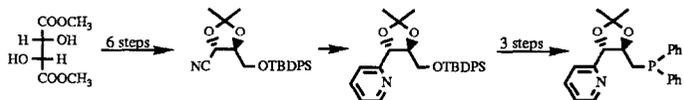


Tetrahedron: Asymmetry 1994, 5, 299

(-)-(4*S*,5*R*)-4-(2-Pyridyl)-5-(diphenylphosphino)methyl-2,2-dimethyl-1,3-dioxolane a new chiral ligand for enantioselective catalysis

Giorgio Chelucci,^{a*} M. Antonietta Cabras,^a Carlo Botteghi^b and M. Marchetti^c

^aDipartimento di Chimica, Università di Sassari, Via Vienna 2, I-07100 Sassari, Italy; ^bDipartimento di Chimica, Università di Venezia, S. Marta 2137, Venezia, Italy; ^cIstituto A.T.C.A.P.A., C.N.R., via Vienna 2, I-07100 Sassari, Italy



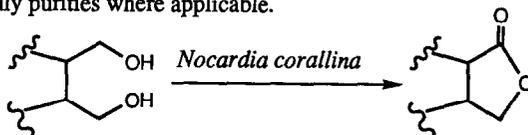
The title compound has been prepared by a ten reaction sequence from L-(+)-tartrate and checked in asymmetric hydroesterification and cross-coupling reactions.

Oxidation of Diols to Lactones by *Nocardia corallina* B-276

H. Luna, K. Prasad,* O. Repic;

Chemical Research & Development, Technical R & D, Sandoz Research Institute, Route 10, East Hanover, NJ 07936, USA

1,4-Diols were oxidized by *Nocardia corallina* B-276, to the corresponding lactones with good to excellent optical purities where applicable.



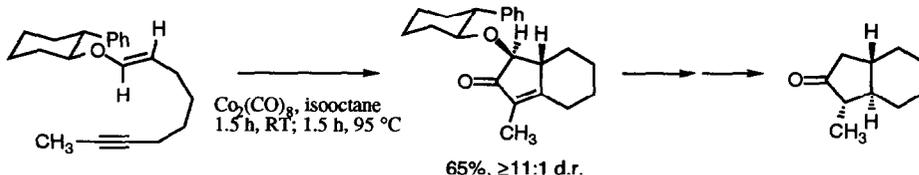
Tetrahedron: Asymmetry 1994, 5, 303

ASYMMETRIC SYNTHESIS OF BICYCLO[4.3.0]NONAN-8-ONES BY INTRAMOLECULAR PAUSON-KHAND REACTION

Jaume Castro, Albert Moyano*, Miquel A. Pericàs*, Antoni Riera and Andrew E. Greene^a

Dep. de Química Orgànica, Universitat de Barcelona, C/ Martí i Franquès, 1-11. 08028-Barcelona, Spain

^aLEDSS, Université J. Fourier, BP 53X, 38041 Grenoble, France

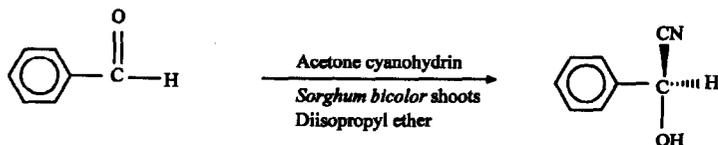


Tetrahedron: Asymmetry 1994, 5, 307

***Sorghum bicolor* Shoots in the Synthesis of (S)-Mandelonitrile**

Eero Kiljunen and Liisa T. Kanerva

Departments of Chemistry and Biomedicine, University of Turku, 20500 Turku, Finland

