleonol and forskolin via structure
- revision, 551
  free stalic acid by aldolcondensation of 2-acetamido-6-0acetyl-2-deoxy-D-mannose with pyruvic acid in presence of immobilised acylneuraminate pyruvate-lyase, 2439

- histamine liberating thansigargin and trilobolide, absolute configurations, 107 isoquinuclidine ring system, prep
- via modified Polonovski reaction, 801 key chiral intermediate for
- key chiral Intermediate for mevinolin and compactin, prep of ethyl 5(S),6-epoxy-3(R)-(methoxymethoxy)hexanoate, 1185 leukotriene D., prep of diketopiperazine analog, 1951 lipopolysaccharides from Aeromonas
- salamonicida, synthesis of oligosaccharide sequences with repeating unit of O-specific chain, 6043

- chain, 6043
  lipoxin A, synthesis and
  stereochemistry, 281
  lipoxin B, synthesis and
  structure, 1919
  lyngbyatoxin A and teleocidin B,
  prep by application of new
  indole synthesis, 4047
  N-acetyl-N-oxo-1,4-benzoquinone
- imine, an acyl nitrone, prep and intramolecular rearrangement to N-acetoxy-1,4-benzoquinone imine, 2467 N-alkylthiazolium halides,
- including thiamine, reaction with KO<sub>2</sub>, chemistry and biological implications, 2917
- new isotetracenone antibiotic capoamycin, structure, 3471 new isotetracenone antibiotics kerriamycins A,B and C, structures, 3475
- new stable analogs of prosta-cyclins,(±)-6a-oxo-6,9-methano-15-hydroxyprosta-5,13-dienoic acids, 4101
- new synthesis of 3.5-diamino-1,2,4,6-thiatriazine 1,1-
- dioxides, selective prep of 2-alkyl or 6-alkyl isomers, 1101 new synthesis of 4-alkyl-3,5-diamino-1,2,4,6-thiatriazine 1,1-dioxides, 1105
- n, 1-dioxides, 1105 new synthesis of a,a-dimethylaryl-alkylamines from 1-diphenyl-phosphinyl-2,2-dimethyl-aziridine, 1249 novel 3-oxocyclopenten-2-
- phosphonate deriv, intermediate
- pnospnonace deriv, intermediace for 2-alkyl substituted cyclo-pentenones, 93 phidolopin from <u>Phidolopora</u> <u>pacifica</u>, synthesis of 7-(4-hydroxy-3-nitrobenzyl)-1,3dimethylxanthine, 2355
- platelet activating factor via cyclic tin intermediate, 1379 platelet activating factor, stereocontrolled synthesis from (S)-(-)-malic acid, 5195
- quinonoid 6-methyl- and 6,7dimethyldihydropterins, synthesis, 4003
- regio-and stereo-selective alkylation at 3'-terminal end of ribonucleotides by N-2-methyl-9-hydroxyellipticinium acetate, 2891
- regiocontrolled H-D exchange of biologically important indoles under UV, 5891
- route to 3-amino-2,3,6-trideoxy-Dor L-hexoses via nitroaldol reaction between O-benzyl D-or L-lactaldehyde and alkyl 3-nitropropionates, 1261
- selective inhibition of 5-lipoxygenase by 1,1-disubstituted 2,5-cyclohexadienes, 2039
- serotonine and bufotenine derivs as potential electrophilic or photoactivable labels for serotoninergic sites, azidoethyl-3 indole deriv most promising compd, 4443

- synthesis of 5-ylidenepyrrole-2one, pukeleimide, from Lyngbya
- majuscula, 3617 synthesis of riboside analog of pyrimidine adduct from Nhydroxy-N-2-acetylaminofluorene and DNA, 2659
- trifluoromethylated diene derivs, Claisen rearrangement of tri-fluoromethylated propargylic and allylic alcohols, 219 trihydroxy derivs of adenosine,
- guanosine, uridine and cytidine, prep using periodate and borohydride supporting resins, 1305

### Bioluminescence

Cypridina biluciferyl, activity, structure, 239

### Biosynthesis

- 5-enolpyruvyl-shikimate-3phosphate, chiral synthesis, 21 aldosterone inhibition by 116hydroxy-18-ethylprogesterone, 1137
- and structure of lipoxin B, 3939 biomimetic route to clavicipitic acids, intermediate in ergot biosynthesis, 4043
- columbamine immediate precursor of
- berberine, 201
  diterpene cystoseirol of mixed
  biosynthesis from Cystoseira
  mediterrranea, 4919 enzyme prep from E.coli in
- formation of o-succinylbenzoic acid from iso-chorismic acid, key in vitamin K<sub>2</sub> biosynthesis, 1487
- gram-negative bacterial lipopolysaccharide, investigation, 677 identification of new eocosanoid from in vitro experiments with Clavularia viridis, biosynthesis of clavulones, 4171
- or clavulones, 41/1 intermediate in menaquinone, 1703 isolation of  $\Delta^{\circ}(^{11})$ -sterols from Psolus fabricii, holothurin biosynthesis, 3513 kinamycin, 4023
- leukotriene, methylene cyclopropane analogs of arachidonic acid, potential mechanism-based inhibitors, 1973 presqualene pyrophosphate-
- squalene, synthesis of bifarnesol, 3303
- terrecyclic acid A, confirmation of cyclisation mechanism and H-shifts using [2-24,] acetate and [2-13C2H,]acetate, 3845

## Bismuth and compounds

Bi-mediated allylation of aldehydes to homoallylic alcohols, 4211

# Bonds see also H-bonds

- 50,80-peroxides, bisfragmentation of C(5)-C(10) and C(8)-C(9) bonds and elimination of AcOH to form new bicyclic 5,10;8,9-
- diseco-steroid derivs, 389 alkyllithium, β C-H bond, hydride abstraction and regioselective insertion by ethyl(phenylthio)-
- methylene carbenoid, 6489 aryl-aryl bonds, construction, 29 northern hemisphere of milbemycins, synthesis via formation of C-15 to C-16 bond, 5837

# Boron and compounds

1,2-dimethylcyclopentene, hydroboration, stereochemistry, 6125 (E)-vinylboronic acids, chlorination to (Z)-vinyl chlorides, 279 a-alkoxy acetylenic ketones

stereoselective reduction with zinc borohydride and Kselectride, 5139

ally1, benzy1 and tertiary alcohols conversion into halides with BF,-etherate/halide ion, 3863

amine-and phosphine-boryl radicals to 2-methyl-2-nitrosopropane, 2,4,6-tri-t-butylnitrosobenzene and phenyl-N-t-butylnitrone to nitroxide spin-adducts, 1349

nitroxide spin-adducts, 1349 anions of alkyldimesitylboranes, reaction with oxiranes followed by oxidation to 1,3-diols, 5097

aryl aldehydes, Wittig reaction with carbanions stabilised with adjacent dimesitylboron group, oxidation of intermediates to erythro-1,2-diols, 5093

