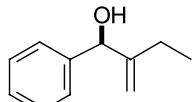


Stereochemistry abstracts

Gelson J. Andrade Conceição, Paulo J. S. Moran and  
J. Augusto R. Rodrigues\*

Tetrahedron: Asymmetry 14 (2003) 43



C<sub>11</sub>H<sub>14</sub>O

(S)-2-Ethyl-1-phenylprop-2-en-1-ol

E.e. >99%

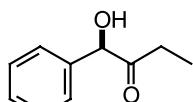
[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -40 (c 2, CHCl<sub>3</sub>)

Source of chirality: biocatalytic reduction

Absolute configuration: *S*

Gelson J. Andrade Conceição, Paulo J. S. Moran and  
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Tetrahedron: Asymmetry 14 (2003) 43



C<sub>10</sub>H<sub>12</sub>O<sub>2</sub>

(R)-1-Hydroxy-1-phenylbutan-2-one

E.e. = 98%

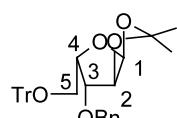
[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -325 (c 2, CHCl<sub>3</sub>)

Source of chirality: biocatalytic reduction

Absolute configuration: *R*

Aymeric Bordier, Philippe Compain, Olivier R. Martin,\*  
Kyoko Ikeda and Naoki Asano

Tetrahedron: Asymmetry 14 (2003) 47



C<sub>34</sub>H<sub>34</sub>O<sub>5</sub>

3-O-Benzyl-1,2-O-isopropylidene-5-O-triphenylmethyl- $\alpha$ -L-xylofuranose

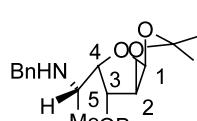
[ $\alpha$ ]<sub>D</sub> = +39 (c 1, CHCl<sub>3</sub>)

Source of chirality: L-xylose

Absolute configuration: 1*S*,2*S*,3*R*,4*S*

Aymeric Bordier, Philippe Compain, Olivier R. Martin,\*  
Kyoko Ikeda and Naoki Asano

Tetrahedron: Asymmetry 14 (2003) 47



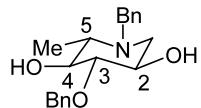
C<sub>23</sub>H<sub>29</sub>NO<sub>4</sub>

3-O-Benzyl-5-benzylamino-5,6-dideoxy-1,2-O-isopropylidene- $\alpha$ -L-glucofuranose

[ $\alpha$ ]<sub>D</sub> = +55 (c 1, CHCl<sub>3</sub>)

Source of chirality: L-xylose and asymmetric nucleophilic addition

Absolute configuration: 1*S*,2*S*,3*R*,4*S*,5*S*



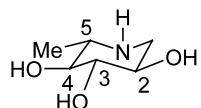
C<sub>20</sub>H<sub>25</sub>NO<sub>3</sub>

N-Benzyl-3-O-benzyl-1,5-imino-1,5,6-trideoxy-L-glucitol

[ $\alpha$ ]<sub>D</sub> = +55 (c 1, CHCl<sub>3</sub>)

Source of chirality: L-xylose and asymmetric nucleophilic addition

Absolute configuration: 2R,3S,4S,5S



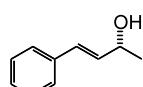
C<sub>6</sub>H<sub>13</sub>NO<sub>3</sub>

1,5-Imino-1,5,6-trideoxy-L-glucitol

[ $\alpha$ ]<sub>D</sub> = -12 (c 1.6, MeOH)

Source of chirality: L-xylose and asymmetric nucleophilic addition

Absolute configuration: 2R,3S,4S,5S



C<sub>10</sub>H<sub>12</sub>O

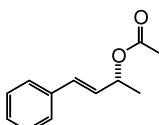
(R)-trans-4-Phenyl-3-butene-2-ol

Ee >99%

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = +19.9 (c 1, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: lipase-catalysed enantioselective acylation

Absolute configuration: 2R



C<sub>12</sub>H<sub>14</sub>O<sub>2</sub>

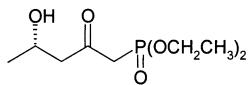
(R)-trans-4-Phenyl-3-butene-2 acetate

Ee >99%

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = +74.2 (c 1, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: lipase-catalysed enantioselective hydrolysis

Absolute configuration: 2R

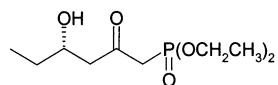


C<sub>9</sub>H<sub>19</sub>O<sub>5</sub>P  
(4S)-Diethyl 4-hydroxy-2-oxo-pentylphosphonate

$[\alpha]_D^{18} = +33.2$  (*c* 1.2, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4S

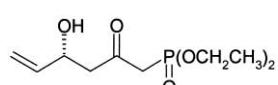


C<sub>10</sub>H<sub>21</sub>O<sub>5</sub>P  
(4S)-4-Diethyl 4-hydroxy-2-oxo-hexylphosphonate

$[\alpha]_D^{18} = +40$  (*c* 0.75, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4S

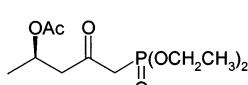


C<sub>10</sub>H<sub>19</sub>O<sub>5</sub>P  
(4R)-Diethyl 4-hydroxy-2-oxo-5-hexenylphosphonate

$[\alpha]_D^{18} = +17.5$  (*c* 0.85, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4R

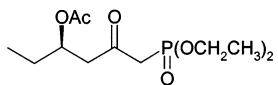


C<sub>11</sub>H<sub>21</sub>O<sub>6</sub>P  
(4R)-Diethyl 4-acetoxy-2-oxo-pentylphosphonate

