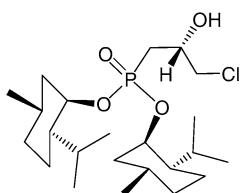


Stereochemistry abstracts

Vitaly V. Nesterov and Oleg I. Kolodiazhnyi*

Tetrahedron: Asymmetry 17 (2006) 1023



$C_{23}H_{44}ClO_4P$
(*2S*)-[Bis(*1R,2S,5R*)-menthyl]-3-chloro-2-hydroxypropylphosphonate

De ~100% (NMR)

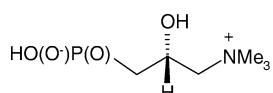
$[\alpha]_D = -97.2$ (*c* 3, CHCl_3)

Source of chirality: double asymmetric synthesis with
(*-*)(*1R,2S,5R*)-menthol and (*R,R*)-tartaric acid

Absolute configuration: *S*

Vitaly V. Nesterov and Oleg I. Kolodiazhnyi*

Tetrahedron: Asymmetry 17 (2006) 1023



$C_6H_{16}NO_4P$
(*R*)-3-(Trimethylammonium)-2-hydroxypropylphosphonic acid

Ee ~99% (NMR)

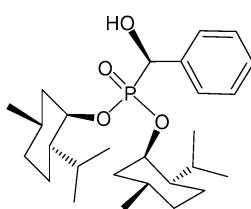
$[\alpha]_D = +25$ (*c* 1, H_2O)

Source of chirality: double asymmetric synthesis with
(*-*)(*1R,2S,5R*)-menthol and (*R,R*)-tartaric acid

Absolute configuration: *R*

Vitaly V. Nesterov and Oleg I. Kolodiazhnyi*

Tetrahedron: Asymmetry 17 (2006) 1023



$C_{27}H_{45}O_4P$
(*1S*)-Bis[*(1R,2R,5S)*-menthyl] hydroxy(phenyl)methylphosphonate

De 99% (NMR)

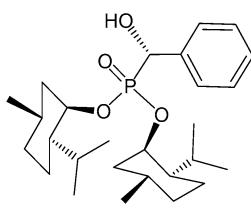
$[\alpha]_D = -70.0$ (*c* 1, CHCl_3)

Source of chirality: double asymmetric synthesis with
(*-*)(*1R,2S,5R*)-menthol and (*S,S*)-tartaric acid

Absolute configuration: *S*

Vitaly V. Nesterov and Oleg I. Kolodiazhnyi*

Tetrahedron: Asymmetry 17 (2006) 1023



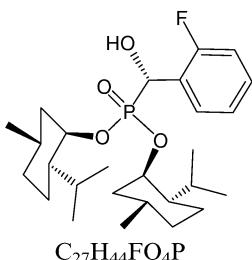
$C_{27}H_{45}O_4P$
(*1R*)-Bis[*(1R,2R,5S)*-menthyl] hydroxy(phenyl)methylphosphonate

De 99% (NMR)

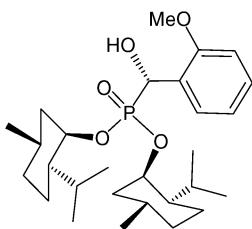
$[\alpha]_D = -87.6$ (*c* 2, CHCl_3)

Source of chirality: double asymmetric synthesis with
(*-*)(*1R,2S,5R*)-menthol and (*R,R*)-tartaric acid

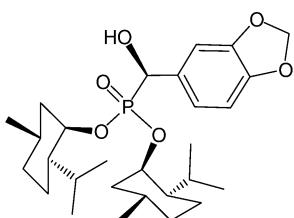
Absolute configuration: *R*

(1S)-Bis[(1*R*,2*R*,5*S*)-menthyl] (2-fluorophenyl)(hydroxyl)methylphosphonate

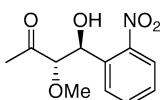
Ee >98% (NMR)

 $[\alpha]_D = -83.7$ (*c* 2, CHCl₃)Source of chirality: double asymmetric synthesis with
(-)-(1*R*,2*S*,5*R*)-menthol and (*R,R*)-tartaric acidAbsolute configuration: *S*(1S)-Bis[(1*R*,2*R*,5*S*)-menthyl] hydroxy(2-methoxyphenyl)methylphosphonate