BF<sub>3</sub>-etherate on alumina catalysis in improved synthesis of cannabidiol, 1083

BF,-etherate ring opening of substituted aziridines by organocopper reagents leading to primary and see amines, 1153

catechol boron halides, cleavage of ether, ester and carbamate protecting groups, 1411

dialkylketones, synthesis from alkenes via hydroborationcarbenoidation, 6361 diborane approach to synthesis of

polyquinanes, 4863 dimethyl(methylthio)sulphonium fluoroborate, sulphenetherification and sulphen-

lactonisation, 6159 dithia[6]radialenes, formation by boron trifluoride-catalysis of "thiiranoradialene", 3019 enantioselective aldol reaction of

enantioselective aldol reaction of chiral acyl thiazolidine thione derived from boron enolates, 4855

enol borates, addition to aldehydes and subsequent metal addition to form 1,3-diols, 1643

haloboration of 1-alkynes, 1065 homogeneous deblocking with 2bromo-1,3,2-benzodioxaborole, 1115

hydroboration of β-acetylenic alcohols followed by oxidation to hemiacetals of Y-aldois and dehydration to 2,3-dihydrofurans, 5683 hydroboration of olefins with

hydroboration of olefins with acetoxyborohydride in presence of reducible functional groups, 1757

iodide ion promotion of benzyl
 chloride-borate ester
 carbonylation, 2273
ionic reactions of N-halo

ionic reactions of N-halo electrophiles with alkenes promoted by BF<sub>3</sub> etherate, 1811

isoprenylation, new reagent 2-dipropylborylmethyl-1,3-butadiene, synthesis of ipsenol and ipsdienol, 2797 Pd-catalysed allylation of lithium

Pd-catalysed allylation of lithium 3-alkenyl-1-cyclopentenolatestriethylborane and application to selective synthesis of methyl (Z)-jasmonate, 2177 pinacol allylboronate,

stereochemistry of reactions with α,β-dialkoxyaldehydes, comparison with reactions of crotylboronates, 3427

potassium triethylborohydride reduction of 2-alkyl-3-oxo 6162

synthesis of 2-substituted methylenecyclopentanes and cyclohexanes, reaction of carbonyl compds and alkoxyacetylenes with 1-(dialkylboryl-methyl)cyclopentene and cyclohexene, 4551

triorganylboranes from organic halides by modified organometallic route using ultrasonics, 4311 unsymmetrical biaryls obtained by

unsymmetrical biaryls obtained by cross coupling of directed metallation-derived arylboronic acid, reaction with aryl

halides, 5997 vinyl ethers, stereoselective hydroboration, 6297

### Carbanions

3,4-epoxy-2-methylenealkanoic acid derivs synthesis,using dianion of N-phenyl-2-(phenylsulphonylmethyl)propenamide and aldehydes, 4751

anion derived from 2-benzenesulphonyltetrahydropyran, reaction with electrophiles to give alkylated and acylated products, 535

anion of 2-nitropropane and 3bromo-1-nitrocyclohex-1-ene, SRN1 substitition, 539

cationic dihexadecyldimethylammonium bromide, permeation of Ellman's anion from surface to interior, 603

chiral nitronate dianion from (+)-D-gyloeraldehyde, generation and use in synthesis of 3-amino-2,3dideoxy sugars, 6269

cyclononatetraenyl dianion, reactions with electrophiles, geometry and conformation of 1,3,6-cyclononatrienes, 5351

dianion metallation of N,Ndimethylvanillylamine and N,Ndimethylisovanillylamine, 2151

dianion of α-cyanoacetone,
 reaction with lactim ethers to
 give bicyclic 2-amino-4(1H)-

pyridones, 259
dianion of N-phenyl-2-(phenylsulphonylmethyl)propenamide,
reagent for synthesis of (E)α,8-unsaturated amides and 5,6-

dihydro-2H-pyrans, 4747 diastereospecific aprotic conjugate additions of allylic anions bearing polar, chargestabilising groups, 1569

efficient propargylic anion equivalent formation for coupling with alkyl and allyl halides, 271

formamidines as α-amino carbanion precursors, synthesis of 2-arylpiperidines, -pyrrolidines and nicotine analogs, 5863

nicotine analogs, 5863 Horner-Wittig reaction with 4hydroxy-2-alkenylphenylphosphine oxides give dianions and hence 3-hydroxydienes, 249

Michael reactions promoted by solid-liquid phase transfer catalysis without solvent, application to addition of acetylacetone, methylaceto-acetate and fluorene anions on hindered acceptors, 4601 new organometallic complex, a

methane anion to form nitromethyl derivs, conversion to nitroaromatic aldehydes, 3599

nitronate anions, of 3-ethyl-5nydroxymethyl-5-nitrotetrahydro-1,3-oxazine, nucleophiles for SRN1 reactions, prep of new tetrahydro-1,3-oxazines with trisubstituted ethylenic double bond in 5-position, 323 optically pure sulphinyl

optically pure sulphinyl carbanions, reaction with α-halocarboxylic esters, 5601

p-toluenethiolate anion, and N-tosyldiarylsulphilimines, irradiation, S-N bond cleavage, 775

phosphorus acyl chlorides phosphonylation, generation of alkylidene-diphosphonate anions convertible to vinylphosphonates, 4435

photooxygenation of nitroaromatic compds involving nitrobenzyl carbanions, 2387

reaction with epoxides derived from tartaric acid. 3121

from tartaric acid, 3121 selective Y-alkylation of triisopropylsilylallyl anion, synthesis of a-triisopropylsilyl aldehydes, 5375

selective methylenation of aldehydes with methylene dianion synthon, 5581 stable radical-carbanion with

stable radical-carbanion with independent radical and anion sites, rapid spin charge exchange, 5589

sterically congested cyclononatetraenyl anion, temp-dependent electrocyclisation to tetracyclic frames, 1695

substituent effect on reverse Menschutkin reaction of 1methylpyridinium cations with iodide anion, 5065

surfactant vesicles, charge type and trans-membrane transport of fluorescent probes, 4827

fluorescent probes, 4827 tertiary nitronate anions, reaction with 1-methtl-2-chloromethyl-5-nitroimidazole to yield 1-methyl-5-nitroimidazoles with trisubstituted ethylenic double bond in 2-position, 1023

tertiary-, steric and electronic limitations of S<sub>RN</sub> reaction with p-nitrobenzylic substrates, whose

tetra anion of acepleiadylene and dianion of pyrene, formation in

, electrochemical reduction, 415 thiocarbonyl compds, desulphurisation to oxo-deriv via peroxy-S intermediate from 2-nitroberznenesulphonyl chloride and superoxide anion, 1079

vinyl selenide anions in stereospecific synthesis of vinyl alkyl selenides from vinyl halides, 2225

with captodative substitution, oxidation to dimers, 4607

### Carbazoles

1-oxygenated-, new route to, synthesis of murrayafoline-A, 5841

6-ethyl-10-methoxy-7H-pyrido[4.3-c]-, new synthesis, 4929 intramolecular cycloaddition to

intramolecular cycloaddition to acetylenic vinylketenimines to carbazoles, synthesis of Nmethyltetrahydroellipticine,

