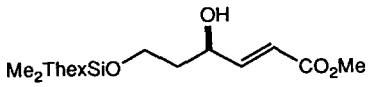
Tetrahedron: Asymmetry **1990**, *1*, 57

K. Burgess and I. Henderson



E.e = 92 % [by ^{1}H NMR with Eu(hfc)$_3$]

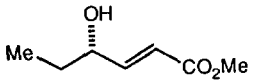
$[\alpha]_D^{25}$ = +5.2 ^{o} (c = 13, $CHCl_3$)

R configuration inferred from correlations of shift experiments with similar compounds and the sequence used to obtain this material.

E-methyl 6-(dimethylthexylsiloxy)-4-hydroxyhex-2-enoate

Tetrahedron: Asymmetry **1990**, *1*, 57

K. Burgess and I. Henderson



E.e = >98 % [by ^{1}H NMR with Eu(hfc)$_3$]

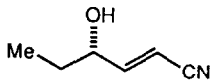
$[\alpha]_D^{25}$ = +23.5 ^{o} (c = 3, $CHCl_3$)

S configuration inferred from correlations of shift experiments with similar compounds and the sequence used to obtain this material.

E-methyl 4-hydroxyhex-2-enoate

Tetrahedron: Asymmetry **1990**, *1*, 57

K. Burgess and I. Henderson



E.e = >98 % [by NMR of MPTA ester]

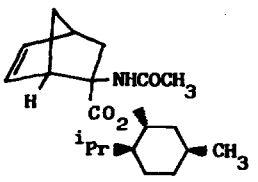
$[\alpha]_D^{25}$ = +47 ^{o} (c = 1.5, $CHCl_3$)

S configuration inferred from correlations of shift experiments with similar compounds and the sequence used to obtain this material.

E-4-hydroxyhex-2-enenitrile

Tetrahedron: Asymmetry **1990**, *1*, 61

C. Cativiela, P. López, J. A. Mayoral.



^{1}**H-NMR:** δ= 2.38(m,1H,H_1), 1.98(s,3H,$COCH_3$), 0.75(d,3H,CH_3)

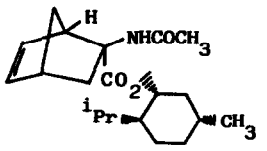
Absolute configuration: 1S,2R,4S
(assigned by comparing with the corresponding hydrogenated amino acid)

$C_{20}H_{31}NO_3$

(-)-menthyl (1S,2R,4S)-2-acetamidobicyclo[2.2.1]hept-5-ene-2-carboxylate

C. Cativiela, P. López, J. A. Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



^{1}H-NMR:δ= 2.32(m,1H,H_1), 1.96(s,3H,$COCH_3$), 0.70(d,3H,CH_3)

Absolute configuration: 1R,2S,4R
(assigned by comparing with the corresponding hydrogenated amino acid)

$C_{20}H_{31}NO_3$

(-)-menthyl (1R,2S,4R)-2-acetamidobicyclo[2.2.1]hept-5-ene-2-carboxylate

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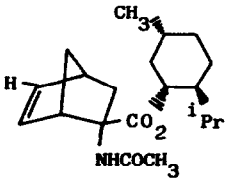
C. Cativiela, P. López, J. A. Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



^{1}H-NMR: δ= 6.04(m,1H,H_5), 2.03(s,3H,$COCH_3$)

Absolute configuration: 1S,2S,4S
(assigned by comparing with the corresponding hydrogenated amino acid)

$C_{20}H_{31}NO_3$

(-)-menthyl (1S,2S,4S)-2-acetamidobicyclo[2.2.1]hept-5-ene-2-carboxylate

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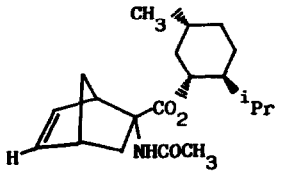
C. Cativiela, P. López, J.A. Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



^{1}H-NMR: δ= 6.10(m,1H,H_5), 1.89(s,3H,$COCH_3$)

Absolute configuration: 1R,2R,4R
(assigned by comparing with the corresponding hydrogenated amino acid)

$C_{20}H_{31}NO_3$

(-)-menthyl (1R,2R,4R)-2-acetamidobicyclo[2.2.1]hept-5-ene-2-carboxylate

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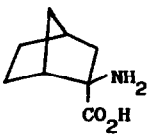
C. Cativiela, P. López, J. A. Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