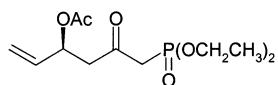
$[\alpha]_D^{18} = +4.0$  (*c* 0.6, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

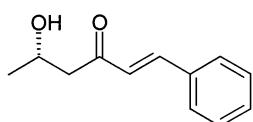
Absolute configuration: 4R

 $C_{12}H_{23}O_6P$ (4*R*)-Diethyl 4-acetyloxy-2-oxo-hexylphosphonate $[\alpha]_D^{18} = +11.5$  (*c* 0.6, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4*R* $C_{12}H_{21}O_6P$ (4*S*)-Diethyl 4-acetyloxy-2-oxo-5-hexenylphosphonate $[\alpha]_D^{18} = -1.3$  (*c* 0.85, CHCl<sub>3</sub>)

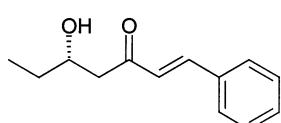
Source of chirality: enzymatic resolution

Absolute configuration: 4*S* $C_{12}H_{14}O_2$ (5*S*,1*E*)-5-Hydroxy-1-phenyl-1-hexen-3-one

E.e. = 99.1%

 $[\alpha]_D^{16} = +41$  (*c* 0.65, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

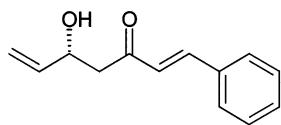
Absolute configuration: 5*S* $C_{13}H_{16}O_2$ (5*S*,1*E*)-5-Hydroxy-1-phenyl-1-hepten-3-one

E.e. = 95%

 $[\alpha]_D^{25} = +50.7$  (*c* 1.65, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

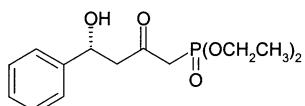
Absolute configuration: 5*S*

 $C_{13}H_{14}O_2$ (5*R*,1*E*)-5-Hydroxy-1-phenyl-1,6-heptadien-3-one

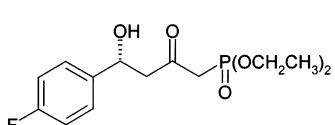
E.e. = 95%

 $[\alpha]_D^{16} = +33$  (*c* 0.25, CHCl<sub>3</sub>)

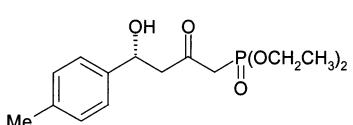
Source of chirality: enzymatic resolution

Absolute configuration: 5*R* $C_{14}H_{21}O_5P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-phenylbutylphosphonate $[\alpha]_D^{27} = +41.6$  (*c* 1.8, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

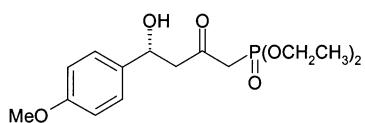
Absolute configuration: 4*R* $C_{14}H_{20}FO_5P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-(4-fluorophenyl)butylphosphonate $[\alpha]_D^{27} = +42.7$  (*c* 1.55, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

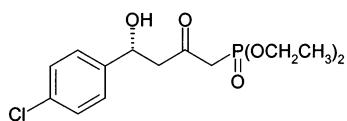
Absolute configuration: 4*R* $C_{15}H_{23}O_5P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-(4-methylphenyl)butylphosphonate $[\alpha]_D^{27} = +38.9$  (*c* 0.9, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

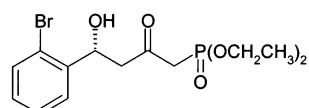
Absolute configuration: 4*R*

 $C_{15}H_{23}O_6P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-(4-methoxyphenyl)butylphosphonate $[\alpha]_D^{27} = +40.1$  (*c* 1.85, CHCl<sub>3</sub>)

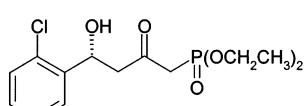
Source of chirality: enzymatic resolution

Absolute configuration: 4*R* $C_{14}H_{20}ClO_5P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-(4-chlorophenyl)butylphosphonate $[\alpha]_D^{27} = +41.2$  (*c* 1.95, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

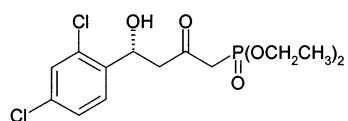
Absolute configuration: 4*R* $C_{14}H_{20}BrO_5P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-(2-bromophenyl)butylphosphonate $[\alpha]_D^{25} = +69.2$  (*c* 1.25, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4*R* $C_{14}H_{20}ClO_5P$ (4*R*)-Diethyl 4-hydroxy-2-oxo-4-(2-chlorophenyl)butylphosphonate $[\alpha]_D^{25} = +74.7$  (*c* 1.5, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4*R*

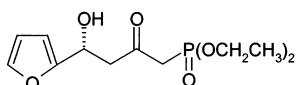


C<sub>14</sub>H<sub>19</sub>Cl<sub>2</sub>O<sub>5</sub>P  
(4*R*)-Diethyl 4-hydroxy-2-oxo-4-(2,4-dichlorophenyl)butylphosphonate

[ $\alpha$ ]<sub>D</sub><sup>27</sup> = +68.7 (*c* 0.7, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4*R*

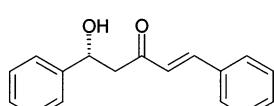


C<sub>12</sub>H<sub>19</sub>O<sub>5</sub>P  
(4*R*)-Diethyl 4-hydroxy-2-oxo-4-(2-furyl)butylphosphonate