### Carbenes

- alkyllithium, ß C-H bond, hydride abstraction and regioselective insertion by ethyl(phenylthio)methylene carbenoid, 6489
- aryloxymethylchlorocarbene, 1,2-H migration, effect of β-aryloxy group on stereochemistry, evidence for "negative" hyperconjugation of carbene lone
- pair, 1651 cyclic disulphides, desulphurisation and insertion reactions with carbenes, 5187
- dialkylketones, synthesis from alkenes via hydroborationcarbenoidation, 6361
- difluoro-, reaction with quadricyclane, 5403
- electrophilic-, addition to electron deficient olefin. kinetics of benzylchlorocarbenediethylfumarate reaction, 3071
- in situ protection in alkyne carbene cycloaddition, 2969 new route to homoaporphines, addition of dichlorocarbene to
- aporphinoids, 1561 persubstituted hexamethoxycyclopropane prep and pyrolysis to dimethoxycarbene and tetra-
- methoxyethylene, 1931 phenylthio-, regioselective intramolecular insertion C-H bond of alkoxides, 4483
- pyrrole derivs, competitive intramolecular carbenoid reactions, 6035
- pyrrole-and phenyl-, Cr-complexes, reaction with alkoxalkyne, dimer formation via o-quinonemethide and subsequent dimerisation, 1159
- reaction with bicyclo[2.1.0]pentane, 5399
- reaction with oxetane, generation of oxonium ylides, Stevens rearrangement and protonation to tetrahydrofurans and 1,3-
- dialkoxycyclopropanes, 193 Se stabilised carbenium ions, bis-(seleno) - and bis(thio) alkylation of ketones and
- aldehydes, 6513 singlet and triplet diphenylcarbene reaction with alcohols, effect of temp on reversible
- ylide formation, 5951 vinyl carbenoids, [3+4] cyclo-addition with furans, 5659

# Carbocations

- bridgehead, C-6-phenylselenyl bicyclo[4.2.2]piperazinediones hydroxylation to 6-peroxy and C-6-hydroxy bicyclic piperazinediones, 37
- NMR isotope shifts as probe of

- soln., 3251
- 2,2,5,5-tetramethylpyrrolidine-1oxyl and  $\beta$ -cyclodextrin, association, equilibrium constant, 741 2,3-dideoxy-3-nitro furanosides
- and pyranosides, enantiospecific
- synthesis, 6269 2,3:5,6-di-O-isopropylidene-Dmanno-furanose, chain extension with 2-lithio-1,3-dithiane to Dglycero-D-galacto-heptose, 3689
- 2,6-dideoxy-and 2,4,6-trideoxysugars via hetero Diels-Alder reaction, diastereoselective
- synthesis, 2065 2-deoxy-D-glucopyranosyl phenyl-sulphones, route to C-glycosides via reductive lithiation, 6185
- 2-0-benzylated cyclic sugar derivs, intramolecular Carylation, useful 1,2-cis-C-gylcosylation, 2055
- 3α-(1"β-D-glucopyranosiduronyl)-8α-isovaleryloxy-scirpen-3,48,15-triol 15-acetate produced in vitro from T-2 toxin, structure, 5231
- 4-0-benzy1-2,3-dideoxy-L-threohex-2-enono-1,5-lactone synthesis, use of L-tartaric acid, 3731
- (+)-castanospermine total
- synthesis from D-mannose, 4617 acetoxylation of 2,3-dihydro-4pyrones, route to glucal stereochemical series, 3411
- ally1-C-glycopyranosides using
- peracetylated glycopyranoses, stereoselective synthesis, 1479 anomeric free OH of protected aldoses and ketoses, reaction with (diethylamino) sulphur trifluoride to give glycosyl
- fluorides, 3 anomeric OH of furanose and pyranose hemiacetals, stereoselective replacement by F atom, 5
- antiherpetic agent, synthesis of  $(\underline{S})$ -9-(2,3-dihydroxy-1-propoxymethyl)guanine, 1815
- antitumour alkaloid, enantiomeric pair of unprotected AB-ring systems of sesbanimide A, synthesis from D- and L-xylose, 4639
- β-cyclodextrin deriv, synthesis, binds guest molecules in heptane because cavity is more polar than solvent, 267
- β-cyclodextrins, unsymmetrically disubstituted 6A,6B- and 6A, 6G-derivs, enzyme-based dsicrimination of clockwise and counterclockwise isomers, 6439
- $\beta\text{-D-glucosides}$  stereoselective synthesis using stannous

- D-glycopyranosyl phenyl sulphones, acylation of lithiated anions and subsequent formation of a-D-C-glucosides, 6193
- D-glycopyranosyl phenylsulphones, use in stereocontrolled synthesis of β-D-C-glycosides,
- deoxymannojirimycin, fagomine and 2R,5R-dihydroxymethyl-3R,4R-dihydroxypyrrolidine from Dglucose, enantiospecific synthesis and configuration,
- deprotection of alloxycarbonyl derivs of sugars in presence of Ir, Rh and Pd catalysts, 5045
- diastereoselective aldol reaction of 2,3-0-isopropylidene-D (and L)-glyceraldehydes leading to 2-deoxy-D (and L)-riboses, 5777
- exciton coupling, excimer emission and unique binding of Y-cyclo-dextrin substituted by two naphthyl residues, 3339
- flexibly capped β-cyclodextrins. prep and characterisation by Hbonded adenine-thymine, 1735
- free sialic acid by aldolcondensation of 2-acetamido-6-0-acetyl-2-deoxy-D-mannose with pyruvic acid in presence of immobilised acylneuraminate
- pyruvate-lyase, 2439 highly functionalised ribofuranose and ribopyranose derivs via iodine cyclisations of Dribohept-2-enonates, 1943
- levulinic acid formation, mechanism, 2111
- lipopolysaccharides from Aeromonas salamonicida, synthesis of oligosaccharide sequences with repeating unit of O-specific
- chain, 6043 methyl 2,4,6-tri-0-benzyl- $\alpha$ -Dglucopyranoside and methyl 3,4,6-tri-0-benzyl-6-D-manno-
- pyranoside, novel synthesis, 807 methyl 3-0-benzyl-2,4,6-trideoxymethyl 3-U-benzyl-c, +, -, - u. Lucus, 6-iodo-a-D-erythro-hexo-pyranoside in synthesis of HMG-CoA reductase inhibitor, 2947 methyl esters of 11S and 12S-HETE,
- synthesis via D-arabinose, 2993 mevinolin and compactin, new prep
- of optically active analogs in which  $\beta$ -hydroxy- $\Delta$ -lactone moiety is drived from D-glucose, 4995
- new synthesis of p-nitro-calix[6]arene, 3343
- novel glucoside of fagomine in Fagopyrum esculentum seeds, isolation and characterisation. 1465
- oligosaccharide structure of de-0acylated LPS from E coli Re mutant F 515, 6321

