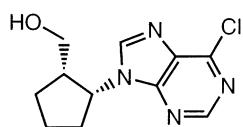


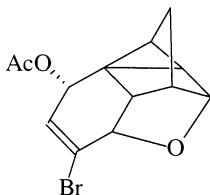
Elias Quezada, Lourdes Santana and Eugenio Uriarte\*

*Tetrahedron: Asymmetry* 12 (2001) 2637cis-6-Chloro-9-[2-(hydroxymethyl)cyclopentyl]-9*H*-purine $[\alpha]_D = -38.5$  (*c* 0.002, MeOH)

Source of chirality: resolution

Absolute configuration: (1'R,2'R)

Fernando D. P. Morisso and Valentim E. U. Costa\*

*Tetrahedron: Asymmetry* 12 (2001) 2641(-)-3-*endo*-Acetoxy-5-bromo-12-oxa-pentacyclo[6.2.1.1<sup>6,9</sup>.0<sup>2,7</sup>.0<sup>2,10</sup>]dodeca-4-ene

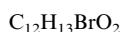
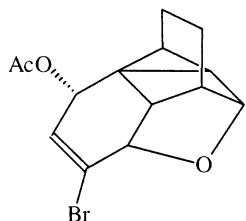
E.e. &gt;99% (by chiral GC)

 $[\alpha]_{D}^{20} = -190$  (*c* 0.5, ethyl acetate)

Source of chirality: enzyme-catalyzed transesterification of racemic mixture

Absolute configuration: unknown

Fernando D. P. Morisso and Valentim E. U. Costa\*

*Tetrahedron: Asymmetry* 12 (2001) 2641(-)-3-*endo*-Acetoxy-5-bromo-13-oxa-pentacyclo[6.2.1.1<sup>6,9</sup>.0<sup>2,7</sup>.0<sup>2,10</sup>]trideca-4-ene

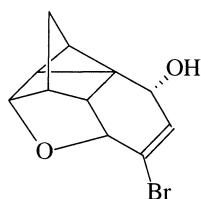
E.e. &gt;99% (by chiral GC)

 $[\alpha]_{D}^{20} = -486$  (*c* 0.5, ethyl acetate)

Source of chirality: enzyme-catalyzed transesterification of racemic mixture

Absolute configuration: unknown

Fernando D. P. Morisso and Valentim E. U. Costa\*

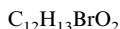
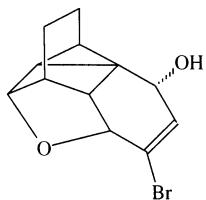
*Tetrahedron: Asymmetry* 12 (2001) 2641(-)-5-Bromo-12-oxa-pentacyclo[6.2.1.1<sup>6,9</sup>.0<sup>2,7</sup>.0<sup>2,10</sup>]dodeca-4-ene-3-*endo*-ol

E.e. &gt;99% (by chiral NMR and GC of the derived acetate (+)-5)

 $[\alpha]_{D}^{20} = +60$  (*c* 0.5, ethyl acetate)

Source of chirality: enzyme-catalyzed transesterification of racemic mixture

Absolute configuration: unknown

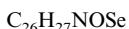
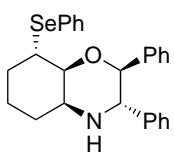
(+) -5-Bromo-13-oxa-pentacyclo[6.2.1.1<sup>6.9</sup>.0<sup>2.7</sup>.0<sup>2.10</sup>]trideca-4-ene-3-*endo*-ol

E.e. &gt;99% (by chiral GC)

 $[\alpha]_{D}^{20} = +140$  (*c* 0.5, ethyl acetate)