[ $\alpha$ ]<sub>D</sub><sup>27</sup> = +28.1 (*c* 1.45, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 4*R*



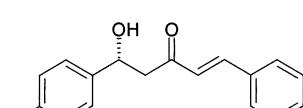
C<sub>17</sub>H<sub>16</sub>O<sub>2</sub>  
(5*R*,1*E*)-5-Hydroxy-1,5-diphenyl-1-penten-3-one

E.e. = 98.7%

[ $\alpha$ ]<sub>D</sub><sup>27</sup> = +67.1 (*c* 1.05, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 5*R*



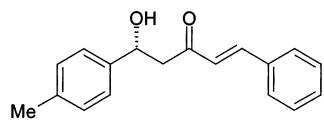
C<sub>17</sub>H<sub>15</sub>FO<sub>2</sub>  
(5*R*,1*E*)-5-Hydroxy-5-(4-fluorophenyl)-1-phenyl-1-penten-3-one

E.e. = 95.9%

[ $\alpha$ ]<sub>D</sub><sup>27</sup> = +59.9 (*c* 1.1, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

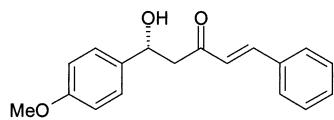
Absolute configuration: 5*R*

 $C_{18}H_{18}O_2$ (5*R*,1*E*)-5-Hydroxy-5-(4-methylphenyl)-1-phenyl-1-penten-3-one

E.e. = 100%

 $[\alpha]_D^{27} = +79.6$  (*c* 0.5, CHCl<sub>3</sub>)

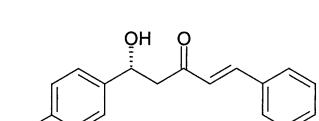
Source of chirality: enzymatic resolution

Absolute configuration: 5*R* $C_{18}H_{18}O_3$ (5*R*,1*E*)-5-Hydroxy-5-(4-methoxyphenyl)-1-phenyl-1-penten-3-one

E.e. = 96.8%

 $[\alpha]_D^{27} = +51.0$  (*c* 0.95, CHCl<sub>3</sub>)

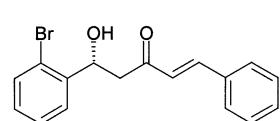
Source of chirality: enzymatic resolution

Absolute configuration: 5*R* $C_{17}H_{15}ClO_2$ (5*R*,1*E*)-5-Hydroxy-5-(4-chlorophenyl)-1-phenyl-1-penten-3-one

E.e. = 99.4%

 $[\alpha]_D^{27} = +58.6$  (*c* 1.3, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

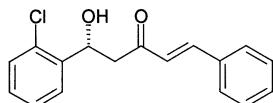
Absolute configuration: 5*R* $C_{17}H_{15}BrO_2$ (5*R*,1*E*)-5-Hydroxy-5-(2-bromophenyl)-1-phenyl-1-penten-3-one

E.e. = 98.0%

 $[\alpha]_D^{27} = +121.2$  (*c* 0.95, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

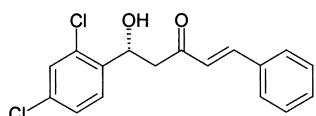
Absolute configuration: 5*R*

 $C_{17}H_{15}ClO_2$ (5*R*,1*E*)-5-Hydroxy-5-(2-chlorophenyl)-1-phenyl-1-penten-3-one

E.e. = 97.0%

 $[\alpha]_D^{27} = +68.9$  (*c* 0.35, CHCl<sub>3</sub>)

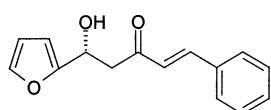
Source of chirality: enzymatic resolution

Absolute configuration: 5*R* $C_{17}H_{14}Cl_2O_2$ (5*R*,1*E*)-5-Hydroxy-5-(2,4-dichlorophenyl)-1-phenyl-1-penten-3-one

E.e. = 97.5%

 $[\alpha]_D^{27} = +92.0$  (*c* 1.0, CHCl<sub>3</sub>)

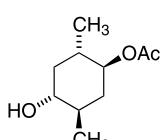
Source of chirality: enzymatic resolution

Absolute configuration: 5*R* $C_{15}H_{14}O_3$ (5*R*,1*E*)-5-Hydroxy-5-(2-furyl)-1-phenyl-1-penten-3-one

E.e. = 85.9%

 $[\alpha]_D^{27} = +33.7$  (*c* 0.7, CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

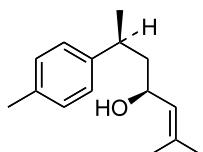
Absolute configuration: 5*R* $C_{10}H_{18}O_3$ (+)-(1*R*,2*R*,4*S*,5*S*)-4-Acetoxy-2,5-dimethyl-1-cyclohexanol

E.e. &gt;99.5%

 $[\alpha]_D = +48.0$  (*c* 0.42, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: enzymatic desymmetrization of centrosymmetric diacetate using pig liver esterase (PLE)

Absolute configuration: 1*R*,2*R*,4*S*,5*S*



$C_{15}H_{22}O$   
(7*S*,9*S*)-1-*epi*-Bisacumol

D.e. 92%

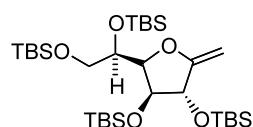
 $[\alpha]_D = +9.4$ , (*c* 9.7,  $CHCl_3$ )

Source of chirality: asymmetric synthesis

Absolute configuration: 7*S*,9*S*

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Tetrahedron: Asymmetry 14 (2003) 79