- stereocontrol in isoxazoline reductions, synthesis of aminodeoxy-DL-xylo-and arabinopentose derivs from furo-isoxazolines, 2997 stereospecific alkylation at C-6
- of pyranose sugar, conversion of glycosyl fluorides into C-
- glycosides, 1823 stereospecific allylation of "ribo" and "galacto" aldosulose derivs, 823

### Carbolines

1-alkyl-β-carbolines, treatment with dialkyl oxalate, synthesis of canthin alkaloid derivs, 385 4-oxo-tetrahydro-β-carboline, hydrazine mediated conversion into 4-amino-β-carboline, 2139 new entry into 4-substituted and 3,4-disubstituted B-carbolines.

### Carbonates

mixed carbonates, reagents for synthesis of active ester of

amino acid derivs, 1721 trimethylene carbonate, formation by cycloaddition of oxetane and CO<sub>2</sub> catalysed by Ph<sub>4</sub>SbI, , 1323

### Carbonium ions

- activated 1-aminopyrazolium cations, regioselective nucleophilic substitution and synthesis of 5-substituted 1-
- methylpyrazoles, 5485 alkylaromatic radical cations, reactivities of i-Pr, Et, and Me groups in gas phase side-chain deprotonation, 4269
- cation binding properties of 12-membered ring N-pivot lariat ethers, 151
- chiral a-siloxyketone, effect of cation on diastereofacial selectivity in aldol reactions,
- complexation by N-(3,6,9-trioxa-decyl)monoaza-12-crown-4 lariat ether, K, cation completely enveloped by donor groups in both ring and side arm, 4035
- crown ethers capped metalloporphyrins, complexes with metal cations, 3739
- doubly charged ions of bridged [4n]annulenes, evaluation of diatropic and current effects, 3091
- ESR characterisation of N.Ndimethyl-p-anisidinium dimer radical cation, 6369
- gem-dibromocyclopropanes, reaction with diphenylphosphide ion to form cyclopropyldiphenylphosphines, 105
- glycine cation equiv, electrophilic, prep and reaction with heteroatom nucleophiles, 695
- glycine cation equiv, reaction with carbon nucleophiles, amino acid synthesis, 699
- hybridisation and structure of aferrocenyl-β-t-butylvinyl cation via <sup>13</sup>C NMR, 1493 initiation of cyclisation with 4-
- substituted cyclohexenyl cations, 2921
- intramplecular reactions of acyclic N-acyliminium ions with propargyl silanes leading to derivs of 3-vinylidenepyrrolidine, 3-vinylidene-piperidine or 1-amino-2vinylidenecyclopentane, 3151 intramolecular reactions of
- acyclic N-acyliminium ions, allyl silanes as nucleophiles,

3155

- methoxides, reactions with (halomethyl)silanes at Si, facilitation by aprotic solvents and cation separation, 1115
- nitrenium ion not involved in production of hydroxamic acid. solvolysis of N-acetoxy-Nacetylaminobiphenyl, 147
- O-protonated acyl-substituted πcomplexes, stabilisation of acarbocationic centre, direct participation of transition
- metal, 3605 oligooxa[n.n]paracyclophane quin-
- hydrones, cation-induced CT absorptions, 6175 oxidation of electron rich compds to radical cations with ClO<sub>2</sub> in dicholoromethane, 1765
- quinone-imonium cation, synthesis using DMSO as methylating agent,
- quinonoid crown ethers, synthesis and cation binding properties,
- radical cation disproportionation in electron-transfer oxidation of dioxenes to a-diketones, 789
- radical cations of aromatic amines and heteroaromatics via action of nitroxides, 4201
- reversible 2-aza-Cope rearrangement in cyclisations of N-acyliminium ions derived from 1'-and 2'-vinyl-N-(3'-butenyl)-
- 5-hydroxy-2-pyrrolidinones, 5105 substituent effect on reverse Menschutkin reaction of 1methylpyridinium cations with iodide anion, 5065
- surfactant vesicles, charge type and trans-membrane transport of fluorescent probes, 4827
- tetra(1-adamantyl)cyclobutadiene radical cation, ESR, 4121 tetrakis(dimethylamino)-p-benzo-quinone, stable radical cation,
- thienium cation, formation from 8ethylthionitroolefin and AlCl,,
- triphenylphosphine radical cation, reaction with cycloalkenes, electro prep of 1-cycloalkenyltriphenylphosphonium salts, 2199

# Carbonylation

- (+)-2-carene, induced by ironpentacarbonyl, 1213
- allylic alcohols, Pd-catalysed oxidative cyclisation and carbonylation to 5-membered ring lactones, 5639
- cis 3-hydroxytetrahydrofuran acetic acid lactones, stereo-selective synthesis via Pdcatalysed oxycarbonylation of 4-
- penten-1,3-diols, 3207 cobalt carbonyl conversion of disulphides to thioesters and desulphurisation to sulphides, 2609
- Y, &-unsaturated carboxylic acid derivs, synthesis by Ni(CO),-induced ring opening carbonylation, 5795
- halides and ethers, Rh-catalysed, 5743
- of benzyl chloride-borate ester, iodide ion promotion, 2273 organo-Al compds, ketone synthesis
- via Pd-catalysis, 4819 silylamine or silylsulphide in Ni(CO),-induced carbonylation of gem-dibromocyclopropanes, 5061