$[\alpha]_D^{25}$ (1% in water)= -61.4°

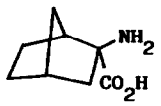
Absolute configuration: 1R,2R,4S

$C_8H_{13}NO_2$

(1R,2R,4S)-2-aminobicyclo[2.2.1]heptane-2-carboxylic acid

C. Cativiela, P. López, J. A. Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



$[\alpha]_D^{25}$ (1% in water)= +61.2°

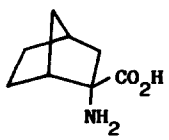
Absolute configuration: 1S,2S,4R

$C_8H_{13}NO_2$

(1S,2S,4R)-2-aminobicyclo[2.2.1]heptane-2-carboxylic acid

C. Cativiela, P. López, J.A.Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



$[\alpha]_D^{25}$ (1% in water)= -24.7°

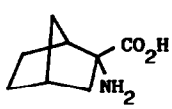
Absolute configuration: 1R,2S,4S

$C_8H_{13}NO_2$

(1R,2S,4S)-2-aminobicyclo[2.2.1]heptane-2-carboxylic acid

C. Cativiela, P. López, J.A. Mayoral.

Tetrahedron: Asymmetry **1990**, *1*, 61



$[\alpha]_D^{25}$ (1% in water)= +24.4°

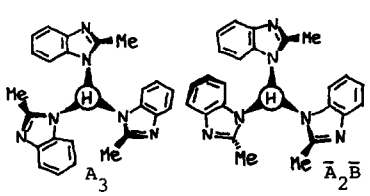
Absolute configuration: 1S,2R,4R

$C_8H_{13}NO_2$

(1S,2R,4R)-2-aminobicyclo[2.2.1]heptane-2-carboxylic acid

Foces-Foces, C.; Hernández Cano, F.; Faure, R.; Roussel, C.; Claramunt, R.M.; López, C.; Sanz, D.; Elguero, J. and Martínez-Ripoll, M.

Tetrahedron: Asymmetry **1990**, *1*, 65



Separation by chromatography on MCT affords enantiomers of TMBM as mixtures of rapid interconverting isomers ($A_3.\bar{A}_2\bar{B}$ and $\bar{A}_3.A_2B$).

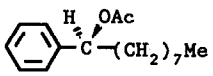
Racemization barrier (28.5 kcal.mol^{-1}) as well as isomerization barrier (9.8 kcal.mol^{-1}) were measured.

The X-Ray structures of the racemic and the (+)enantiomer were determined.

Tris-(2'-methylbenzimidazol-1'-yl)methane (TMBM)

K. Mori and R. Bernotas.

Tetrahedron: Asymmetry **1990**, *1*, 87



E.e. = 100% (by HPLC on a Chiralcel® OB column)

$[\alpha]_D^{25}$ = -63.57 (*c* 1.565, 99.5% EtOH)

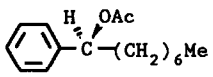
$C_{17}H_{26}O_2$

Source of chirality: resolution by lipase P
Absolute configuration: *S*
[assigned by chemical correlation with (*S*)-1,2-decanediol]

1-Phenylnonyl acetate

K. Mori and R. Bernotas.

Tetrahedron: Asymmetry **1990**, *1*, 87



E.e. = 99.6% (by HPLC on a Chiralcel® OB column)

$[\alpha]_D^{25}$ = -66.51 (*c* 3.95, 99.5% EtOH)

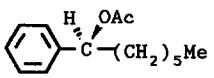
$C_{16}H_{24}O_2$

Source of chirality: resolution by lipase P
Absolute configuration: *S*
(assigned by the comparison of the sign of the specific rotation and the stereospecificity of the enzyme.)

1-Phenyloctyl acetate

K. Mori and R. Bernotas.

Tetrahedron: Asymmetry **1990**, *1*, 87



E.e. = 100% (by HPLC on a Chiralcel® OB column)

$[\alpha]_D^{25}$ = -72.29 (*c* 1.32, 99.5% EtOH)

$C_{15}H_{22}O_2$

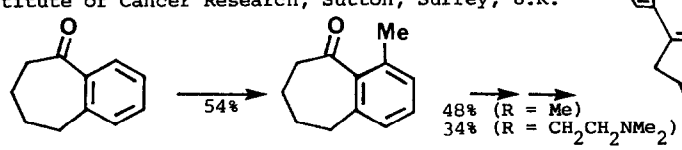
Source of chirality: resolution by lipase P
Absolute configuration: *S*
(assigned by the comparison of the sign of the specific rotation and the stereospecificity of the enzyme.)

1-Phenylheptyl acetate

Tetrahedron: Asymmetry **1990**, *1*, 97

An Analogue of the Antioestrogen Tamoxifen of Sufficient Rigidity to Exist as Distinct Enantiomers: Synthesis and Conformational Dynamics Studies

Raymond McCague
Institute of Cancer Research, Sutton, Surrey, U.K.



When R = Me enantiomeric atropisomers are separable on (+)-poly(triphenylmethyl)-methacrylate.

Racemisation has $\Delta G^{\ddagger}$ = 20.9 kcal mol^{-1}

Tetrahedron: Asymmetry **1990**, *1*, 97

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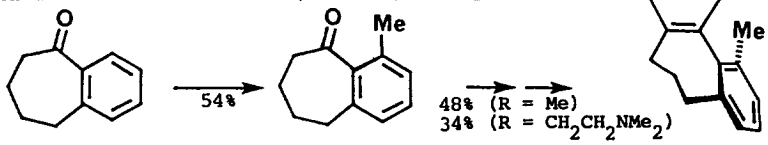
Raymond McCague
Institute of Cancer Research, Sutton, Surrey, U.K.