$C_{31}H_{68}O_5Si_4$   
2,5-Anhydro-3,4,6,7-tetra-*O*-(*tert*-butyldimethylsilyl)-1-deoxy-D-gluco-hept-1-enitol

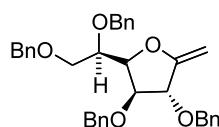
 $[\alpha]_D = +14.3$  (*c* 1.5,  $CHCl_3$ )

Source of chirality: D-gluconolactone starting material

Absolute configuration: D-gluco

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Tetrahedron: Asymmetry 14 (2003) 79



$C_{35}H_{36}O_5$   
2,5-Anhydro-3,4,6,7-tetra-*O*-benzyl-1-deoxy-D-gluco-hept-1-enitol

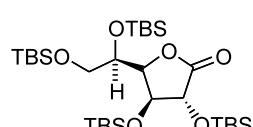
 $[\alpha]_D = +17.0$  (*c* 1.0,  $CHCl_3$ )

Source of chirality: D-gluconolactone starting material

Absolute configuration: D-gluco

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$C_{30}H_{66}O_6Si_4$   
2,3,5,6-Tetra-*O*-(*tert*-butyldimethylsilyl)-D-glucono-1,4-lactone

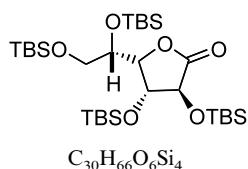
 $[\alpha]_D = +36.1$  (*c* 1.0,  $CHCl_3$ )

Source of chirality: D-gluconolactone starting material

Absolute configuration: D-gluco

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Duncan E. Paterson and Richard J. K. Taylor

Tetrahedron: Asymmetry 14 (2003) 79



2,3,5,6-Tetra-O-(*tert*-butyldimethylsilyl)-L-glucono-1,4-lactone

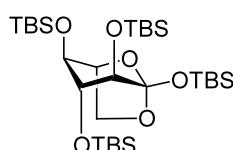
$[\alpha]_D = -36.1$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: L-gluconolactone starting material

Absolute configuration: L-gluco

Paul V. Murphy,\* Ciaran McDonnell, Ludger Hämig,  
Duncan E. Paterson and Richard J. K. Taylor

Tetrahedron: Asymmetry 14 (2003) 79



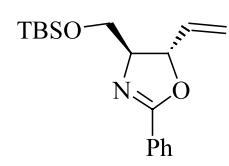
1-(*tert*-Butyldimethylsilyloxy)-2,3,4-tri-O-(*tert*-butyldimethylsilyl)-1,6-anhydro-L-glucopyranose

$[\alpha]_D = +15.4$  (*c* 0.9, CHCl<sub>3</sub>)

Source of chirality: L-gluconolactone starting material

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

Tetrahedron: Asymmetry 14 (2003) 87



(4*S*,*trans*)-4,5-Dihydro-4-(*tert*-butyl-dimethylsilyloxy)methyl)-2-phenyloxazoline

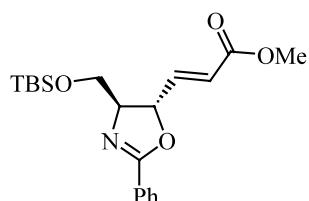
$[\alpha]_D^{25} -3.7$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular cyclization

Absolute configuration: 4*S,trans*

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

Tetrahedron: Asymmetry 14 (2003) 87



(2*E*)-3-((4*S*,*trans*)-4,5-Dihydro-4-(*tert*-butyl-dimethylsilyloxy)methyl)-2-phenyloxazol-5-yl)-acrylic acid methyl ester

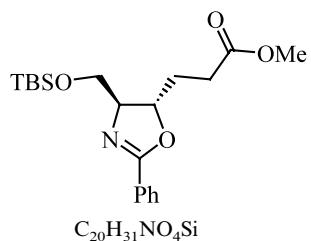
$[\alpha]_D^{24} +53.9$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular cyclization

Absolute configuration: 2*E*,4*S,trans*

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

Tetrahedron: Asymmetry 14 (2003) 87



3-((4*S,trans*)-4,5-Dihydro-4-(*tert*-butyl-dimethylsilanyloxymethyl)-2-phenyloxazol-5-yl)-propionic acid methyl ester

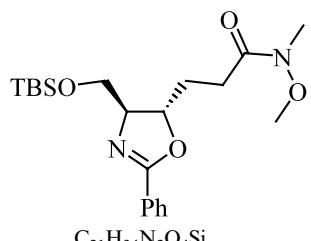
$[\alpha]_D^{24} -15.7$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular cyclization

Absolute configuration: 4*S,trans*

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

Tetrahedron: Asymmetry 14 (2003) 87



3-((4*S,trans*)-4,5-Dihydro-4-(*tert*-butyl-dimethylsilanyloxymethyl)-2-phenyloxazol-5-yl)-*N*-methoxy-*N*-methyl-propionamide

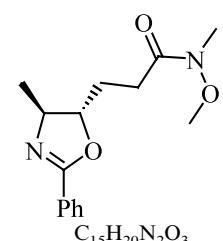
$[\alpha]_D^{24} -32.2$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular cyclization

Absolute configuration: 4*S,trans*

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

Tetrahedron: Asymmetry 14 (2003) 87



3-((4*S,trans*)-4,5-Dihydro-4-methyl-2-phenyloxazol-5-yl)-*N*-methoxy-*N*-methyl-propionamide

