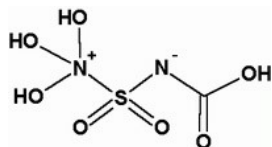


Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu
Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3



Burgess reagent

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt

Purity:	98%
CAS	29684-56-8
FW	238,30
Melting point	78-80 °C

Literature

Aminalester. Zur Stabilität und Umsetzung mit aromatischen Aldehyden

Bredereck, H.; Simchen, G.; Hoffmann, H.; Horn, P.; Wahl, R.; *Angew. Chem.*, 79 (1967) 311;

Darstellung und Eigenschaften der Amidacetale und Aminalester,

Hellmut Bredereck, Gerhard Simchen, Siegfried Rebsdatt, Willi Kantlehner, Peter Horn, Rudolf Wahl, Horst Hoffmann und Peter Grieshaber, *Chem. Ber.*, 101 (1968) 41

New synthetic reactions. Generalalkylation via α -trimethylenedithiocyclobutanones

Trost, B.M.; Preckel, M.; *J. Am. Chem. Soc.*; 97 (1975) 2224;

A novel method for converting ketones to α -diketones. The reaction of enamino ketones with singlet oxygen
Wassermann, H.H.; Ives, J.L.; *J. Am. Chem. Soc.*; 98 (1976) 7868;

Asymmetric synthesis of *allo*-heteroyohimbine alkaloids

Uskokovic, M.R.; Lewis, R.L.; Partridge, J.J.; Despreaux, C.W.; Pruess, D.L.; *J. Am. Chem. Soc.*; 101 (1979) 6742;

Orthoamide, XXXII. Umsetzungen von *tert*-Butoxy-*N,N,N',N'*-tetramethylmethandiamin mit NH- und CH-aciden Verbindungen

Kantlehner, W.; Wagner, F.; Bredereck, H.; *Liebigs Ann. Chem.*, (1980) 344;

Tetracarbonylferrate aus Orthoameisensäurederivaten und Pentacarbonylisen

Daub, J.; Hasenhündl, A.; Krenkler, K.P.; Schmetzer, J.; *Liebigs Ann. Chem.*, (1980) 997;

Darstellung und Reaktionen von Trimethoxyacetamid und Trimethoxyacetamidderivaten

Kantlehner, W.; Kapassakalidis, J.J.; Maier, T.; *Liebigs Ann. Chem.*, (1980) 1448;

Synthesis of 2'-methylated pyrimidine C-nucleosides

Sato, T.; Kobayashi, H.; Noyori, R.; *Tetrahedron Lett.*, 21 (1980) 1971;

Synthesis of 4'-hydroxymethylated pyrimidine ribo-C-nucleosides

Sato, T.; Noyori, R.; *Tetrahedron Lett.*, 21 (1980) 2535;

Total synthesis of racemic ajmalicine and 19-*epi*-ajmalicine

Gutzwiller, J.; Pizzolato, G.; Uskokovic, M.R.; *Helv. Chim. Acta*, 64 (1981) 1663;

Orthoamide, XXXVI. Zur Formylierung von 2,5-Dialkyl-1,3,4-thiadiazolen und 2,3,5-Trialkyl-1,3,4-thiadiazoliumsalzen durch Orthoamide der Ameisensäure

Kantlehner, W.; Haug, E.; Hagen, H.; *Liebigs Ann. Chem.*, (1982) 298;

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu

Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

Einfache Synthese von α -substituierten Tetronsäure-Derivaten

Schmidt, R.R.; Betz, R.; *Synthesis*, (1982) 748;

Stereocontrolled general synthesis of pyrimidine C-nucleosides having branched-chain sugar moieties

Sato, T.; Watanabe, M.; Kobayashi, H.; Noyori, R.; *Bull. Chem. Soc. Jpn.*, 56 (1983) 2680;

Fungal metabolites. Part 15. Structure and chemical correlations of uvidin C, D, and E, new drimane sesquiterpenes from *Lactarius uvidus* Fries

De Bernardi, M.; Mellerio, G.; Vidari, G.; Vita-Finzi, P.; Fronza, G.; *J. Chem. Soc., Perkin Trans. 1*, (1983) 2739;