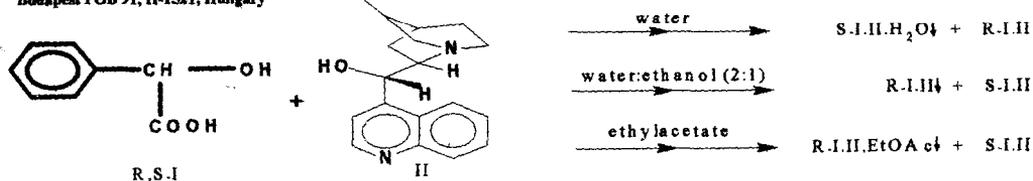


Tetrahedron: Asymmetry 1994, 5, 311

OPTICAL RESOLUTION OF MANDELIC ACID BY CINCHONINE IN DIFFERENT SOLVENTS

Dávid Kozma^a, Ágnes Nyéki^a, Mária Ács^b, Elemér Fogassy^a

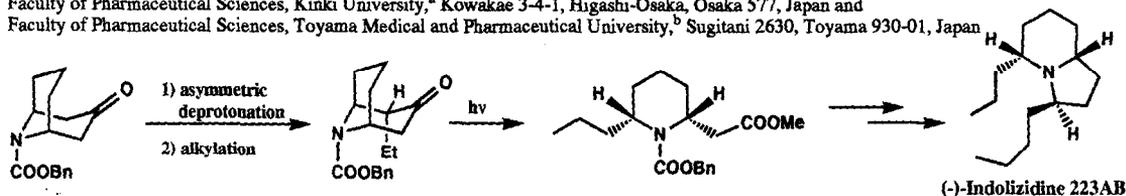
a) Department of Organic Chemical Technology and b) Institute of General and Analytical Chemistry, Technical University of Budapest, Budapest POB 91, H-1521, Hungary



The Norrish Type I Photo-cleavage of (+)-2 β -Ethyl-9-azabicyclo[3.3.1]nonan-3-one: A Short, Enantioselective Formal Synthesis of (-)-Indolizidine 223 AB

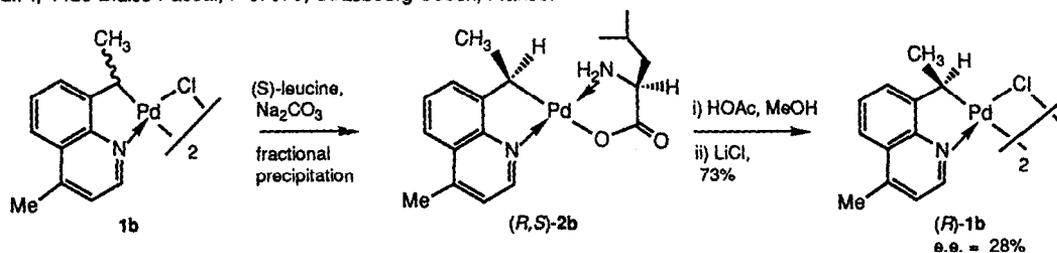
Osamu Muraoka,^a Kazuhito Okumura, Tomomi Maeda, Genzoh Tanabe,^a and Takefumi Momose^{ab}

Faculty of Pharmaceutical Sciences, Kinki University,^a Kowakae 3-4-1, Higashi-Osaka, Osaka 577, Japan and Faculty of Pharmaceutical Sciences, Toyama Medical and Pharmaceutical University,^b Sugitani 2630, Toyama 930-01, Japan



Resolution of a Cyclopalladated Complex Containing an Asymmetric Metallated Carbon Atom

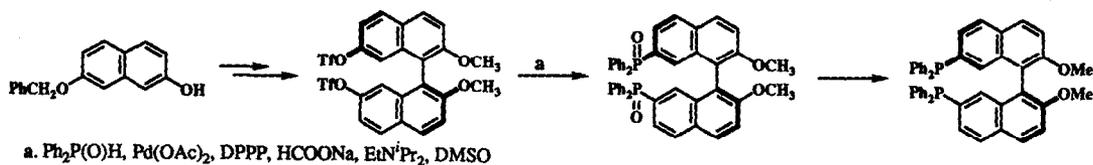
John Spencer, Fida Maassarani, Michel Pfeffer^{*}, André De Cian, Jean Fischer U.L.P., 4 rue Blaise Pascal, F-67070, Strasbourg Cedex, France.