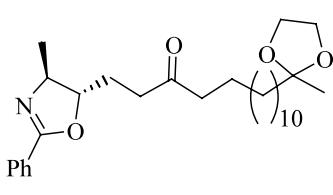
$[\alpha]_D^{25} -53.9$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular cyclization

Absolute configuration: 4*S,trans*

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

Tetrahedron: Asymmetry 14 (2003) 87



3-((4*S,trans*)-4,5-Dihydro-4-methyl-2-phenyloxazol-5-yl)-heptadecan-3-one-16-ethylenacetal

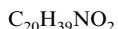
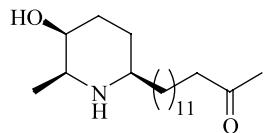
$[\alpha]_D^{23} -21.0$  (*c* 1.0, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular cyclization

Absolute configuration: 4*S,trans*

Yiu-Suk Lee, Yong-Ho Shin, Yong-Hyun Kim, Kee-Young Lee,  
Chang-Young Oh, Sung-Jae Pyun, Hyun-Ju Park, Jin-Hyun Jeong  
and Won-Hun Ham\*

*Tetrahedron: Asymmetry* 14 (2003) 87



(+)-Spectraline

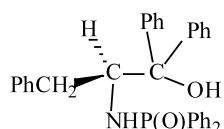
[ $\alpha$ ]<sub>D</sub><sup>26</sup> +8.8 (*c* 1.3, CHCl<sub>3</sub>)

Source of chirality: stereoselective intramolecular  
reductive amination

Absolute configuration: 2S,3S,6R

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95



(2S)-1,1,3-Triphenyl-2-(*N*-diphenylphosphinyl)amino-1-propanol

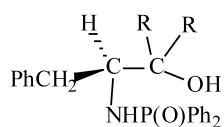
Mp=223~225°C

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=-31.1 (*c* 1.0, CHCl<sub>3</sub>)

Absolute configuration: 2S

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95



(2S)-1,1-Di(4-fluorophenyl)-2-(*N*-diphenylphosphinyl)amino-3-phenyl-1-propanol

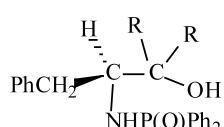
Mp=233~236°C

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=-28.7 (*c* 1.0, CHCl<sub>3</sub>)

Absolute configuration: 2S

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95



(2S)-1,1-Di(4-methylphenyl)-2-(*N*-diphenylphosphinyl)amino-3-phenyl-1-propanol

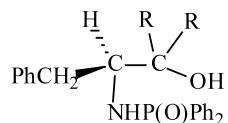
Mp=218~220°C

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=-22.5 (*c* 0.4, CHCl<sub>3</sub>)

Absolute configuration: 2S

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95

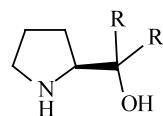


(2*S*)-3-Ethyl-2-(*N*-diphenylphosphinylamino)-1-phenyl-3-pentanol

Mp = 147 ~ 149°C  
 $[\alpha]_D^{20} = -69.5$  (*c* 1.0, CH<sub>2</sub>Cl<sub>2</sub>)  
 Absolute configuration: 2*S*

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95

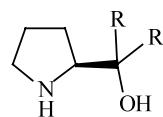


(2*S*)-2-[Di-(4-fluorophenyl)hydroxymethyl]pyrrolidine

Mp = 68 ~ 70°C  
 $[\alpha]_D^{20} = -57.0$  (*c* 0.73, CH<sub>2</sub>Cl<sub>2</sub>)  
 Absolute configuration: 2*S*

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95

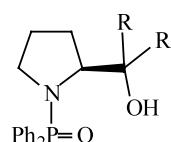


(2*S*)-2-[Di-(4-methylphenyl)hydroxymethyl]pyrrolidine

Mp = 93 ~ 94°C  
 $[\alpha]_D^{20} = -58.0$  (*c* 1.0, CHCl<sub>3</sub>)  
 Absolute configuration: 2*S*

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95

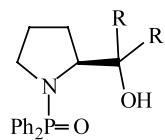


*N*-Diphenylphosphinyl-(2*S*)-2-[di(4-fluorophenyl)hydroxymethyl]pyrrolidine

Mp = 158 ~ 160°C  
 $[\alpha]_D^{20} = -44.4$  (*c* 1.1, CH<sub>2</sub>Cl<sub>2</sub>)  
 Absolute configuration: 2*S*

Kangying Li, Zhenghong Zhou, Lixin Wang, Qifa Chen,  
Guofeng Zhao, Qilin Zhou and Chuchi Tang\*

*Tetrahedron: Asymmetry* 14 (2003) 95

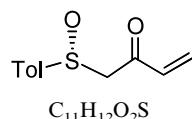


*N*-Diphenylphosphoryl-(2*S*)-2-[di(4-methylphenyl)hydroxymethyl]pyrrolidine

Mp = 158 ~ 160°C  
 $[\alpha]_D^{20} = -38.2$  (*c* 1.1, CHCl<sub>3</sub>)  
 Absolute configuration: 2*S*

Sadagopan Raghavan\* and S. C. Joseph

*Tetrahedron: Asymmetry* 14 (2003) 101

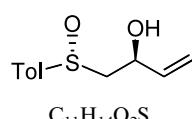


1(*R*<sub>S</sub>)-(4-Methylphenylsulfinyl)-3-buten-2-ol

Ee = 100%  
 $[\alpha]_D^{24} = +184.6$  (*c* 1, CHCl<sub>3</sub>)  
 Source of chirality: asymmetric synthesis  
 Absolute configuration: 1*R*<sub>S</sub>