Stereoselective synthesis of racemic trichodiene

Schlessinger, R.H.; Schulz, J.A.; *J. Org. Chem.*, 48 (1983) 407;

Synthetic strategies toward the synthesis of 2,4-dimethoxypyrrolo[3,2-d]pyrimidine

Cupps, T.L.; Wise, D.S.; Townsend, L.B.; *J. Org. Chem.*, 48 (1983) 1060;

Bromination of some pyridine and diazine N-oxides

Paudler, W.W.; Jovanovic, M.V.; *J. Org. Chem.*, 48 (1983) 1064;

Synthesis of porphyrinotakis(dialkylcarboxamides)

Kaesler, R.W.; LeGoff, E.; *J. Org. Chem.*, 48 (1983) 4399;

Benz[c,d]indoles – I. The use of *tert*-butoxy-bis-(dimethylamino)-methane

Haeflinger, W.; Knecht, H.; *Tetrahedron Lett.*, 24 (1983) 285;

Synthese de cyclopena[c]tetrahydropyridines precuseurs d'alcaloides monoterpeniques

Brayer, J.-L.; Alazard, J.-P.; Thal, C.; *Tetrahedron Lett.*, 24 (1983) 4193;

Reaction of singlet oxygen with enamino carbonyl systems. A general method for the synthesis of α -keto derivatives of lactones, esters, amides, lactams and ketonem

Wassermann, H.H.; Ives, J.L.; *J. Org. Chem.*, 50 (1985) 3573;

Simple enols. 4. Generation of some New simple enols In solution and the kinetics and mechanism of their ketonization

Capon, B.; Siddhanta, A.K.; Zucco, C.; *J. Org. Chem.*, 50 (1985) 3580;

O-Glycosylimidate, 19. Reaktionen von Glycosyl-trichloracetimidaten mit silylierten C-Nucleophilen

Hoffmann, M.G.; Schmidt R.R.; *Liebigs Ann. Chem.*, (1985) 2403;

Total synthesis of acosamine and daunosamine utilizing a diastereoselective intramolecular [3+2] cycloaddition

Wovkulich, P.M.; Uskokovic, M.R.; *Tetrahedron*, 41 (1985) 3455;

An enantiospecific synthesis of estrone

Hutchinson, J.H.; Money, T.; *Tetrahedron Lett.*, 26 (1985) 1819;

New piperidinic synthons *via* ring contraction. Formal synthesis of (\pm)-perhydrohistrionicotoxin

Duhamel, P.; Kotera, M.; Monteil, T.; *Bull. Chem. Soc. Jpn.*, 56 (1986) 2353;

Reaktionen mit 2-aza-1,3-butadien-Derivaten, 1. Eine neue und besonders einfache Synthese von zentralaktiven β -Carbolin-Derivaten

Biere, H.; Russe, R.; Seelen, W.; *Liebigs Ann. Chem.*, (1986) 1749;

The synthesis of Mannich bases from ketonem and esters *via* enamionones

Schuda, P.F.; Ebner, C.B.; Morgan, T.M.; *Tetrahedron Lett.*, 27 (1986) 2567;

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu
Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

Enantiospecific synthesis of estrone

Hutchinson, J.H.; Money, T.; *Can. J. Chem.*, 65 (1987) 1;

Electron-rich and electron-poor pentalene derivatives

Frank, W.; Gompper, R.; *Tetrahedron Lett.*, 28 (1987) 3083;

Thermolysis of polyazapentadienes. Part 11. Concerted and free radical mechanisms in 2-aza enone and 2-aza
enthione pyrolyses: crystal and molecular structures of 3-dimethylamino-1-*p*-tolyl-2-azaprop-2-en-1-one and 3-
dimethylamino-1-phenyl-2-azaprop-2-ene-1-thione

Blake, A.J.; McNab, H.; Murray, M.E.-A.; *J. Chem. Soc., Perkin Trans. 1*, (1989) 589

New 7-substituted quinolone antibacterial agents. II. The synthesis of 1-ethyl-1,4-dihydro-4-oxo-7-(pyrazoyl,
isoxazolyl, and pyrimidinyl)-1,8-naphthyridine and quinolone-3-carboxylic acids