Source of chirality: enzyme-catalyzed transesterification of racemic mixture

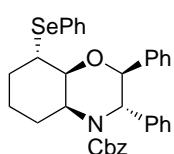
Absolute configuration: unknown

Kwan Soo Kim,\* Sung Ook Choi, Jong Myun Park, Yong Joo Lee  
and Jin Hwan Kim

(2S,3S,8S,9S,10S)-2,3-Diphenyl-8-phenylselenenyloctahydrobenzo-1,4-oxazine

 $[\alpha]_D = -32.6$  (*c* 0.54, CHCl<sub>3</sub>)

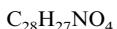
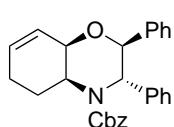
Source of chirality: asymmetric synthesis

Kwan Soo Kim,\* Sung Ook Choi, Jong Myun Park, Yong Joo Lee  
and Jin Hwan Kim

(2S,3S,8S,9S,10S)-4-Benzoyloxycarbonyl-2,3-diphenyl-8-phenylselenenyloctahydrobenzo-1,4-oxazine

 $[\alpha]_D = -39.6$  (*c* 0.56, CHCl<sub>3</sub>)

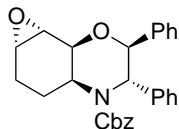
Source of chirality: asymmetric synthesis

Kwan Soo Kim,\* Sung Ook Choi, Jong Myun Park, Yong Joo Lee  
and Jin Hwan Kim

(2S,3S,9R,10S)-4-Benzoyloxycarbonyl-2,3-diphenyl-2,3;5,6;9,10-hexahydrobenzo-1,4-oxazine

 $[\alpha]_D = -16.9$  (*c* 0.52, CHCl<sub>3</sub>)

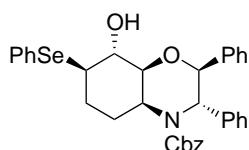
Source of chirality: asymmetric synthesis



C<sub>28</sub>H<sub>27</sub>NO<sub>3</sub>  
(5S,6S,8R,9S,10S,11S)-4-Benzylloxycarbonyl-5,6-diphenyloctahydro-1,7-dioxa-4-azacyclopropa[a]naphthalene

[ $\alpha$ ]<sub>D</sub> = -14.4 (c 0.50, CHCl<sub>3</sub>)

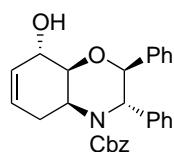
Source of chirality: asymmetric synthesis



C<sub>34</sub>H<sub>33</sub>NO<sub>4</sub>Se  
(2S,3S,7R,8R,9S,10S)-4-Benzylloxycarbonyl-8-hydroxy-2,3-diphenyl-7-phenylselenenyloctahydrobenzo-1,4-oxazine

[ $\alpha$ ]<sub>D</sub> = -40.9 (c 0.55, CHCl<sub>3</sub>)

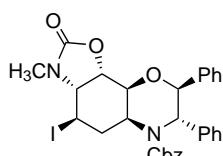
Source of chirality: asymmetric synthesis



C<sub>28</sub>H<sub>27</sub>NO<sub>4</sub>  
(2S,3S,8S,9S,10S)-4-Benzylloxycarbonyl-8-hydroxy-2,3-diphenyl-2,3;5,10;8,9-hexahydrobenzo-1,4-oxazine

[ $\alpha$ ]<sub>D</sub> = -3.9 (c 0.54, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis



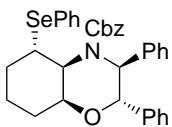
C<sub>30</sub>H<sub>29</sub>IN<sub>2</sub>O<sub>5</sub>  
(4R,7S,8S,10S,11R,12S,13S)-6-Benzylloxycarbonyl-4-iodo-3-methyl-2-oxo-7,8-diphenyldecahydro-1,9-dioxa-3,6-diazacyclopenta[a]naphthalene

[ $\alpha$ ]<sub>D</sub> = -3.3 (c 0.51, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