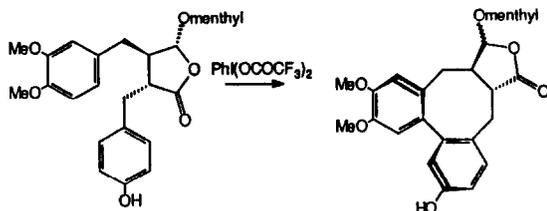
Synthesis of (R)- and (S)-7,7'-Bis(diphenylphosphino)-2,2'-dimethoxy-1,1'-binaphthyl, a New Axially Dissymmetric Bis(triarylphosphine)

Toshihide Horiuchi, Tetsuo Ohta, Massoud Stephan and Hidemasa Takaya^{*}

Division of Material Chemistry, Faculty of Engineering, Kyoto University, Sakyo-ku, Kyoto 606-01, Japan



Asymmetric Syntheses of Isotegane Derivatives using Trivalent Iodine Reagents.
Andrew Pelter, Robert S. Ward and Atef Abd-el-Ghali
Chemistry Department, University of Wales, Singleton Park, Swansea SA2 8PP, UK.

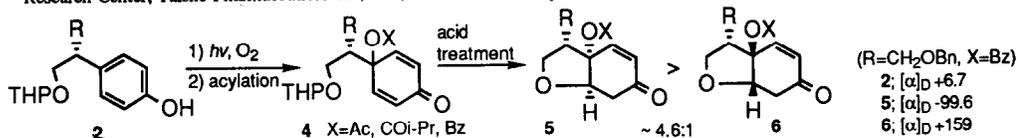


Diastereotopic Group Selective Intramolecular Conjugate Addition of
4-(2-Hydroxyethyl)-*p*-Quinol Derivatives: Synthesis of the Optically
Pure *cis*-7-Oxabicyclo[4,3,0]non-2-en-4-one Skeleton

Hiroichi Fujioka^a, Shinji Kitagaki^a, Naoko Ohno^a, Hidetoshi Kitagawa^a, Yasuyuki Kita^a and Keita Matsumoto^b

^a Faculty of Pharmaceutical Sciences, Osaka University, 1-6 Yamada-oka, Suita, Osaka 565, Japan

^b Research Center, Taisho Pharmaceutical Co., Ltd., Yoshino-cho, Omiya, Saitama 330, Japan

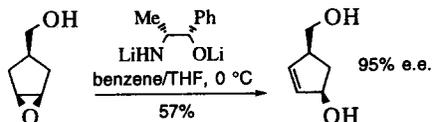


Highly Enantioselective Rearrangement of a *meso*-Epoxide to an Allyl Alcohol
for Carbo-cyclic Nucleoside Synthesis: an Internal Alkoxide Effect

David M. Hodgson,^{*a} Jason Witherington^a and Brian A. Moloney^b

^a Department of Chemistry, University of Reading,
Whiteknights, PO Box 224, Reading RG6 2AD, U.K.

^b Schering Agrochemicals Limited, Chesterford Park,
Saffron Walden, Essex CB10 1XL, U.K.



A New Asymmetric Bridging Annulation Reaction Involving the
Intramolecular Michael Addition of Chiral Imines to Enoates

Françoise Dumas, Véronique Maine, Christian Cavé, Jean d'Angelo

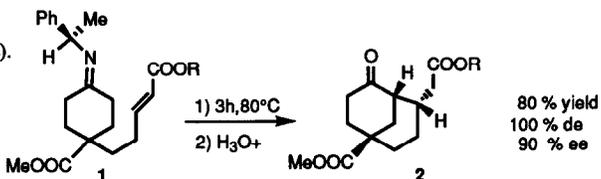
Laboratoire de Chimie Organique, associé au CNRS, Faculté de Pharmacie, 5 rue J.-B. Clément, 92296 Châtenay-Malabry (France)

Angèle Chiaroni, Claude Riche

Institut de Chimie des Substances Naturelles,

CNRS, Avenue de la Terrasse, 91198 Gif-sur-Yvette (France).

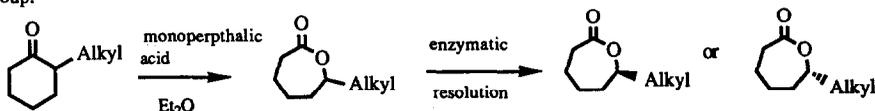
Thermal cyclization of imine **1** led to bicyclic
derivative **2** with high stereocontrol.



R.FELLOUS*, L.LIZZANI-CUVELIER, M.A.LOISEAU and E.SASSY

Laboratoire de Chimie des Aromes, Université de Nice Sophia-Antipolis, Parc Valrose - 06108 - Nice Cedex 2

ϵ -lactones obtained by Baeyer-Villiger oxidation from 2-n-alkylcyclohexanones is regioselective. Enzymatic resolution of 6-n-alkyl ϵ -lactones with PLE gives preferentially lactone of configuration *R*; with HLE enantiomer *S* of lactone is collected from the butyl group.