Domagala, J.M.; Peterson, P.; *J. Heterocyclic Chem.*, 26 (1989) 1147;

Highly stereoselective ring contraction of heterocyclic enamines: total synthesis of perhydrohistrionicotoxin and
its 2,6-epimer

Duhamel, P.; Kotera, M.; Monteil, T.; Marabout, B.; *J. Org. Chem.*, 54 (1989) 4419;

Total synthesis of (-)-jolkinalide A, B, and E utilizing a new mild esterification followed by intramolecular
Wittig-Horner reaction

Katsumura, S.; Kimura, A.; Isoe, S.; *Tetrahedron*, 45 (1989) 1337;

Tricyclic Pyridine Derivatives with High Affinity to the Central Benzodiazepine Receptor

Fischer, U.; Möhler, H.; Schneider, F.; Widmer, U.; *Helv. Chim. Acta*, 73 (1990) 763;

3,4-Disubstituted γ -lactam rings as conformationally constrained mimics of peptide derivatives containing
aspartic acid or norleucine

Garvey, D.S.; May, P.D.; Nadzan, A.M.; *J. Org. Chem.*, 55 (1990) 936;

Synthesis of advanced quinocarcin intermediate from L-glutamic acid

Lessen, T.A.; Demko, D.M.; Weireb, S.M.; *Tetrahedron Lett.*, 31 (1990) 2105;

Synthesis of 7-ethyl-4,7-dihydro-4-oxo-2-(4-pyridinyl)hieno[2,3-*b*]pyridine-5-carboxylic acid

Bacon, E.R.; Daum, S.J.; *J. Heterocyclic Chem.*, 28 (1991) 1953;

Expanded heterohelicenes: molecular coils that form chiral complexes

Bell, T.W.; Jousselein, H.; *J. Am. Chem. Soc.*; 113 (1991) 6283;

β -Carbolin-Alkaloide, II. Tributyl(1-ethoxyvinyl)stannan als C2-Baustein für die Synthese von β -Carbolin-
Alkaloiden

Bracher, F.; Hildebrand, D.; *Liebigs Ann. Chem.*, (1993) 837;

Enantioselective synthesis of 5-substituted- and 3,5-disubstituted-2-formylpyrrolidine derivatives, the key D-ring
fragments of (-)-quinocarcin and (-)-decarboxyquinocarcin

Katoh, T.; Nagata, Y.; Kobayashi, Y.; Arai, K.; Minami, J.; Tereshima, S.; *Tetrahedron Lett.*, 34 (1993) 5743;

Synthesis of 3-(*N*-alkylamino)acetophenones via a benzyne intermediate

Albright, J.D.; Liebermann, D.F.; *J. Heterocyclic Chem.*, 31 (1994) 537;

Total synthesis of drimane sesquiterpenes from *S*(+)-carvone (part 5)

Swarts, H.J.; Versteegen-Haaksma, A.A.; Jansen B.J.M.; de Groot, A.; *Tetrahedron*, 50 (1994) 10083;

Synthesis and stereochemistry of some thiazolidines related to 6-(hydroxyethyl)-penams

Urban, F.J.; Bordner, J.; DeCosta, D.; Dee, M.F.; Vincent, L.A.; *Tetrahedron: Asymmetry*, 5 (1994) 215;

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu

Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

The serotonin 5-HT₄ receptor. 2. Structure-activity studiem of the indole carbazimidamide class of agonists
Buchheit, K.-H.; Gamse, R.; Giger, R.; Hoyer, D.; Klein, F.; Kloppner, E.; Pfannkuche, H.-J.; Maties, H.; *J. Med. Chem.*, 38 (1995) 2331;

Cyclic variations of 3-quinolinecarboxamides and effects on antiherpetic activity
Wentland, M.P.; Carlson, J.A.; Dorff, P.H.; Aldous, S.C.; Perni, R.B.; Young, D.C.; Woods, M.G.; Kingsley, S.D.; Ryan, K.A.; Rosi, D.; Drozd, M.L.; Dutko, F.J.; *J. Med. Chem.*, 38 (1995) 2541;