Ee >98% (NMR)

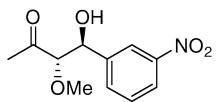
 $[\alpha]_D = -75.2$ (*c* 1, CHCl₃)Source of chirality: double asymmetric synthesis with
(-)-(1*R*,2*S*,5*R*)-menthol and (*R,R*)-tartaric acidAbsolute configuration: *S*(1*R*)-Bis[(1*R*,2*R*,5*S*)-menthyl] (benzo[*d*][1,3]dioxol-5-yl)(hydroxyl)methylphosphonate

Ee >98% (NMR)

 $[\alpha]_D = -74$ (*c* 1, CHCl₃)Source of chirality: double asymmetric synthesis with
(-)-(1*R*,2*S*,5*R*)-menthol and (*R,R*)-tartaric acidAbsolute configuration: *R* $C_{11}H_{13}NO_5$
(3*S*,4*S*)-4-Hydroxy-3-methoxy-4-(2-nitrophenyl)butan-2-one $[\alpha]_D^{20} = +78$ (*c* 0.83, CHCl₃)

Ee 98%

Source of chirality: asymmetric synthesis

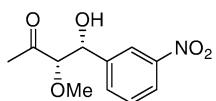


$C_{11}H_{13}NO_5$
(*3S,4S*)-4-Hydroxy-3-methoxy-4-(3-nitrophenyl)butan-2-one

$[\alpha]_D^{20} = -22$ (*c* 0.4, CHCl₃)

Ee 91%

Source of chirality: asymmetric synthesis

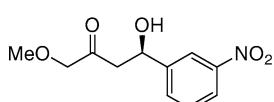


$C_{11}H_{13}NO_5$
(*3S,4R*)-4-Hydroxy-3-methoxy-4-(3-nitrophenyl)butan-2-one

$[\alpha]_D^{20} = -18$ (*c* 0.52, CHCl₃)

Ee 27%

Source of chirality: asymmetric synthesis



$C_{11}H_{13}NO_5$
(*R*)-4-Hydroxy-1-methoxy-4-(3-nitrophenyl)butan-2-one

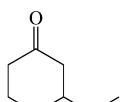
$[\alpha]_D^{20} = +18$ (*c* 0.23, CHCl₃)

Ee 39%

Source of chirality: asymmetric synthesis

Ee = 76%

$[\alpha]_D = +119.7$ (*c* 1, CHCl₃)

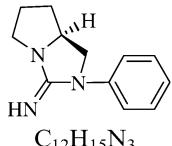


$C_8H_{14}O$
3-Ethlcyclohexanone

$[\alpha]_D^{25} = -48.7$ (*c* 1.1, CHCl₃)

Source of chirality: L-glutamic acid

Absolute configuration: *S*



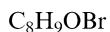
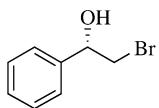
(*S*)-1,3-Diaza-2-imino-3-phenylbicyclo(3.3.0)octane

Ee = 91%

$[\alpha]_D^{25} = +39.8$ (*c* 1.2, CHCl₃)

Source of chirality: asymmetric reduction

Absolute configuration: *S*



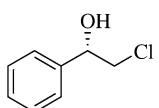
(*S*)-2-Bromo-1-phenylethanol

Ee = 87%

$[\alpha]_D^{25} = +43.8$ (*c* 1.0, cyclohexane)

Source of chirality: asymmetric reduction

Absolute configuration: *S*



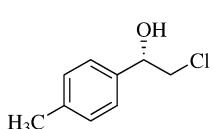
(*S*)-2-Chloro-1-phenylethanol

Ee = 84%

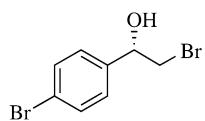
$[\alpha]_D^{25} = +43.1$ (*c* 0.8, CHCl₃)

Source of chirality: asymmetric reduction

Absolute configuration: *S*



(*S*)-2-Chloro-1-(4-methylphenyl)ethanol



C₈H₈OBr₂
(S)-2-Bromo-1-(4-bromophenyl)ethanol

Ee = 90%

[α]_D²⁵ = +30.5 (*c* 1.0, CHCl₃)

Source of chirality: asymmetric reduction

Absolute configuration: *S*