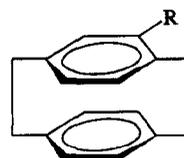


Resolution of Chiral [2.2]Paracyclophanes by Enantioselective Gas Chromatography

Tetrahedron: Asymmetry 1994, 5, 347

Wilfried A. König*¹, Bärbel Gehrcke¹, Detlev H. Hochmuth¹,
Cornelia Mlynek² and Henning Hopf²

- 1) Institut für Organische Chemie, Universität Hamburg, Martin-Luther-King-Platz 6, D-20146 Hamburg, Germany
- 2) Institut für Organische Chemie, Technische Universität Braunschweig, Hagenring 30, D-38106 Braunschweig, Germany

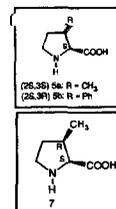
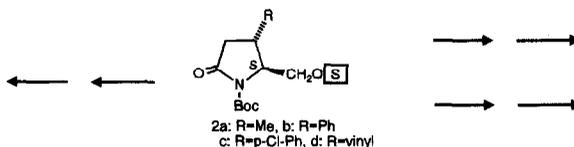
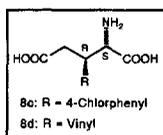


Tetrahedron: Asymmetry 1994, 5, 351

Synthesis of Homochiral 3-Substituted Glutamic Acids and Prolines from *S*-Pyroglutamic Acid

Claus Herdeis*, Hans Peter Hubmann and Hermann Lotter

Institut für Pharmazie und Lebensmittelchemie der Universität D-97074 Würzburg, Am Hubland



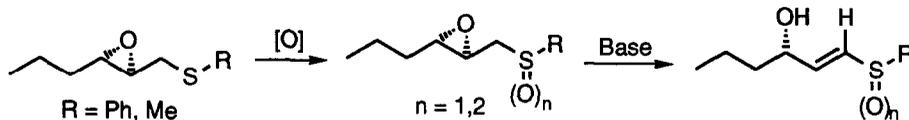
HOMOCHIRAL 2,3-EPOXY SULPHIDES AS PRECURSORS TO E - γ -HYDROXY- α,β -UNSATURATED SULPHOXIDES AND SULPHONES.

Tetrahedron: Asymmetry 1994, 5, 355

(Christopher M. Rayner* and Andrew D. Westwell)

School of Chemistry, University of Leeds, Leeds LS2 9JT, U.K.

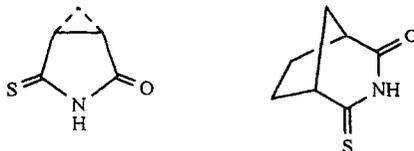
Abstract: The synthesis of *E*- γ -hydroxy- α,β -unsaturated sulphoxides and sulphones from the corresponding homochiral 2,3-epoxy sulphides is described with good to excellent stereochemical control.



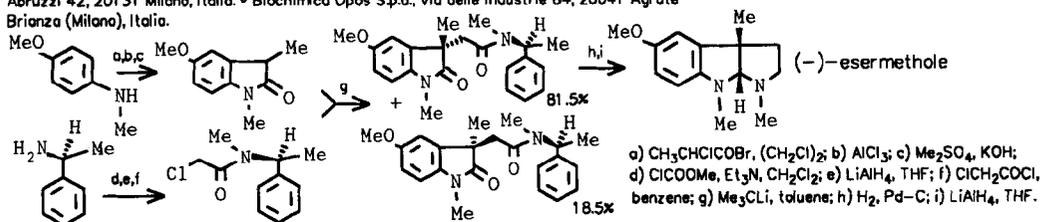
SYNTHESIS OF OPTICALLY ACTIVE MONOTHIOIMIDES WITH CHIRALITY DUE TO SULPHUR SUBSTITUTION.