Novel and potent adenosine 3',5'-cyclic phosphate phosphodiesterase III inhibitors: thiazolo [4,5-b] [1,6]naphthyridin-2-ones
Singh, B.; Bacon, E.R.; Leshner, G.Y.; Robinson, S.; Pennock, P.O.; Bode, D.C.; Pagani, E.D.; Bentley, R.G.; Connell, M.J.; Hamel, L.T.; Silve, P.J.; *J. Med. Chem.*, 38 (1995) 2546;

New synthetic routes to 3-, 5-, and 6-aryl-2-chloropyridines
Chuch, R.; Trust, R.; Albright, J.D.; Powell, D.W.; *J. Org. Chem.*, 60 (1995) 3750;

Synthesis of functionalized aryloxy 1,3-butadienes and their transformation to diaryl ethers *via* Diels - Alder cycloaddition reactions
Olsen, R.K.; Feng, X.; Campbell, M.; Shao, R.; Math, S.K.; *J. Org. Chem.*, 60 (1995) 6025;

Mannich biscyclization. Total synthesis of (-)-ajamalicine
Logers, M.; Ogerman, L.E.; Welmaker, G.S.; *J. Am. Chem. Soc.*; 117 (1995) 9139;

A novel approach to the synthesis of 1-substituted-3,6-diaryl-imidazo[1,5-*b*]pyridazines
Zhao, X.; Zhang, R.; *Synth. Comm.*, 25 (1995) 3271;

Synthesis of kainoid analogues
Barraclough, P.; Hudhomme, P.; Spray, C.A.; Young, D.W.; *Tetrahedron*, 51 (1995) 4195;

Stereospecific synthesis of (2*S*,4*R*)-[5,5,5-²H₃]leucine
August, R.A.; Khan, J.A.; Moody, C.M.; Young, D.W.; *J. Chem. Soc., Perkin Trans. 1*, (1996) 507;

Study on the preparation of heteroaryl substituted enamines. A simple synthesis of heteroaryl substituted acetaldoximes from enamines
Copar, A.; Stanovnik, B.; Eisler, M.; *J. Heterocyclic Chem.*, 33 (1996) 465;

Concise syntheses of novel 1*H*-pyrrolo[3,2-*g*]quinazoline ring system and its [2,3-*f*] angular isomer
Showalter, H.D.H.; Sun, L.; Sercel, A.; Winters, R.T.; Denny, W.A.; Palmer, B.D.; *J. Org. Chem.*, 61 (1996) 1155;

A novel synthesis of the antipsychotic agent ziprasidone
Urban, F.J.; Breitenbach, R.; Gonyaw, D.; *Synth. Comm.*, 26 (1996) 1629;

Chemistry and nonlinear optical properties of new 2*H*-benzotriazole derivatives
Gompper, R.; Walther, P.; Brauchle, C.; Stadle, S.; *Tetrahedron*, 52 (1996) 14607;

Ring contraction of a two-carbon bridged spiropentane
Wiberg, K.B.; Snoonian, J.R.; Lahti, P.M.; *Tetrahedron Lett.*, 37 (1996) 8285;

Synthesis of (η⁵-cyclopentadienyl)-1-(4-benzyloxycarbonyl-3,4-dihydroquinoxalin-2-yl)et hene-1,2-dithiolatocobalt(III) and (η⁵-cyclopentadienyl)-1-[2-(*N,N*-dimethylaminomethyleneamino)-3-methyl-4-oxopteridin-6-yl]ethene-1,2-di thiolatocobalt(III)
Dinsmore, A.; Birks, J.H.; Garner, C.D.; Joule, J.A.; *J. Chem. Soc., Perkin Trans. 1*, (1997) 801;

Synthesis of agonists and antagonists for central glutamate receptors by a novel 'ring switching' strategy

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu
Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

Bowler, A.N.; Dinsmore, A.; Doyle, P.M.; Young, D.W.; *J. Chem. Soc., Perkin Trans. 1*, (1997) 1297;

Nitrosation of methyl 2-cinnamoylamino-3-dimethylaminopropenoates. Alkil N-cinnamoyloxalic acid hydroxyimidic amides, intermediates in the synthesis of alkil 5-styryl-1,2,4-oxodiazole-3-carboxylates
Kmetec, M.; Stankovic, B.; *J. Heterocyclic Chem.*, 34 (1997) 1705;