Sadagopan Raghavan\* and S. C. Joseph

*Tetrahedron: Asymmetry* 14 (2003) 101

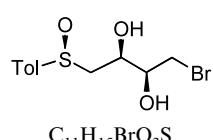


1(*R*<sub>S</sub>)-(4-Methylphenylsulfinyl)-(2*S*)-3-buten-2-ol

De >95%  
 $[\alpha]_D^{24} = +134.0$  (*c* 1, acetone)  
 Source of chirality: asymmetric synthesis  
 Absolute configuration: 1*R*<sub>S</sub>, 2*S*

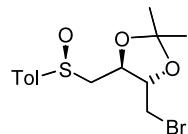
Sadagopan Raghavan\* and S. C. Joseph

*Tetrahedron: Asymmetry* 14 (2003) 101



1-Bromo-4(*S*<sub>S</sub>)-(4-methylphenylsulfinyl)-(2*S*,3*S*)-3-butane-2,3-diol

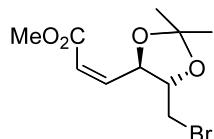
De >95%  
 $[\alpha]_D^{24} = -208.2$  (*c* 0.5, MeOH)  
 Source of chirality: asymmetric synthesis  
 Absolute configuration: 4*S*<sub>S</sub>, 2*S*, 3*S*

 $C_{14}H_{19}BrO_3S$ 4-Bromomethyl-2,2-dimethyl-5-(4-methyl-(*S*)-phenylsulfinylmethyl)-(4*S*,5*S*)-1,3-dioxolane

De &gt;95%

 $[\alpha]_D^{24} = -202.3$  (*c* 1, CHCl<sub>3</sub>)

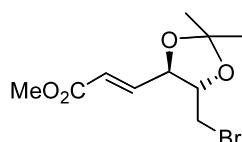
Source of chirality: asymmetric synthesis

Absolute configuration: *S*<sub>S</sub>,4*S*,5*S* $C_{10}H_{16}BrO_4$ Methyl 3-[5-bromomethyl-2,2-dimethyl-(4*R*,5*S*)-1,3-dioxolan-4-yl]-(*Z*)-2-propenoate

Ee &gt;95%

 $[\alpha]_D^{24} = +118.5$  (*c* 0.14, CHCl<sub>3</sub>)

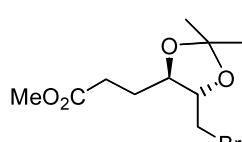
Source of chirality: asymmetric synthesis

Absolute configuration: 4*R*,5*S* $C_{10}H_{16}BrO_4$ Methyl 3-[5-bromomethyl-2,2-dimethyl-(4*R*,5*S*)-1,3-dioxolan-4-yl]-(*E*)-2-propenoate

Ee &gt;95%

 $[\alpha]_D^{24} = -9.2$  (*c* 0.42, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

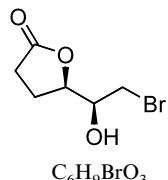
Absolute configuration: 4*R*,5*S* $C_{10}H_{18}BrO_4$ Methyl 3-[5-bromomethyl-2,2-dimethyl-(4*R*,5*S*)-1,3-dioxolan-4-yl]propanoate

Ee &gt;95%

 $[\alpha]_D^{24} = -15.7$  (*c* 1.38, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

Absolute configuration: 4*R*,5*S*

5-[2-Bromo-1-hydroxy-(1*S*)-ethyl]-(5*R*)-2*H*,3*H*,4*H*-2-furanone

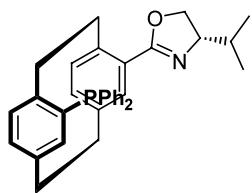
Ee &gt;95%

 $[\alpha]_D^{24} = -35.6$  (*c* 0.19, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

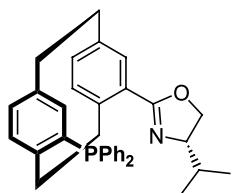
Absolute configuration: 1*S*,5*R*

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and Xue-Long Hou\*

C34H34NOP(S,4*R*<sub>p</sub>,13*S*<sub>p</sub>)-4-Diphenylphosphino-13-(4-*iso*-propyloxazoline-2-yl)[2.2]paracyclophane $[\alpha]_D^{20} = -92.7$  (*c* 0.575, CHCl<sub>3</sub>)

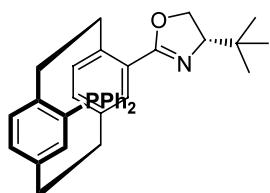
Source of chirality: (S)-valinol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and Xue-Long Hou\*

C34H34NOP(S,4*S*<sub>p</sub>,13*R*<sub>p</sub>)-4-Diphenylphosphino-13-(4-*iso*-propyloxazoline-2-yl)[2.2]paracyclophane $[\alpha]_D^{20} = +4.9$  (*c* 0.65, CHCl<sub>3</sub>)

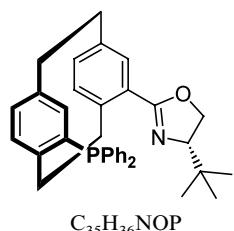
Source of chirality: (S)-valinol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and Xue-Long Hou\*

C35H36NOP(S,4*R*<sub>p</sub>,13*S*<sub>p</sub>)-4-Diphenylphosphino-13-(4-*tert*-butyloxazoline-2-yl)[2.2]paracyclophane $[\alpha]_D^{20} = -80.6$  (*c* 0.565, CHCl<sub>3</sub>)Source of chirality: (S)-*tert*-leucinol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and  
Xue-Long Hou\*