### Carbonyl compounds 1,2-dicarbonyl compds, reaction with dimethyl 3-keto

- glutarate, steric and electronic effects, 2163
- 1,5-dicarbonyl compds, prep from enol ethers, methyl vinyl ketone and BF, etherate in presence of
- an alcohol, 6201 1-ethoxy-3-trimethylsilyl-1propyne regio- and stereoselective reaction with saturated and ethylenic carbonyl derivs giving new a-functional allylic, conjugated dienic or tienic silanes, 1861 α,β-unsaturated-, reaction with
- ethyl isocyanoacetate promoted by ZnCl2 and CuCl, 5781
- α,β-unsaturated nitroalkenes reduction by Cr(11)chloride, route to carbonyl compds, 3777
- a-hydroxylation of dicarbonyl compds as their silyl enol ethers by MCPBA, 3563 addition of 3-bromo-1,1,1-tri-
- chloroalkanes to give 3.3dichlorotetrahydrofurans, 4899
- β-dicarbonyl compds, novel method for isoprenylation, 6337 β-substituted, new synthesis, 657
- β-trimethylsilyl-α,β-unsaturated carbonyl compds, new synthesis using 1-methoxy-3-phenylthio-3-trimethylsilyl-1-propene, 2677
- benzyne-induced fragmentation of 1,3-oxathiolanes, deprotection of CO groups, prep of phenyl vinyl sulphides and 1,2-carbonyl
- transposition, 2195 carbonyl oxides, rearrangement to esters using 180, 3723 carbonyl participation in
- solvolysis of a-keto mesylates, cleavage of π-bonds with 4-t-butyl
- iodoxybenzene, 4955 conjugated-, addition of organocuprate-chlorotrimethylsilane
- reagents, 6019
  methylenecyclopentanes and cyclohexanes, 2-substituted, via reaction of carbonyl compds and alkoxyacetylenes with 1-(dialkyl-boryl-methyl)cyclopentene and -cyclohexene,
- mono- and di-compds prep by fluoride ion induced reactions
- of β-ketosilanes, 787 monocyclopentadienyl titanium alkyls thermal stability in reaction with carbonyl groups,
- ozonation of dimethyl ester of endo cis-bicyclo[2.2.1]hept-5-en-2,3-dicarbonic acid, intramolecular interaction of zwitterion and methoxycarbonyl group, 5843
- reaction with \$-phenylthioalkyltitanium reagent, 5313
- regiospecific nucleophile-induced fragmentation of α,α-dichloro-βiminocarbonyl compds, 2709
- selective reduction with diphenylstibine, 5171 selective transformation of N-t-
- butoxycarbonyl group into N-alkoxycarbonyl group via Ncarboxylate ion equivalent, 5543 stereochemistry of addition to Υ-alkoxy-α,β-unsaturated carbonyl systems, 6313
- systems, conversion to carbonyls by clay-supported ferric nitrate, 2327 unsaturated-, and butenolide via
- isomerisation of 1,3-diene monoepoxide and α-alkylidene-Ybutyrolactone, regio-and stereo-control, 1527

xviii Tetrahedron Letters

### Carboxylation

- electro, of organic halides and CO<sub>2</sub>, 1509
- homologation of ketones to α,βunsaturated carboxylic esters or amides via Pd-catalysed enol triflates, 1109
- reductive-, of sec amines, synthesis of tertiary Nmethylated amines, 5367

### Carboxylic acids anhydrides

- 11-deoxyanthracyclinones via cycloaddition of substituted tetrahydrohomophthalic anhydride, regiospecific synthesis, 1549 2-alkylated-3-deuteriated-1-amino-
- 2-alkylated-3-deuteriated-1-aminocyclopropane-1-carboxylic acids, regiospecific synthesis, 481
- 2-butanoic acid dienolate, addition to 1,3-diphenylpropenone, almost exclusive formation of R,R(S,R)-3,5diphenyl-2-ethenyl-5-oxopentanoic acid, 2485
- 3,4-epoxy-2-methylenealkanoic acid derivs synthesis using dianion of N-phenyl-2-(phenylsulphonyl-methyl)propenamide and aldehydes, 4751
- 3-(3,3-dimethyltriazen-1-yl)pyridine-4-carboxylic acid, precursor of 3,4-didehydronyridine prep 2123
- pyridine, prep, 2123 3-substituted coumarins via sequential Claisen-ene reactions of α-aryloxymethylacrylic acids and enophiles, 4393
- (+)-pederin, total synthesis, stereocontrolled synthesis of (+)-benzoylselenopederic acid, 6465
- (+)-thienamycin from 3(R)-hydroxybutyric acid, stereocontrolled synthesis,.1523
- α,β-unsaturated acylsilanes as reactive carboxylic acid equivs, 2509
- α-hydroxy acids by alkylation of chiral N-(benzyloxyacetyl)-trans 2,5-bis(methoxymethoxymethyl)-
- pyrrolidine, 1343 α-hydroxy carboxylic acids, asymmetric synthesis, direct oxidation of chiral amide enolates using 2-sulphonyloxaziridines, 3539
- acids, reaction with 2-chloro-4,6-disubstituted 1,3,5triazines, useful acylating reagents in prep of esters, amides, acid anhydrides and peptides, 2901
- activated carboxylic acids intramplecular lactone annulation onto olefins by Mn(111) yields bicyclo[3.3.0] and [4.3.0]lactones, 3761
- adducts of 2,3-bis(bromomethy1)-1,3-butadiene with propiolic acids, precursors of 2Hisoindoles and 2,3-dithianaphthalenes, 4093
- allenic acids ,cyclisation, synthesis of  $\beta$ -substituted  $\alpha$ , $\beta$ -unsaturated butenolides, 4811 and 1,2-amino alcohols, stereochemistry of Ph,P-CC1, modisted evolution to
  - stereochemistry of Ph,P-CCl, mediated cyclisation to oxazolines with inversion of carbinol center, 4687

- chiral (S)-glutamic acid use as template to construct 7-carbon subunits with alternating OH groups with stereochemical control, prep of enantiomeric and/or diastereomeric 3,5dideoxy-heptitols, 5631
- crystalline stable phosphaethyne derivs via acid chlorides and tris(trimethylsilyl)phosphine, 5507
- different product distributions in reactions of cyclopentyl-and isopropylmagnesiumbromides with cyclohexane-, cyclohex-4-ene and cyclobutane-1,2-dicarboxylic anhydrides, 1297
- dinydriaes, 127, dihydroxy-, stereospecific synthesis via E-2-(phenylsulphonyl)vinyl ethers of 2.3-epoxyalcohols, 6301
- Z,3-epoxyaloohols, 6301
  E and Z C-4 deuterated 3-methyl-zbutenoic acid, stereoselective synthesis and and regioselective deprotonation syn to acid
- function, 1939
  enantiomeric 3,6-hexanooxepin-4carboxylic acid and [6]paracyclophan-8-carboxylic acid,
  absolute configurations, 721
- enzyme prep from <u>E.coli</u> in formation of <u>o</u>-succinylbenzoic acid from iso-chorismic acid, key in vitamin K<sub>2</sub> biosynthesis, 1487
- esters and amides from aziridine 2-carboxylic acid salts, 4439 Y,6-unsaturated carboxylic acid derivs, synthesis by Ni(CO),induced ring opening carbonylation, 5795
- gem-dimethyl cyclopropane carboxylic acids, 2-lithio-2phenylsulphonyl propane as precursor, 6115 large ring lactone, synthesis by
- large ring lactone, synthesis by acid-catalysed cyclisation of (Z)-ene-diyne hydroxy acid
- precursor, 3811 Li dianion of 4-oxo-2-phenylthiobutanoic acid ethylene acetal addition to aldehydes and cyclisation to 2-alkyl-3-furoic acids 2625
- acids, 3635 lithium trienolate of sorbic acid as d\* synthon, prep of 7-hydroxy dienoic acids from ketones, 3625
- methylbenzoic acids and acrylic acids, 2-substituted, oxidation by S<sub>1</sub>O<sub>2</sub>-Ag(I), conversion to phthalides and butenolides via rearrangement of acyloxyl radicals. 1209
- radicals, 1209
  new chiral acid for resolution of racemic bases, (S)-(-)-(2-phenylcarbamoyloxy)propionic acid, 4451
- new juvenile hormone analogs, synthesis of alkenoic acid derivs with cyclopropane ring, 4261
- new stable analogs of prostacyclins,(±)-6a-oxo-6,9-methano-15-hydroxyprosta-5,13-dienoic acids, 4101
- novel synthesis of 3-cyanoindoles, new route to indole-3-carboxylic acid derivs, 1827
- o-substituted benzoic acids enantioselective conversion to chiral cyclohexane derivs, 4575 olefins, conversion to Y-lactones