Synthesis of oligo(diazaphenyls). Tailor-made fluorescent heteroaromatics and pathways to nanostructures
Gompper, R.; Mair, H.-J.; Polborn, K.; *Synthesis*, (1997) 696;

A convenient and efficient synthesis of (2*S*,4*R*)- and (2*S*,4*S*)-4-methylglutamic acid
Coudert, E.; Acher, F.; Azerad, R.; *Synthesis*, (1997) 863;

4-Substituted protoanemonin in intramolecular cycloaddition reactions of non-stabilised azomethine ylides
Grigg, R.; Samic, V.; Thornton-Pett, M.; *Tetrahedron*, 53 (1997) 10633;

A convergent route for the total synthesis of the eleuthesides
Chen, X.-T.; Gutteridge, C.E.; Bhattacharya, S.K.; Zhou, B.; Pettus, T.R.R.; Hascall, T.; Danishefsky, S.J.;
Angew. Chem. Int. Ed., 37 (1998) 185;

Stereoselective 1,3-Dipolar Cycloadditions to (S)-1-Benzoyl-3-(cyanomethylidene)-5-(methoxycarbonyl)pyrrolidin-2-one
Skof, M.; Svete, J.; Stanovnik, B.; Golic, L.; Golic-Grdadolnik, S.; Selic, L.; *Helv. Chim. Acta*, 81 (1998) 2332;

Synthesis and structures of 1,3,1',3'-tetrabenzyl-2,2'-biimidazolidinylidenes (electron-rich alkenes), their amination intermediates and their degradation products
Çetinkaya, B.; Çetinkaya, E.; Chamizo, J.A.; Hitchcock, P.B.; Jasim, H.A.; Küçükbay, H.; Lappert, M.F.; *J. Chem. Soc., Perkin Trans. 1*, (1998) 2047;

Ring expansion and contraction of a two-carbon bridged spiropentane
Wilberg, K.W.; Snoonian, J.R.; *J. Org. Chem.*, 63 (1998) 1390;

Molecular architecture. 2. Synthesis and metal complexation of heptacyclic terpyridyl molecular clefts
Bell, T.W.; Cragg, P.J.; Firestone, A.; Kwok, A.D.-I.; Liu, J.; Ludwig, R.; Sodoma, A.; *J. Org. Chem.*, 63 (1998) 2232;

4-(2,2-Dimethyldioxalan-4-yl)-5-(pterin-6-yl)-1,3-dithiol-2-ones proligands relating to the cofactor of the oxomolybdoenzymes
Dinsmore, A.; Garner, C.D.; Joule, J.A.; *Tetrahedron*, 54 (1998) 9559;

The synthesis and transformations of substituted 2-hydroxy-3-dimethylaminopropenoates. The preparation of condensed 3-hydroxypyran-2-ones
Smodis, J.; Stanovnik, B.; *Tetrahedron*, 54 (1998) 9799;

Solid phase synthesis of 5-aminopyrazoles and derivatives part II
Wilson, R.D.; Watson, S.P.; Richards, S.A.; *Tetrahedron Lett.*, 39 (1998) 2827;

Spiro-imidazo[1,2-a]indeno[1,2-c]pyrazine-4-one derivatives are mixed AMPA and NMDA glycine-site antagonists active in vivo
Jimonet, P.; Boireau, A.; Chev e, M.; Damour, D.; Genevois-Borella, A.; Imperato, A.; Pratt, J.; Randle, J.C.R.; Ribeill, Y.; Stutzmann, J.-M.; Mignani, S.; *Bioorg. Med. Chem. Lett.*, 9 (1999) 2921;

Stereoselective amination of 5-substituted γ -lactones and γ -lactams - a convenient route for the preparation of 5-substituted (3*S*,5*S*)-3-acetylamino-tetrahydrofuran-2-ones and (3*S*,5*S*)-3-acetylamino-pyrrolidin-2-ones
Skof, M.; Svete, J.; Kmetec, M.; Golic-Grdadolnik, S.; Stanovnik, B.; *Eur. J. Org. Chem.*; (1999) 1581;