Kwan Soo Kim,\* Sung Ook Choi, Jong Myun Park, Yong Joo Lee  
and Jin Hwan Kim

*Tetrahedron: Asymmetry* 12 (2001) 2649



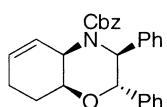
C<sub>34</sub>H<sub>33</sub>NO<sub>3</sub>Se  
(2S,3S,5S,9S,10S)-4-Benzyl-2,3-diphenyl-5-phenylselenenoxyoctahydrobenzo-1,4-oxazine

[ $\alpha$ ]<sub>D</sub> = -49.6 (*c* 1.20, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

Kwan Soo Kim,\* Sung Ook Choi, Jong Myun Park, Yong Joo Lee  
and Jin Hwan Kim

*Tetrahedron: Asymmetry* 12 (2001) 2649



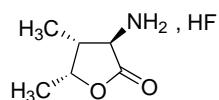
C<sub>28</sub>H<sub>27</sub>NO<sub>3</sub>  
(2S,3S,9S,10R)-4-Benzyl-2,3-diphenyl-2,3,7,8,9,10-hexahydrobenzo-1,4-oxazine

[ $\alpha$ ]<sub>D</sub> = -141.0 (*c* 0.50, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

Tarek Kassem, Jonhy Wehbe, Valérie Rolland-Fulcrand,  
Marc Rolland, Marie-Louise Roumestant\* and Jean Martinez

*Tetrahedron: Asymmetry* 12 (2001) 2657



C<sub>6</sub>H<sub>12</sub>FNO<sub>2</sub>  
(3R,4R,5R)-4-Hydroxy isoleucine lactone

E.e. = 99±1%

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = +51.2 (*c* 0.43, MeOH)

Source of chirality: oxazinone derived from  
(1*R*,2*R*,5*R*)-2-hydroxypinan-3-one

Francesca Clerici, Maria Luisa Gelmi,\* Donato Pocar and  
Tullio Pilati

*Tetrahedron: Asymmetry* 12 (2001) 2663



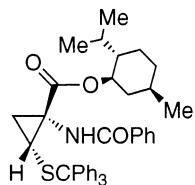
C<sub>39</sub>H<sub>41</sub>NO<sub>3</sub>S  
(-)Menthyl-2-benzoylamino-3-tritylsulfanylacrylate

Mp = 198°C (CH<sub>2</sub>Cl<sub>2</sub>/iPr<sub>2</sub>O)

Ee = 100%

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -88.6 (*c* 5.2×10<sup>-3</sup>, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis



C<sub>40</sub>H<sub>43</sub>NO<sub>3</sub>S

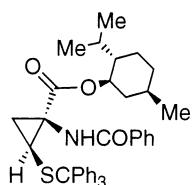
(-)-Menthyl (1*R*,2*R*)-1-benzoylamino-2-tritylsulfanyl-cyclopropylcarboxylate

Mp = 179°C (CH<sub>2</sub>Cl<sub>2</sub>/iPr<sub>2</sub>O)

De = 100%

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = +152 (*c* 2.8 × 10<sup>-3</sup>, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis



C<sub>40</sub>H<sub>43</sub>N<sub>3</sub>O<sub>3</sub>S

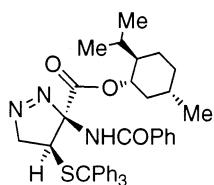
(-)-Menthyl (1*S*,2*S*)-1-benzoylamino-2-tritylsulfanyl-cyclopropylcarboxylate

Mp = 205°C (CH<sub>2</sub>Cl<sub>2</sub>/iPr<sub>2</sub>O)

De = 100%

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -235 (*c* 2.8 × 10<sup>-3</sup>, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis



C<sub>40</sub>H<sub>43</sub>N<sub>3</sub>O<sub>3</sub>S

(-)-Menthyl (3*R*,4*S*)-3-benzoylamino-4-tritylsulfanyl-4,5-dihydro-3*H*-pyrazole-3-carboxylate

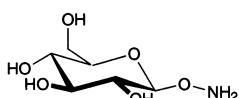
Oil

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -125.4 (*c* 2.1 × 10<sup>-3</sup>, CHCl<sub>3</sub>)