Tetrahedron: Asymmetry 14 (2003) 107



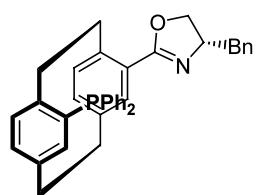
(*S,4S<sub>p</sub>,13R<sub>p</sub>*)-4-Diphenylphosphino-13-(4-*tert*-butyloxazoline-2-yl)[2.2]paracyclophane

$[\alpha]_D^{20} = +21.0$  (*c* 0.56, CHCl<sub>3</sub>)

Source of chirality: (*S*)-*tert*-leucinol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and  
Xue-Long Hou\*

Tetrahedron: Asymmetry 14 (2003) 107



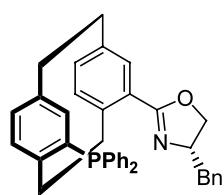
(*S,4R<sub>p</sub>,13S<sub>p</sub>*)-4-Diphenylphosphino-13-(4-benzyloxazoline-2-yl)[2.2]paracyclophane

$[\alpha]_D^{20} = -47.5$  (*c* 0.46, CHCl<sub>3</sub>)

Source of chirality: (*S*)-phenylanol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and  
Xue-Long Hou\*

Tetrahedron: Asymmetry 14 (2003) 107



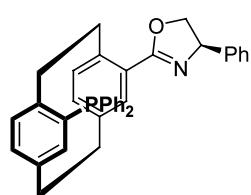
(*S,4S<sub>p</sub>,13R<sub>p</sub>*)-4-Diphenylphosphino-13-(4-benzyloxazoline-2-yl)[2.2]paracyclophane

$[\alpha]_D^{20} = +8.2$  (*c* 0.75, CHCl<sub>3</sub>)

Source of chirality: (*S*)-phenylanol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and  
Xue-Long Hou\*

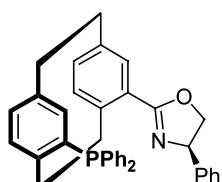
Tetrahedron: Asymmetry 14 (2003) 107



(*R,4R<sub>p</sub>,13S<sub>p</sub>*)-4-Diphenylphosphino-13-(4-phenyloxazoline-2-yl)[2.2]paracyclophane

$[\alpha]_D^{20} = -23.3$  (*c* 0.535, CHCl<sub>3</sub>)

Source of chirality: (*R*)-phenylglycinol



C<sub>37</sub>H<sub>32</sub>NOP

(R,4S<sub>p</sub>,13R<sub>p</sub>)-4-Diphenylphosphino-13-(4-phenyloxazoline-2-yl)[2.2]paracyclophane

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = +46.7 (c 0.525, CHCl<sub>3</sub>)

Source of chirality: (R)-phenylglycinol

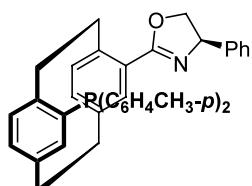


C<sub>39</sub>H<sub>36</sub>NOP

(R,4R<sub>p</sub>,13S<sub>p</sub>)-4-Di(o-toyl)phosphino-13-(4-phenyloxazoline-2-yl)[2.2]paracyclophane

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -58.8 (c 0.325, CHCl<sub>3</sub>)

Source of chirality: (R)-phenylglycinol



C<sub>39</sub>H<sub>36</sub>NOP

(R,4R<sub>p</sub>,13S<sub>p</sub>)-4-Di(p-toyl)phosphino-13-(4-phenyloxazoline-2-yl)[2.2]paracyclophane

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -29.5 (c 0.40, CHCl<sub>3</sub>)

Source of chirality: (R)-phenylglycinol



C<sub>39</sub>H<sub>36</sub>NO<sub>3</sub>P

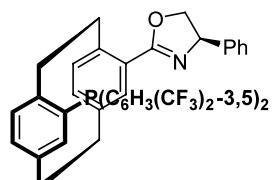
(R,4R<sub>p</sub>,13S<sub>p</sub>)-4-Di(p-methoxyphenyl)phosphino-13-(4-phenyloxazoline-2-yl)[2.2]paracyclophane

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -31.2 (c 0.32, CHCl<sub>3</sub>)

Source of chirality: (R)-phenylglycinol

Xun-Wei Wu, Ke Yuan, Wei Sun, Ming-Jie Zhang and Xue-Long Hou\*

*Tetrahedron: Asymmetry* 14 (2003) 107



$\text{C}_{41}\text{H}_{28}\text{F}_{12}\text{NOP}$

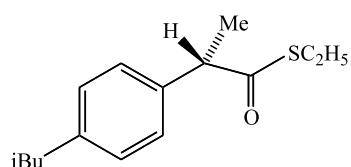
(*R*,4*R*<sub>p</sub>,13*S*<sub>p</sub>)-4-Di(3,5-di(trifluoromethyl)phenyl)phosphino-13-(4-phenyloxazoline-2-yl)[2.2]paracyclophane

$[\alpha]_D^{20} = +10.6$  (*c* 0.355,  $\text{CHCl}_3$ )

Source of chirality: (*R*)-phenylglycinol

Marco Clericuzio,\* Iacopo Degani,\* Stefano Dughera and Rita Fochi

*Tetrahedron: Asymmetry* 14 (2003) 119



$\text{C}_{15}\text{H}_{22}\text{OS}$

*S*-Ethyl (*S*)-2-(4-isobutylphenyl)thiopropionate

E.e. = 78%

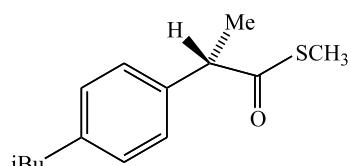
$[\alpha]_D^{22} = +73.3$  (*c* 0.4,  $\text{CH}_2\text{Cl}_2$ )