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu
Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

Synthesis of Highly Soluble Annelated Polypyridines

Bejan, E.; Fontenas, C.; Aït-Haddou, H.; Daran, J.-C.; Balavoine, G.G.A.; *Eur. J. Org. Chem.*; (1999) 2485;

The synthesis of methyl 2-(benzyloxycarbonyl)amino-3-dimethylaminopropenoate. The synthesis of trisubstituted pyrroles, 3-amino-2*H*-pyran-2-ones, Fusem 2*H*-pyran-2-ones and 4*H*-pyridin-4-ones
Toplak, R.; Svete, J.; Stankovnik, B.; Grdadolnik, S.G.; *J. Heterocyclic Chem.*, 36 (1999) 225;

Attempts to prepare some 3-substituted azolo[1,2-*x*]azines, intermediates In the synthesis of azaaplysinopsin derivatives

Jukic, L.; Copar, A.; Malesic, M.; Krabavcic, A.; Svete, J.; Stankovnik, B.; *J. Heterocyclic Chem.*, 36 (1999) 1147;

Regioselective Preparation of N⁷- and N⁹-Alkyl Derivatives of N⁶-[(Dimethylamino)methylene]adenine Bearing an Active Methylene Group and Their Further Derivatization Leading to α -Branched Acyclic Nucleoside Analogues

Hockova, D.; Budesinsky, M.; Marek, R.; Marek, J.; Holy, A.; *Eur. J. Org. Chem.*; (1999) 2675;

The total synthesis of eleutherobin

Chen, X.-T.; Bhattacharya, S.K.; Zhou, B.; Gutteridge, C.E.; Pettus, T.R.R.; Danishefsky, S.J.; *J. Am. Chem. Soc.*; 121 (1999) 6563;

Transformations of alkyl 2-(2,2-disubstituted-ethenyl)amino-3-dimethylaminoprop-2-enoates: synthesis of alkyl 3,4-disubstituted- and alkyl 1-acyl-3,4-disubstituted pyrrole-2-carboxylates

Selic, L.; Stanovnik, B.; *Synthesis*, (1999) 479;

An effective route to polysubstituted symmetric terpyridines

Sasaki, I.; Daran, J.C.; Balavoine, G.G.A.; *Synthesis*, (1999) 815;

Selective synthesis of β -substituted aspartic acids *via* tetrahydro-1,3-oxazin-6-ones

Burtin, G.; Corringier, P.-J.; Hitchcock, P.B.; Young, D.W.; *Tetrahedron Lett.*, 40 (1999) 4275;

The synthesis and vasopressin (AVP) antagonist activity of a novel series of n-aro-yl-2,4,5,6-tetrahydropyrazolo[3,4-*d*]thieno[3,2-*b*]azepines

Albright, J.D.; Santos, E.G.D.; Dusza, J.P.; Chan, P.S.; Coupet, J.; Ru, X.; Mazandarani, H.; *Bioorg. Med. Chem. Lett.*, 10 (2000) 695;

Synthesis of New Soluble Annelated Polypyridines

Aït-Haddou, H.; Fontenas, C.; Bejan, E.; Daran, J.-C.; Balavoine, G.G.A.; *Eur. J. Org. Chem.*; (2000) 987

Synthesis of (2*S*)-2-(Benzoylamino)-3-(heteroaryl)propyl Benzoates

Skof, M.; Svete, J.; Stanovnik, B.; Golic-Grdadolnik, S.; *Helv. Chim. Acta*, 83 (2000) 760;

A Simple Stereoselective Synthesis of Aplysinopsin Analogs

Selic, L.; Jakse, R.; Lampic, K.; Golic, L.; Golic-Grdadolnik, S.; Stanovnik, B.; *Helv. Chim. Acta*, 83 (2000) 2802;

Tetrahydro-1,3-oxazin-6-ones as templates for the stereoselective synthesis of β -substituted L-aspartic acids

Burtin, G.; Corringier, P.-J.; Young, D.W.; *J. Chem. Soc., Perkin Trans. 1*, (2000) 3451;