- platelet activating factor, synthesis from  $(\underline{S})$ -(-)-malic acid, 5195
- regioselective openings of 2,3epoxy acids with organocuprates, 4683
- regioselectivity in halolactonisation of Υ,δ-unsaturated acids, 5497
- resolution of β-hydroxycarboxylic acid derivs on chiral stationary phase derived from (R,R)tetramide, 4217
- ring opening and ring enlargement of 3,3-dimethyl-2-phenylcyclopropene carboxylic acid, 4911
- sepiomelanin, treatment with NaBH., isolation of 5,6dihydroxyindole-2-carboxylic acid, 2801
- series of 2,6-disubstituted unsaturated pimelic acids, synthesis, 3115
- spiro hydrindandione ring system of Fredericamycin A via condensation of 1,4-dimethoxybenzene with 1,1-cyclopentane dicarboxylic acid or 1,1-indan dicarboxylic acid deriv, 4723
- stereoselective annelation of trimethylsiloxyacetic acids and imines into 3-hydroxy-β-lactams, 4239
- stereoselectivity, reaction of sec alkylMg compds with bicyclo-[2.2.1]hept-5-ene-2,3dicarboxylic anhydride and lactones, formation of trans diastereomeric lactones and erythro diols, 4719 symmetrical anhydrides via action
- symmetrical anhydrides via actio of tetracyanoethylene on carboxylic acids, 1503
- threo-or erythro-a-methyl-β-hydroxy acids via chiral propionate enolate equivs, stereoselective synthesis, 2125
- two carboxylic functions, chemodifferentiation by selective lactonisation, 473
- vinylketenes stereo- and regiospecific prep from  $\alpha,\beta$ unsaturated acid chlorides and their intramolecular [2+2] cycloadditions, 5619

### Carbynes

phenols from metal-carbynes and diynes, 2159

# Carcinogens

- carcinogenic aza-arene dihydro diols and diol epoxides, novel synthesis of trans-tetrahydro diol diacetates as precursors, 6417
- sugar as component of embryonic and tumor antigens, 2439 synthesis of riboside analog of pyrimidine adduct from Nhydroxy-N-2-acetylaminofluorene and DNA, 2659

# Carotenoids

ophioxanthin, new carotenoid sulphate from Ophioderma longicaudum, 1871

### Catalvais

alkylation of aromatic compds, silica-alumina supported transition metal oxide catalyst,

- acids, intermediate in ergot biosynthesis, 4043
- CD of β-cyclodextrin with amino moiety, binding ability for ferrocenecarboxylic acid in DMSO, 899
- presence of Mn(111), 4291 oxidative removal of 2,6dimethoxybenzyl and phenyl ester derivs as protecting groups,
- ester and prop-z-enylidene acetal protecting groups, relative stability towards Ir, Rh and Pd catalysts, 5045 BF,-etherate on alumina catalysis in improved synthesis of cannabidiol, 1083

- bovine serum albumin, Darsens condensation of aromatic aldehydes with phenacyl halides,
- HLAD-oxidation of alcohols with acetaldehyde as coenzyme recycling substrate, 4527

hydroxylation of saturated hydrocarbons, oxone as oxygen donor, 4459

Knoevenagel condensation, catalysis, 4453 macropolycyclic molecules with extended basket-shaped skeleton including benzene ring, syn-thesis: THF retained in hydrochloridecavity and accelerated guest-selective H/D exchange in aromatic diols, 1483

Michael reaction, catalysis, 2645 Michael reactions promoted by solid-liquid phase transfer catalysis without solvent, application to addition of acetylacetone, methylaceto-acetate and fluorene anions on hindered acceptors, 4601

Montmorillonite clay, unsymmetrical pyrromethanes via coupling of  $\alpha$ -acetoxymethyl-pyrroles with  $\alpha$ -free pyrroles , 793

nickel boride and NaBH, reduction of aliphatic nitro compds, 6413 rearranged cholest-13(17)-enes catalysed by montmorillonite clay, synthesis, 2135 reduction of aryl ketones with Zn

powder, viologen as electron transfer catalyst, 4525 sec alcohols, new Cr(V1) reagent

for catalytic oxidation to ketones, 5855

stable 2-acinitro-1,3-propanediol, prep from nitromethane and paraformaldehyde with fluoride ion catalysis, 1423

use of Yb(fod), in Diels-Alder reactions of acrolein with dienes, stereoselectivity, 2507

### Cephalosporins

7α-amino-1-oxacephems epimerisatio to 7β-amino epimers, 339 benzyl 6-β-aminopenicillinate conversion into crystalline 6-aformido deriv without chromatography, 377 carbocephalosporin nucleus, enantioselective synthesis, 3787

Cerium and compounds 4-methylcoumarins, 5-substituted and 5,6-disubstituted, synthesis using organo-Ce reagents, 6477 alkylbenzenes, photo-autoxidation catalysed by cerium(1V)ammonium

nitrate, 3353 new halogenated marine prostanoids, antitumour chloro-vulones from <u>Clavularia</u> <u>viridis</u>,

nitrate free radicals, role in photo side-chain nitroxylation

of alkylbenzenes by cerium(1V) ammonium nitrate, 541 tertiary alcohols via CeCl,-promoted addition of Grignard reagents to ketones, 4763

# Cesium and compounds

depsipeptides with D-a-hydroxy carboxylic acids, synthesis via L-α-halo-carboxylic esters and cesium salts of N-protected amino acids, 5257