Source of chirality: (*R*)-BINOL· $\text{SnCl}_4$  complex (asymmetric protonation)

Absolute configuration: *S*

Marco Clericuzio,\* Iacopo Degani,\* Stefano Dughera and Rita Fochi

*Tetrahedron: Asymmetry* 14 (2003) 119



$\text{C}_{14}\text{H}_{20}\text{OS}$

*S*-Methyl (*S*)-2-(4-isobutylphenyl)thiopropionate

E.e. = 82%

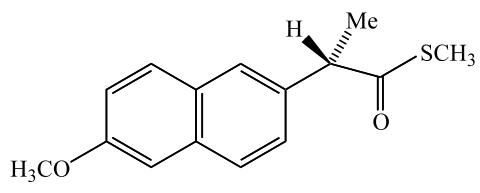
$[\alpha]_D^{22} = +80.5$  (*c* 0.35,  $\text{CH}_2\text{Cl}_2$ )

Source of chirality: (*R*)-BINOL· $\text{SnCl}_4$  complex (asymmetric protonation)

Absolute configuration: *S*

Marco Clericuzio,\* Iacopo Degani,\* Stefano Dughera and Rita Fochi

*Tetrahedron: Asymmetry* 14 (2003) 119



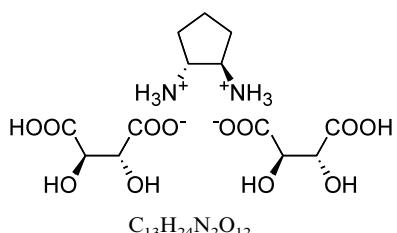
$\text{C}_{15}\text{H}_{16}\text{O}_2\text{S}$   
*S*-Methyl (*S*)-2-(6-methoxy-2-naphthyl)thiopropionate

E.e. = 64%

$[\alpha]_D^{22} = +81.2$  (*c* 0.6,  $\text{CH}_2\text{Cl}_2$ )

Source of chirality: (*R*)-BINOL· $\text{SnCl}_4$  complex (asymmetric protonation)

Absolute configuration: *S*

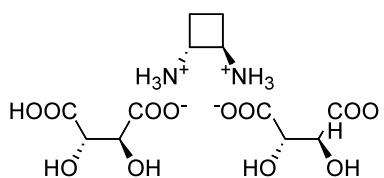


(R,R)-(-)-trans-Cyclopentane-1,2-diamine di-(+)-tartrate

Ee &gt;99%

 $[\alpha]_D^{20} = +10.0$  (*c* 2, H<sub>2</sub>O, lit. +10.1°)

Source of chirality: resolution

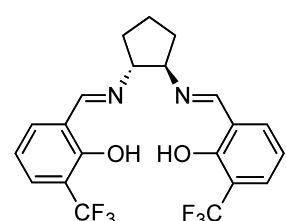


(+)-trans-Cyclobutane-1,2-diamine di-(+)-tartrate

Ee &gt;99%

 $[\alpha]_D^{20} = +28.0$  (*c* 1, H<sub>2</sub>O)

Source of chirality: resolution

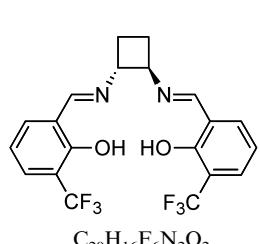


(R,R)-(-)-N,N'-Bis(3-trifluoromethylsalicylidene)-trans-cyclopentane-1,2-diamine

Ee &gt;99%

 $[\alpha]_D^{20} = -437$  (*c* 1, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: resolution

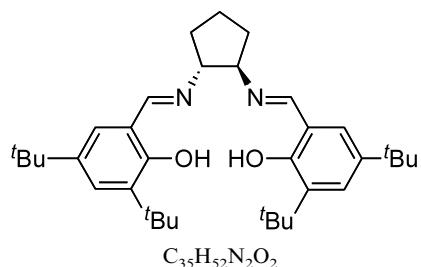


(+)-N,N'-Bis(3-trifluoromethylsalicylidene)-trans-cyclobutane-1,2-diamine

Ee &gt;99%

 $[\alpha]_D^{20} = +485$  (*c* 1, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: resolution

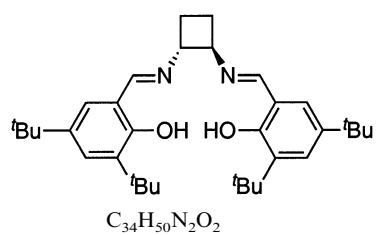


(R,R)-(-)-N,N'-Bis(3,5-di-tert-butylsalicylidene)-trans-cyclopentane-1,2-diamine

Ee &gt;99%

 $[\alpha]_D^{20} = -365$  (*c* 1, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: resolution

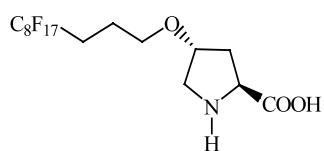


(+)-N,N'-Bis(3,5-di-tert-butylsalicylidene)-trans-cyclobutane-1,2-diamine

Ee &gt;99%

 $[\alpha]_D^{20} = +400$  (*c* 1, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: resolution



$C_{16}H_{15}F_{17}NO_3$   
trans-4-(Perfluoroctyl)propyloxy-L-proline

 $[\alpha]_D^{25} = -11.7$  (*c* 0.47, EtOH)Source of chirality: *trans*-4-hydroxy-L-proline

Absolute configuration: 2S,4R