Synthesis of 1,4,5,6-tetrahydropyrazolo[3,4-*d*]pyrido[3,2-*b*]azepine

Albright, J.D.; Du, X.; *J. Heterocyclic Chem.*, 37 (2000) 41;

Synthesis of transition state analogue inhibitors for purine nucleoside Phosphorylase and *N*-ribidose hydrolases

Evans, G.B.; Furneaux, R.H.; Gainsford, G.J.; Schramm, V.L.; Tyler, P.C.; *Tetrahedron*, 56 (2000) 3053;

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu

Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

New synthesis of 2-aminobicyclo[2.1.1]hexane-2,5-dicarboxylic acid-I (ABHxD-I), a potent metabotropic receptor agonist

Conti, P.; Kozikowski, A.P.; *Tetrahedron Lett.*, 41 (2000) 4053;

Synthetic studiem towards (+)-Dihydroampullicin. Michael addition of *N*-Boc-2-(*tert*-butyldimethylsiloxy)-3-methyl-pyrrole to α -methylene lactones

Marcos, I.; Rudero, E.; Bermejo, F.; *Tetrahedron Lett.*, 41 (2000) 8451;

NO-Independent stimulators of soluble guanylate cyclase

Straub, A.; Stasch, J.-P.; Alonso-Alija, C.; Benet-Buchholz, J.; Ducke, B.; Feurer, A.; Furstner, C.; *Bioorg. Med. Chem. Lett.*, 11 (2001) 781;

Synthesis of anticonvulsive AMPA antagonists - 4-Oxo-10-substituted-imidazo[1,2-a]indeno[1,2-e]pyrazin-2-carboxylic acid derivatives

Stutzmann, J.-M.; Bohme, G.A.; Boireau, A.; Damour, D.; Debono, M.W.; Genevois-Borella, A.; Jimonet, P.; Pratt, J.; Randle, J.C.R.; Ribeill, Y.; Vuilhorgne, M.; Mignani, S.; *Bioorg. Med. Chem. Lett.*, 11 (2001) 1205;

1,3-Dipolar cycloadditions to (5*Z*)-1-acyl-5-(cyanomethylidene)-imidazolidine-2,4-diones: synthesis and transformations of spirohydantoin derivatives

Groselj, U.; Drobnic, A.; Recnik, S.; Svete, J.; Stanovnik, B.; Golobic, A.; Lah, N.; Leban, I.; Meden, A.; Golic-Grdadolnik, S.; *Helv. Chim. Acta*, 84 (2001) 3403;

Synthesis of the organic ligand of the molybdenum cofactor, in protected form

Bradshaw, B.; Dinsmore, A.; Ajana, W.; Collison, D.; Garner, C.D.; Joule, J.A.; *J. Chem. Soc., Perkin Trans. 1*, (2001) 3239;

The synthesis and transformations of 2-[2-ethoxycarbonyl-2-(2-pyridinyl)ethenyl]amino-3-

dimethylaminopropenoates. The synthesis of substituted β -amino- α,β -didehydro- α -amino acid derivatives
Jukic, L.; Recnik, S.; Grdadolnik, S.G.; Svete, J.; Stanovnik, B.; *J. Heterocyclic Chem.*, 38 (2001) 859;

Synthesis of (S)-3-heteroaryl-2-hydroxy-1-propyl benzoates by "Ring switching" methodology

Mihelic, D.; Jakse, R.; Svete, J.; Stanovnik, B.; Grdadolnik, S.G.; *J. Heterocyclic Chem.*, 38 (2001) 1307;

A New synthetic router to 3,4-bridged 1,6,6a λ^4 -trithiapentalenes

Hang, W.; Henry, Y.; *Synlett*, (2001) 1129;

Enaminones acylation: competitive formation of quinolin-4-one and isoquinolin-1-one derivatives

Vales, M.; Lokshin, V.; Pepe, G.; Samat, A.; Guglielmetti, R.; *Synthesis*, (2001) 2419;

Asymmetric synthesis of α -amino acids from α,β -(*Z*)-didehydroamino acid derivatives with 1,2,3,6-tetrahydropyrazin-2-one structure

Abellan, T.; Mancheno, B.; Najera, C.; Sansano, J.M.; *Tetrahedron*, 57 (2001) 6627;