Chain reactions
chiral 4-hydroxymethyl butenolide use as template for stereocontrolled conjugate addition of C-Me group leading to 7-carbon chain with predictable 1,3-C-Me

substitution, 5627 construction of acyclic chains of 7 carbons (or more) with predisposed, alternating and/or remote C-Me substitution of 1.5type with or without intervening OH groups, 5623

enolate derived from chiral butyrolactone deriv use as template for stereocontrolled introduction of OH groups, 2carbpon chain extension to give diastereomeric or enantiomeric lactones, 5631

### Challeones

photolysis of phenyl cinnamates, synthesis of 2'-hydroxychalcones, 4125

# Charge transfer absorptions

oligooxa[n.n]paracyclophane quinhydrones, cation-induced CT absorptions, 6175

### Chelation

new structural units for prep of chelating and macrocyclic host

molecules, 5749
synthesis of bicyclo[2.1.1]hexenes-5-d by transmetallation,
regio-and stereocontrol by chelation of a neighboring OH function? 1141

Chemiluminescence decomposition of 1,3-dinydro-peroxy-1,1,3,3-tetraphenyl-2-propanone, 173

of intermediate in photooxy-genation of 1,1,3,3-tetramethyl-2-indanone triphenylphosphazine, 843

Chiral compounds 3-,5-,6-and 7-membered carbocyclic compds with extracyclic chiral center by stereoselective synthesis intramolecular alkylation, 1723

(+)(S)-3-trichloromethylbutyric acid via stereoselective conjugate addition of Cl\_CMgCl to crotonate of chiral auxiliary with sulphonylamino substituent, 6047

α-aminonitriles from new N-cyanomethyl-1,3-oxazolidine synthon, 3567

on diastereofacial selectivity in aldol reactions, 6009

α,β-unsaturated iron acyls, useful chiral enoate synthons, 3075

aldehydes, reaction with preformed lithium enolate of methyl 2methylpropanoate, 973 aldehydes, stereoselective aldol

condensation of protected glycine and 3-butyryl thia-zolidine-2-thiones to give βhydroxyesters, intermediates for B-lactam antibiotics, 977

alkoxy-allenes via reaction of chiral acetylenic acetals with Grignard reagents in presence of Cu salt; diastereoselective βelimination, 4197

asymmetric alkylation and sulphenylation of chiral 0silylated imide enclates, 5339 asymmetric synthesis of (R) or

(S)-4,4-dialkyl-2-cyclopentenones, 2047

asymmetric synthesis of a-hydroxy carboxylic acids, direct oxidation of chiral amide

enolates using 2-sulphonyloxaziridines, 3539 bifunctional chiral synthons via

biochemical methods, C, isoprenoid units, 961

bisazetidines via hydroalane reduction of bis-8-lactams, 2035 building block (3S,4R)-3-[(R)--1-hydroxyethyl]-4-acetyloxyazetidin-2-one, stereocontrolled

total synthesis, 937 chirality transfer in [2.3] sigmatropic rearrangement of anions derived from trialkylstannylmethyl allylic ethers, stereospecific synthesis of (2R)-3-benzyloxy-2-methyl-propanol, 5013

cryptocaryone, revised structure,

cyclohexane derivs, by enantio-selective conversion of o-substituted benzoic acids, 4575

DEF-ring system of nogalmycin, chiral synthesis, 2693

diastereoface selectivity, aldol condensation of ethyl 1,3dithiolane and 1,3-dithiane-2-carboxylates with chiral aldehydes, 2977

diastereofaceselection, condensation of aldehydes with Zr enolate of chiral N-propionyl-trans-2,5-disubstituted pyrrolidine, 5807

diastereomers of chiral β,β'dimethyl-bis-homoallylic alcohol with 3 contiguous chiral centers, stereoselective prep via diastereoselective addition of nucleophiles to α-methyl-β, Yunsaturated carbonyl compds, 3707

enantiocomplimentary route to PGA<sub>2</sub> and PGE<sub>2</sub> synthons, inversion of chirality of 2,5-dihydroxy-cyclopent-3-enyl-acetic acid

lactone derivs, 3141 enantioselective syntheses of (+)dictyopterenes A and C' by active allylic benzoate with Pd(0) catalyst, 2779

intermediates for prostanoids, asymmetric synthesis of cis-4-tbutyldimethylsiloxy-2-cyclo-penten-1-ol and cis-4-tetra-hydropyranyloxy-2-cyclopenten-1-01, 5803

key intermediate for mevinolin and compactin, prep of ethyl 5(S),6-epoxy-3(R)-(methoxymethoxy)= hexanoate, 1185 measurement of enantiomeric

excesses of phosphine oxides, use of chiral shift reagent,

methyl esters of N(3,5-dinitro-benzoyl)amino acids, chiral solvating agents for NMR determination of enantiomeric

purity, 2989 new chiral acid for resolution of racemic bases, (S)-(-)-(2-phenylcarbamoyloxy)propionic acid, 4451

new flavin with axial chirality and redox-dependent racemisation, synthesis and resolution, 5183

new route to chiral β-hydroxy-αamino acids, cleavage of oxirane with hydrazoic acid, 5309

novel (4+1)fragment combination approach to chiral cyclopentanoids from tartaric acid, 3121

P-chiral O-isopropyl oligo-nucleotide triesters, synthesis and absolute configuration, 2191

polypropionate units, intermediates for construction, 263

porcine liver esterase, simple, effective, covalent immobilisation and application in prep of chiral building blocks, 407 prochiral C centers, diastereotopic selectivity,

stereodivergent synthesis of talaromycins, 17

resolution of β-hydroxycarboxylic acid derivs on chiral stationary phase derived from (R.R)tetramide, 4217

stereoselective additions to symmetrical ketones via chiral propionate enolate equivs, 2129 stereoselective synthesis of

threo-or erythro-α-methyl-β-hydroxy acids via chiral propionate enolate equivs, 2125

stereospecific displacement of S from chiral centers, activation

via thiaphosphonium salts, 4867 tetracyclo[6.6.0], 5.0°, 12]tetra-decane-3,6,10,13-tetraene, via Weiss reaction, 2275

Troger's base analogs, new units for prep of chiral hosts and metal ligands, 5749

### Chlorophy11

chlorophyll b, synthesis and properties of 3-viny1-3desmethyl chlorophyll a, 4875 model compd for photosynthetic electron transfer, synthesis,