Source of chirality: asymmetric synthesis

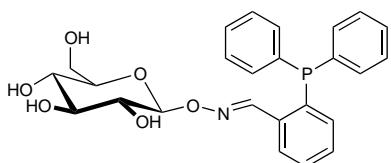
[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -40.0 (*c* 1, H<sub>2</sub>O)

Source of chirality: homochiral starting material

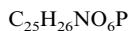
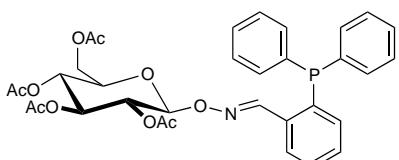


C<sub>6</sub>H<sub>13</sub>NO<sub>6</sub>

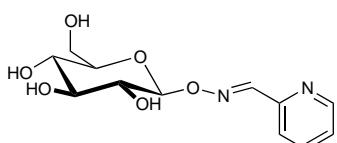
O-β-D-Glucopyranosylhydroxylamine


 $[\alpha]_D^{25} = -1.8 \text{ (c 3, CH}_2\text{Cl}_2)$ 

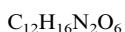
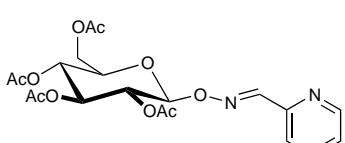
Source of chirality: homochiral starting material

*O*-( $\beta$ -D-Glucopyranosyl)-2-diphenylphosphanylbenzaldoxime
 $[\alpha]_D^{25} = -17.3 \text{ (c 3, CH}_2\text{Cl}_2)$ 

Source of chirality: homochiral starting material

*O*-(2,3,4,6-Tetra-*O*-acetyl- $\beta$ -D-glucopyranosyl)-2-diphenylphosphanylbenzaldoxime
 $[\alpha]_D^{25} = -16.3 \text{ (c 2, CH}_3\text{OH)}$ 

Source of chirality: homochiral starting material

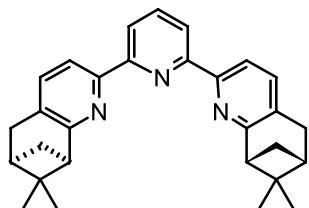
*O*-( $\beta$ -D-Glucopyranosyl)pyridine-2-carbaldoxime
 $[\alpha]_D^{25} = -22.0 \text{ (c 2, CH}_2\text{Cl}_2)$ 

Source of chirality: homochiral starting material

*O*-(2,3,4,6-Tetra-*O*-acetyl- $\beta$ -D-glucopyranosyl)pyridine-2-carbaldoxime

Hoi-Lun Kwong,\* Wing-Leung Wong, Wing-Sze Lee,  
Leung-Shi Cheng and Wing-Tak Wong

*Tetrahedron: Asymmetry* 12 (2001) 2683



C<sub>29</sub>H<sub>31</sub>N<sub>3</sub>

2,6-Bis(7,7-dimethyl-5,6,7,8-tetrahydro-6,8-methanoquinolin-2-yl)pyridine

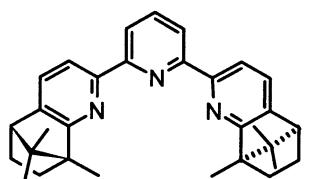
[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -49.4 (*c* = 0.50, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: (1*R*)-(+)-nopolone

Absolute configuration: 6*R*,8*R*

Hoi-Lun Kwong,\* Wing-Leung Wong, Wing-Sze Lee,  
Leung-Shi Cheng and Wing-Tak Wong