Maria J. Milewska and Tadeusz Poloński*, Department of Chemistry, Technical University of Gdańsk, 80-952 Gdańsk, Poland

Bicyclic monothioimides of the known absolute configuration were obtained by treatment of corresponding monoamides with Lawesson's reagent



New asymmetric synthesis of (-)-esermethole. M.Palavicini^a, E.Valati^a, L.Villa^a and I.Resto^b. ^aIstituto di Chimica Farmaceutica e Tossicologica dell'Università di Milano, viale Abruzzi 42, 201 31 Milano, Italia. ^bBiochimica Opos S.p.a., via delle Industrie 84, 20041 Agrate Brianza (Milano), Italia.



LEWIS ACID CATALYZED DIELS-ALDER CYCLOADDITIONS OF CHIRAL BUTENOLIDES TO CYCLOPENTADIENE: *endo:exo* STEREOSELECTIVITY

Zhiyu Chen and Rosa M. Ortuño*

Departament de Química, Universitat Autònoma de Barcelona, 08193 Bellaterra, Barcelona, Spain.



catalyst: $\text{ZnCl}_2 / \text{EtAlCl}_2$ for Z = H

ZnI_2 for Z = OH

endo isomers: 80 - 90% yield

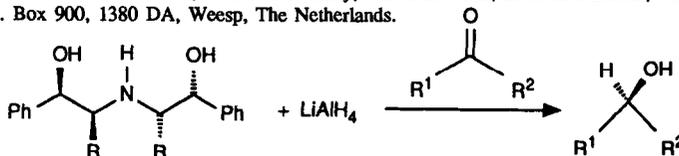
Stereoselective Reduction of Prochiral Ketones, Using Aluminum Hydride Reagents Prepared from LiAlH_4 and Chiral Diethanolamines.

Erik F.J. de Vries¹, Johannes Brussee¹, Chris G. Kruse² and Arne van der Gen¹.

1) Department of Chemistry, Gorlaeus Laboratories, Leiden University, P.O. Box 9502, 2300 RA Leiden, The Netherlands.

2) Solvay Duphar B.V., P.O. Box 900, 1380 DA, Weesp, The Netherlands.

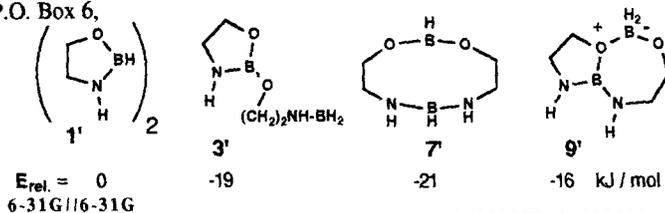
The asymmetric reduction of prochiral ketones by LiAlH_4 , modified with chiral diethanolamines, was studied.



Quantum Chemical Modeling of Chiral Catalysis. Part 16. On the Isomerism of Dimers of Chiral Oxazaborolidines Used in the Catalytic Enantioselective Reduction of Ketones

Vesa Nevalainen, Division of Organic Chemistry, P.O. Box 6, SF-00014 University of Helsinki, Finland.

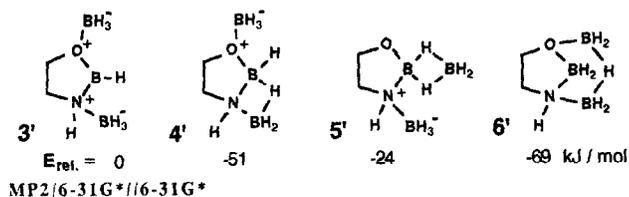
Abstract: Relative stabilities of dimeric isomers of oxazaborolidines were studied by means of *ab initio* MO (RHF) methods (1' as a model of oxazaborolidines). Formation of several dimeric isomers (e.g. 3', 7' and 9') was found to be energetically advantageous.



Quantum Chemical Modeling of Chiral Catalysis. Part 17. On the Diborane Derivatives of Chiral Oxazaborolidines Used as Catalysts in the Enantioselective Reduction of Ketones

Vesa Nevalainen, Division of Organic Chemistry, P.O. Box 6, SF-00014 University of Helsinki, Finland.

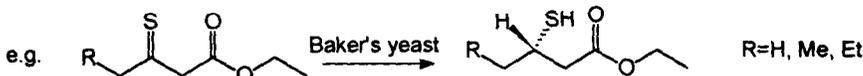
Abstract: Relative stabilities of isomers of borane (H_3B) diadducts (e.g. 3' - 6') of oxazaborolidines were investigated by means of *ab initio* MO (RHF) methods (1,3,2-oxazaborolidine as a model of oxazaborolidines). A new plausible regeneration pathway for the CBS reduction was proposed.