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Racemisation has $\Delta G^{\ddagger}$ = 20.9 kcal mol^{-1}

Tetrahedron: Asymmetry **1990**, *1*, 97

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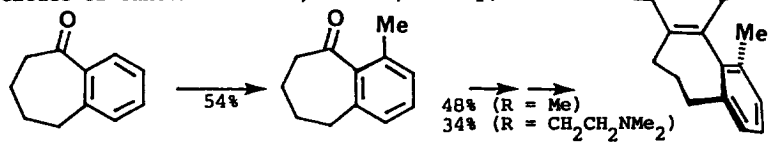
Raymond McCague
Institute of Cancer Research, Sutton, Surrey, U.K.

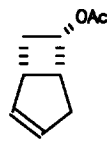


When R = Me enantiomeric atropisomers are separable on (+)-poly(triphenylmethyl)-methacrylate.

Racemisation has $\Delta G^{\ddagger}$ = 20.9 kcal mol^{-1}

N. Klempier, P. Geymayer, P.Stadler, K.Faber. H.Griengl

Tetrahedron: Asymmetry **1990**, *1*, 111

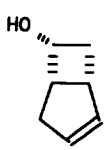


E.e.=99.3% [by (-)-menthylchloroformate]
$[\alpha]_D^{20}$=-36.3 (c 2.27, $CHCl_3$)
Source of chirality: enzymatic resolution
Absolute configuration 1*S*,5*R*,6*S* by $[\alpha]_D^{20}$ of lit.

$C_9H_{12}O_2$
Bicyclo[3.2.0]hept-2-en-6-yl acetate

N. Klempier, P. Geymayer, P.Stadler, K.Faber. H.Griengl

Tetrahedron: Asymmetry **1990**, *1*, 111

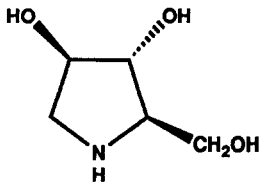


E.e.=98.4% [by (-)-menthylchloroformate]
$[\alpha]_D^{20}$=-75.8 (c 2.46, $CHCl_3$)
Source of chirality: enzymatic resolution
Absolute configuration 1*R*,5*S*,6*R* by $[\alpha]_D^{20}$ of lit.

$C_7H_{10}O$
Bicyclo[3.2.0]hept-2-en-6-ol

G. W. J. Fleet and D. R. Witty

Tetrahedron: Asymmetry **1990,** *1*, 119



E.e. = 100%

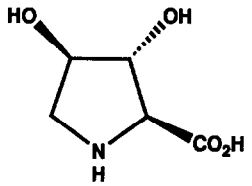
$[\alpha]_D^{20}$ = +36.7° (c, 1.25 in water)

Source of chirality: D-glucose as starting material

$C_5H_{11}NO_3$. HCl
1,4-Dideoxy-1,4-imino-D-arabinitol hydrochloride

G. W. J. Fleet and D. R. Witty

Tetrahedron: Asymmetry **1990,** *1*, 119



E.e. = 100%

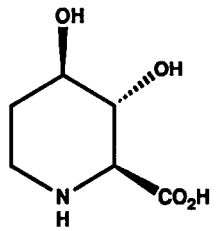
$[\alpha]_D^{20}$ = -12.2° (c, 0.83 in water)

Source of chirality: D-glucose as starting material

$C_5H_9NO_4$
(2S,3R,4R)-3,4-Dihydroxyproline

G. W. J. Fleet and D. R. Witty

Tetrahedron: Asymmetry **1990,** *1*, 119



E.e. = 100%

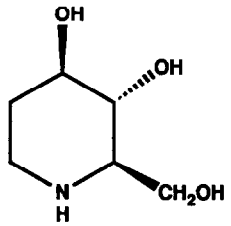
$[\alpha]_D^{20}$ = -1.3° (c, 0.54 in water)

Source of chirality: D-glucose as starting material

$C_6H_{11}NO_4$. H_2O (2S,3R,4R)-3,4-Dihydroxypipecolic Acid

G. W. J. Fleet and D. R. Witty

Tetrahedron: Asymmetry **1990,** *1*, 119



E.e. = 100%

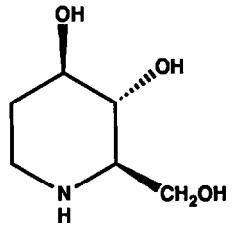
$[\alpha]_D^{20}$ = +21.0° (c, 0.3 in water)

Source of chirality: D-glucose as starting material

$C_6H_{13}NO_3$ Fagomine 1,5,-Imino-1,2,5-trideoxy-D-arabino-hexitol

Tetrahedron: Asymmetry **1990,** *1*, 119

G. W. J. Fleet and D. R. Witty



E.e. = 100%

$[\alpha]_D^{20} = +17.9°$ (c, 0.78 in water)

Source of chirality: D-glucose as starting material

$C_6H_{13}NO_3$. HCl Fagomine Hydrochloride, 1,5,-Imino-1,2,5-trideoxy-D-arabino-hexitol hydrochloride