*Tetrahedron: Asymmetry* 12 (2001) 2683



C<sub>31</sub>H<sub>35</sub>N<sub>3</sub>

2,6-Bis(8,9,9-trimethyl-5,6,7,8-tetrahydro-5,8-methanoquinolin-2-yl)pyridine

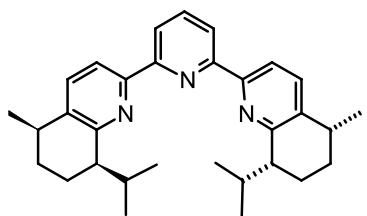
[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -15.8 (*c* = 0.53, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: (1*R*)-(+)-camphor

Absolute configuration: 5*S*,8*R*

Hoi-Lun Kwong,\* Wing-Leung Wong, Wing-Sze Lee,  
Leung-Shi Cheng and Wing-Tak Wong

*Tetrahedron: Asymmetry* 12 (2001) 2683



C<sub>31</sub>H<sub>39</sub>N<sub>3</sub>

2,6-Bis(5-methyl-8-isopropyl-5,6,7,8-tetrahydroquinolin-2-yl)pyridine

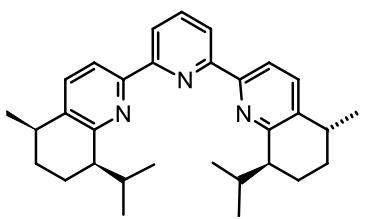
[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -58.8 (*c* = 0.50, CH<sub>2</sub>Cl<sub>2</sub>), mp 155–158°C

Source of chirality: (-)-menthone

Absolute configuration: 5*R*,8*R*

Hoi-Lun Kwong,\* Wing-Leung Wong, Wing-Sze Lee,  
Leung-Shi Cheng and Wing-Tak Wong

*Tetrahedron: Asymmetry* 12 (2001) 2683



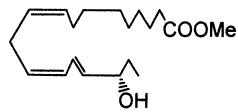
C<sub>31</sub>H<sub>39</sub>N<sub>3</sub>

2,6-Bis(5-methyl-8-isopropyl-5,6,7,8-octahydroquinolin-2-yl)-pyridine

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -49.0 (*c* = 0.53, CH<sub>2</sub>Cl<sub>2</sub>), mp 124–125°C

Source of chirality: (-)-menthone

Absolute configuration: 5*R*,5*'R*,8*R*,8*'S*



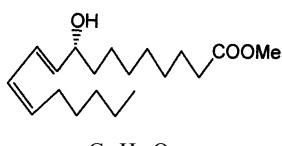
(16*S*)-Hydroxy- $\alpha$ -linolenic acid methyl ester

E.e.=97% (by NMR of MTPA ester of related bromohydrin of  $\alpha$ -linolenic acid methyl ester)

[ $\alpha$ ]<sub>D</sub><sup>27</sup>=+19 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: resolution with Lipase PS of ( $\pm$ )-bromohydrin of  $\alpha$ -linolenic acid methyl ester

Absolute configuration: 16*S* (assigned by Kusumi-Mosher method)



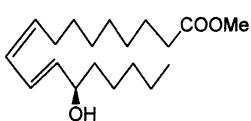
(9*R*)-Hydroxylinoleic acid methyl ester

E.e. >98% (by chiral HPLC of benzoate)

[ $\alpha$ ]<sub>D</sub><sup>29</sup>=-8.4 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: partial resolution of DL-alcohol followed by chiral HPLC separation of the benzoate

Absolute configuration: 9*R* (assigned by optical rotation of the benzoate)



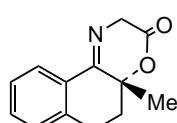
(13*R*)-Hydroxylinoleic acid methyl ester

E.e. >98% (by chiral column of benzoate)

[ $\alpha$ ]<sub>D</sub><sup>30</sup>=-8.6 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: partial resolution of DL-alcohol followed by chiral HPLC separation of the benzoate

Absolute configuration: 13*R* (assigned by optical rotation of the benzoate)



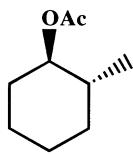
(4a*R*)-4a-Methyl-2,4a,5,6-tetrahydro-3H-naphtho[2,1-*b*][1,4]oxazin-2-one