### Chromones

2-substituted chroman-4-ones rearrangement to 3,4-alkenylchromones, 1879

# Chromium and compounds

α,β-unsaturated nitroalkenes reduction by Cr(11)chloride, route to carbonyl compds, 3777

alkynyl-Cr compds, selective addition to aldehyde moiety without affecting CO group of substrate, 5585

cross aldolisation of  $\alpha$ -bromoketones with aldehydes in presence of chromium(11)-chloride, 4371

high asymmetric induction during reduction of pyruvate and phenylglyoxylate esters of chromium complexed carbinol, 429

pyrrole-and phenyl-carbene Crcomplexes, reaction with alkoxalkyne, dimer formation via o-quinonemethide and subsequent dimerisation, 1159 regioselective aromatic

substitution on diaryloxidetricarbonyl-Cr complex, 3989 sec alcohols, new Cr(V1) reagent

for catalytic oxidation to ketones, 5855

stereoselective alkylation of benzylic acetates and free alcohols of (II-arene)tri-carbonyl-Cr complexes to exoalkyl Cr-complexes, 767

### Circular dichroism

absolute configuration of (-)solanapyrone, 2453

absolute stereochemistry of panaxynol by exciton chirality method, 5775 β-cyclodextrin with amino moiety,

binding ability for ferrocene-carboxylic acid in DMSO, 899 CD homoallylic benzoate method, tool for determination of

absolute configuration, 2213 exciton coupling, excimer emission

and unique binding of Y-cyclodextrin substituted by two naphthyl residues, 3339 optically active α-pyridyl

substituted diazocines, Cu(1) selective chiral ligands, CD and cyclic voltammetry, 2077

spectra of chiral α-hydroxy-phosphonates and phosporic acids, 493

stereostructures of halenaquinol and halenaquinol sulphate from Xeatospongia sapra via calculations of CD spectra, 3833

tridecadeoxynucleotide, synthesis, conformation, NMR, IR and CD studies, 2085

### Cleavage reactions

[1,2,4-]triazole[1.5-a]pyrimidine N-ylides, new ring cleavage of triazole moiety to pyrimidine deriv, 1321

(3-chloro-2-methylenecycloalkyl)palladium chloride dimers, synthesis of (±)-13-methyltridecanolide, 4163

activated vinylcyclopropanes, conjugate nucleophilic ring

opening by PD catalysis, 3049 allylic functional groups, reductive removal by nickel boride, 2581

allylic trimethylsilyl ethers with

allylic trimethylsilyl ethers wi nickel boride, 371 anodic oxidation of polycyclic cyclopropanes followed by hydrolysis, regio- and stereo-selective formation of homoallylic alcohols, 4513

axially disymmetric tertiary amines and quaternary ammonium salts, synthesis of new 1,1'-binaphthyl substit amines, 421

benzyl-Si and aryl-Si rupture in reaction of alcohols with benzosilacyclobutenes, formation of (dialkyl)alkoxy-o-tolylsilanes and (dialkyl)alkoxybenzylsilanes, 4761

bicyclic ketals of 6,8-dioxa-bicyclo[3.2.1]octane series to δ, ε-unsaturated ketones by treatment with acetyl iodide, 3895

biphenyl-fused 1,2-diphenylcyclo-butanes, orientation of bond cleavage initiated by electron transfer, 3579

bromine in acetone ring opening of 2,5-disubstituted furan ring,

ether, ester and carbamate protecting groups with catechol boron halides, 1411 hydroquinone dimethyl ethers

oxidative demethylation with nitric acid impregnated MnO2,

Li enclate opening of <u>cis</u> and <u>trans</u>-but-2-ene oxides in <u>presence</u> of <u>BF</u><sub>3</sub>.OEt<sub>2</sub> by S<sub>N</sub>2 mechanism, 4815

N- and S-substituted dioxetanes, activation parameters and rates of decomposition, 3183 new route to chiral β-hydroxy-α-

amino acids, cleavage of oxirane

with hydrazoic acid, 5309 nucleophilic ring of crown ethers for building up crown ether

rings, 2705 nucleophilic, of esters and ethers with phenyltellurotrimethyl-silane, 453

peptides from benzhydrylamine resins, cleavage using tri-fluoromethanesulphonic acid,

peptides, solid phase synthesis of signal sequence fragments using

transesterification cleavage.

photolysis of acyclic monothio-imides to β-lactams and thioamides, involving Y-H abstract ion by thiocarbonyl group, 4475

photosensitised oxygenation of cyclic sulphides, selective C-S bond cleavage, 4609 polypeptides, Edman degradation using new basic cleavage

conditions, 4375

conditions, 4375
protected oligonucleotides,
removal of internucleotidic
phenylthio group by bis(tributyltin) oxide, 4621
m-bonds of carbonyl compds, with
4-t-butyl iodoxybenzene, 4955
regioselective openings of 2,3approxy adds with operancements.

epoxy acids with organocuprates,

regioselective ring opening of epoxides with alcohols catalysed by organotin phosphate

by organotin phosphate condensates, 3219 regiospecific opening of oxetanes with trimethylsilyl cyanide-zinc iodide, approach to Y-amino alcohols, 4971 reversible Claisen rearrangement

of 3-(3,3-dimethylally1)-4-(3,3dimethylallyloxy)quinolin-2-one to buchapsine and loss of its

1,1-dimethylallyl group, 4253 S-N bond, by irradiation of N-tosyldiarylsulphilimines and p-

toluenethiolate anion, 775 sodium hydrogentelluride cleavage of epoxides and quaternary ammonium salts, 6197

thioether protection via selectively cleavable sulphonium salts, 6143

# Cobalt and compounds

alkylation of β-diketones with 1bromoadamantane to a-(1adamanty1)- $\beta$ -diketones via Co(11), Co(111) and Zn(11)

complexes, 3735 bicyclo[3.3.0]octenones, substituted, synthesis via dicobaltoctacarbonyl mediated alkene-alkyne cyclisations, 4851

carboxylation of organic halides electro-catalysed by Co-salen complex, 2633 Co(11)TPP-catalysed isomerisation

of electronegative substituted quadricyclanes to norbornadiene,

desulphurisation of mercaptans to hydrocarbons by CO and water in presence of CO<sub>2</sub>(CO)<sub>a</sub>, 1935

direct entry to perhydrotriquinacene skeleton by Comediated intramolecular cyclisation, 2475

enol silyl ethers via cobalt carbonyl catalysed reaction of tetrahydrofurans with hydrosilane and CO, 2683

reduction of aromatic nitriles with a hydrosilane leading to N,N-disilylamines, cobalt

carbonyl catalysis, 5145 trialkylsilane controlled CO2(CO). catalysed silyl-hydroformylation of oxetanes, 5753

trialkylsilanes, reaction with styrene oxide, Co<sub>2</sub>(CO), catalysed, 5415

### Coenzymes

HLAD-catalysed oxidation of alcohols with acetaldehyde as coenzyme recycling substrate, 4527

intramolecular linkage of flavinporphyrin, rate enhancing effect