A simple synthesis of aplysinopsin analogues by dimethylamine substitution In *N,N*-(dimethylamino)methylidene derivatives of five-membered heterocycles

Jakse, R.; Recnik, S.; Svete, J.; Golobic, A.; Golic, L.; Stanownik, B.; *Tetrahedron*, 57 (2001) 8395;

Extension of the "ring switch" approach to glutamate antagonists to δ -lactam urethanes

Coe, D.; Drysdale, M.; Philips, O.; West, R.; Young, D.W.; *J. Chem. Soc., Perkin Trans. 1*, (2002) 2459;

Synthesis of spiro lactones by 1,3-dipolar cycloadditions to methyl (S)-3-[(*E*)-cyanomethylidene]-2-oxotetrahydrofuran-5-carboxylate

Pirc, S.; Recnik, S.; Škof, M.; Svete, J.; Golic, L.; Meden, A.; Stanovnik, B.; *J. Heterocyclic Chem.*, 39 (2002) 411

Methoxycarbonylsulfamoyl triethylammonium hydroxide, inner salt, Burgess reagent, CAS 29684-56-8
Item available at : www.bujno.eu
Specific web site : http://www.bujno.eu/index.php?sel_odd=CEN&page=8&szuk_tekst=green&id_of=3

Synthesis of 1,2-dithiolane analogues of leucine for potential use in peptide chemistry
Morera, E.; Pinnen, F.; Lucente, G.; *Org. Lett.*, 4 (2002) 1139;

An Expedient Synthesis of Highly Functionalized Naphthyridones and Quinolines from a Common *N*-Aryl Pyridinone Template
Savarin, C.G.; Murry, J.A.; Dormer, P.G.; *Org. Lett.*, 4 (2002) 2071;

Practical synthesis of 8-acyl-7-alkyl-1,6-naphthyridin-5(6*H*)-ones
Vales, M.; Lokshin, V.; Pepe, G.; Guglielmetti, R.; Samat, A.; *Tetrahedron*, 58 (2002) 8543;

Stereoselective synthesis of (1*R*,3*R*,4*R*)-3-(1,2,4-triazolo[4,3-*x*]azin-3-yl)-1,7,7-trimethylbicyclo[2.2.1]heptan-2-ones
Groselj, U.; Recnik, S.; Svete, J.; Meden, A.; Stanownik, B.; *Tetrahedron: Asymmetry*, 13 (2002) 821;

Novel B-ring modified allocolchicinoids of the NCME series: design, synthesis, antimicrotubule activity and cytotoxicity.
Bergemann, S.; Brecht, R.; Buttner, F.; Guenard, D.; Gust, R.; Seitz, G.; Stubbs, M.T.; Thoret, S.; *Bioorg. Med. Chem.*, 11 (2003) 1269;

Total synthesis of ingenol
Tanino, K.; Onuki, K.; Asano, K.; Miyashita, M.; Nakamura, T.; Takahashi, Y.; Kuwajima, I.; *J. Am. Chem. Soc.*; 125 (2003) 1498;

Traceless solid phase synthesis of 2-substituted pyrimidines using an 'off-the-shelf' chlorogermane-functionalised resin.
Spivey, A.C.; Srikaran, R.; Diaper, C.M.; Turner, D.J.; *Org. Biomom. Chem.*, 1 (2003) 1638;

An alternative to the use of δ -lactam urethanes in the "ring switch" approach to higher homologues of AMPA-type glutamate antagonists.
Hitchcock, P.B.; Masood, S.R.; Young, D.W.; *Org. Biomom. Chem.*, 1 (2003) 2682;

Enamine-functionalized oligopyridines as convenient intermediates for the synthesis of carbaldehyde derivatives
Viau, L.; Senechal, K.; Maury, O.; Guegan, J.-P.; Dupau, P.; Toupet, L.; Le Bozec, H.; *Synthesis*, (2003) 577;

Electrochemical probing of ground state electronic interactions in polynuclear complexes of a new heteroditopic ligand
Konstable, E.C.; Figgemeier, E.; Housecroft, C.E.; Olsson, J.; Zimmermann, Y.C.; *Dalton Trans.*, (2004) 1918