E.e. >99%

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=+298 (*c*=1, CHCl<sub>3</sub>)

Source of chirality: resolution via supercritical fluid chromatography on a semi-preparative Chiraldak AS column

Absolute configuration: *R* (using VCD method)



C<sub>9</sub>H<sub>16</sub>O<sub>2</sub>

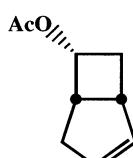
*trans*-2-Methylcyclohexanol acetate

Ee = 100% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl- $\beta$ -cyclodextrin in OV 1701]

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -69 (c 0.64, EtOH)

Source of chirality: microbial hydrolysis

Absolute configuration: 1*R*,2*R*



C<sub>9</sub>H<sub>12</sub>O<sub>2</sub>

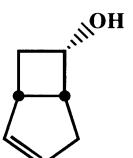
*endo*-Bicyclo[3.2.0]hept-2-en-6-ol acetate

Ee >95% [by GLC analysis on a 25 m dimethyl-*n*-pentyl- $\beta$ -cyclodextrin in OV 1701]

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = 35 (c 2.27, CHCl<sub>3</sub>)

Source of chirality: microbial hydrolysis

Absolute configuration: 1*R*,5*S*,6*R*



C<sub>7</sub>H<sub>10</sub>O

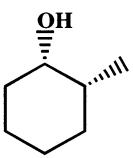
*endo*-Bicyclo[3.2.0]hept-2-en-6-ol

Ee = 67% [by GLC analysis on a 25 m dimethyl-*n*-pentyl- $\beta$ -cyclodextrin in OV 1701]

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = 45 (c 1.1, CHCl<sub>3</sub>)

Source of chirality: microbial hydrolysis

Absolute configuration: 1*S*,5*R*,6*S*



C<sub>7</sub>H<sub>14</sub>O

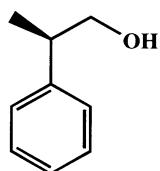
*cis*-2-Methylcyclohexanol

Ee = 100% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl- $\beta$ -cyclodextrin in OV 1701]

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = 18 (c 1, MeOH)

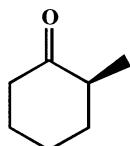
Source of chirality: microbial hydrolysis and reduction

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



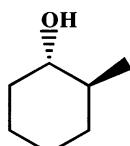
C<sub>9</sub>H<sub>12</sub>O  
2-Phenylpropanol

Ee = 57% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl- $\beta$ -cyclodextrin in OV 1701]  
 $[\alpha]_D^{25} = -9.7$  (neat)  
Source of chirality: microbial hydrolysis and reduction  
Absolute configuration: *S*



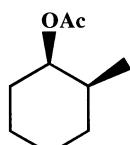
C<sub>7</sub>H<sub>12</sub>O  
2-Methylcyclohexanone

Ee = 100% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl- $\beta$ -cyclodextrin in OV 1701]  
 $[\alpha]_D^{25} = -14$  (*c* 0.23, MeOH)  
Source of chirality: microbial hydrolysis  
Absolute configuration: *S*



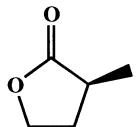
C<sub>7</sub>H<sub>14</sub>O  
trans-2-Methylcyclohexanol

Ee = 39% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl- $\beta$ -cyclodextrin in OV 1701]  
 $[\alpha]_D^{25} = 14.9$  (*c* 9.6, EtOH)  
Source of chirality: microbial hydrolysis  
Absolute configuration: 1*S*,2*S*



C<sub>9</sub>H<sub>16</sub>O<sub>2</sub>  
cis-2-Methylcyclohexanol acetate

Ee = 100% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl- $\beta$ -cyclodextrin in OV 1701]  
 $[\alpha]_D^{25} = -37.4$  (*c* 1.50, CHCl<sub>3</sub>)  
Source of chirality: microbial hydrolysis  
Absolute configuration: 1*R*,2*S*