Stereoselective Reduction of Thiocarbonyl Compounds with Baker's Yeast

Jens Kærsgaard Nielsen and Jørgen Øgaard Madsen, Department of Organic Chemistry, The Technical University of Denmark, DK-2800 Lyngby, Denmark

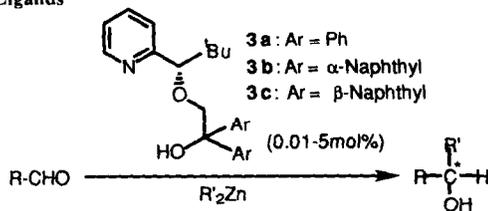
Optically active thiols are produced by baker's yeast reduction of open-chain and cyclic β -thioxo esters



Catalysed Asymmetric Reaction of Aldehydes with Dialkylzinc in the Presence of Chiral Pyridyl Alcohols as Ligands

Miyuki Ishizaki, Ko-ichi Fujita, Mahoko Shimamoto, and Osamu Hoshino*
 Faculty of Pharmaceutical Sciences, Science University of Tokyo, Shinjuku-ku, Tokyo 162, Japan

Catalysed asymmetric addition of dialkylzinc to various aldehydes using 3a-c proceeded in a short reaction time to give optically active alcohols (up to 95% e.e.).

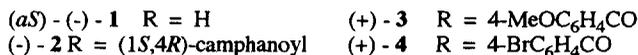
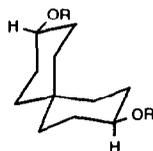


Resolution and Absolute Configuration of the Enantiomeric

Spiro[5.5]undecane-3,9-diols

Gundula Voß and H. Gerlach

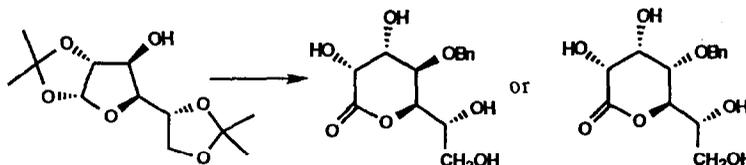
Laboratory of Organic Chemistry, University of Bayreuth, D-95440 Bayreuth, Germany

The absolute configuration of **3** and **4** was derived from their chiroptical properties.ACETONIDES OF HEPTONOLACTONES: KILIANI ASCENSION OF 3-*O*-BENZYL-D-GLUCOSE AND 3-*O*-BENZYL-D-ALLOSE

Claire J. F. Bichard, Joseph R. Wheatley and George W. J. Fleet

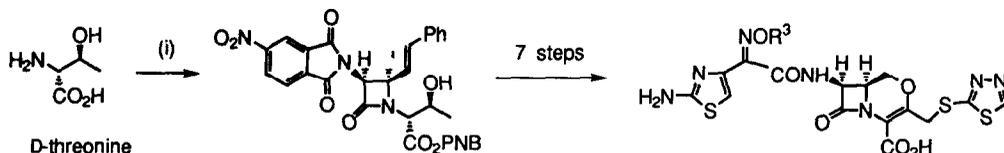
Dyson Perrins Laboratory, Oxford Centre for Molecular Sciences, South Parks Road, Oxford OX1 3QY, UK

Two δ -heptonolactones, epimeric at C-4, and their acetonides have been prepared from diacetone glucose.

A Convenient One Pot Asymmetric Synthesis of *cis*- β -Lactams:

Key Precursors for Optically Active 2-oxaisocephems

Hidetsugu Tsubouchi,* Koichi Tsuji, Koichi Yasumura, Nobuhito Tada, Shinji Nishitani, Jun-ichi Minamikawa, and Hiroshi Ishikawa



(i) one pot asymmetric synthesis : 63% yield , PNB = p-nitrobenzyl

SYNTHESIS OF (3*R*,6*R*)- AND (3*S*,6*S*)-3,6-DIALKYLPIPERAZIN-2,5-DIONE DERIVATIVES AS USEFUL INTERMEDIATES TO BOTH (*R*)-AND (*S*)- α -AMINOACIDS

Gianni PORZI and Sergio SANDRI-Dipartimento di Chimica "G.Ciamician" Università di Bologna Italy