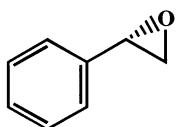
α-Methyl-γ-butyrolactone

Ee = 40% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl-β-cyclodextrin in OV 1701]

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = -8.9 (c 6.5, EtOH)

Source of chirality: microbial hydrolysis

Absolute configuration: *S*



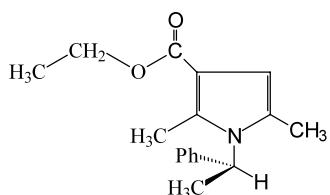
Styrene oxide

Ee = 52% [by GLC analysis on a 25 m diethyl-*tert*-butylsilyl-β-cyclodextrin in OV 1701]

[ $\alpha$ ]<sub>D</sub><sup>25</sup> = 17.1 (neat)

Source of chirality: microbial hydrolysis

Absolute configuration: *R*

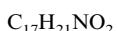


E.e. = 98%

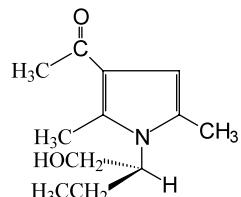
[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -35.8 (c 0.4, CHCl<sub>3</sub>)

Source of chirality: (*S*)-1-phenylethylamine as starting material

Absolute configuration: *S*



(*S*)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)pentanedioic acid diethyl ester



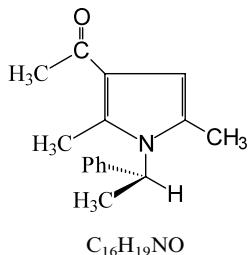
(*S*)-1-[1-(Hydroxymethylpropyl)-2,5-dimethyl-1*H*-pyrrol-3-yl]ethanone

E.e. = 98%

[ $\alpha$ ]<sub>D</sub><sup>20</sup> = -16.4 (c 0.7, CHCl<sub>3</sub>)

Source of chirality: (*S*)-2-amino-1-butanol as starting material

Absolute configuration: *S*



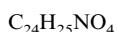
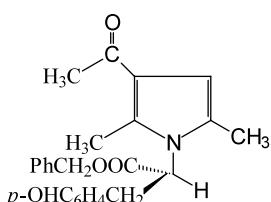
(*S*)-1-[2,5-Dimethyl-1-(1-phenylethyl)-1*H*-pyrrol-3-yl]ethanone

E.e.=99%

$[\alpha]_D^{20} = -12.8$  (*c* 0.6, CHCl<sub>3</sub>)

Source of chirality: (*S*)-1-phenylethylamine as starting material

Absolute configuration: *S*



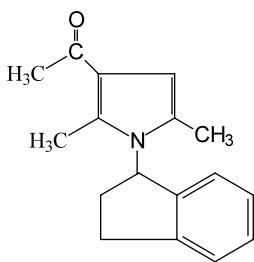
(*S*)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)-3-(4-hydroxyphenyl)propionic acid phenyl ester

E.e.=98%

$[\alpha]_D^{20} = -101.7$  (*c* 0.6, CHCl<sub>3</sub>)

Source of chirality: L-tyrosine benzyl ester *p*-toluenesulfonate salt as starting material

Absolute configuration: *S*



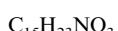
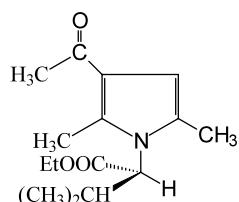
(*R*)-1-(1-Indan-1-yl-2,5-dimethyl-1*H*-pyrrol-3-yl)ethanone

E.e.=98%

$[\alpha]_D^{20} = -1.5$  (*c* 0.9, CHCl<sub>3</sub>)

Source of chirality: (*R*)-1-aminoindane as starting material

Absolute configuration: *R*



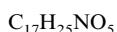
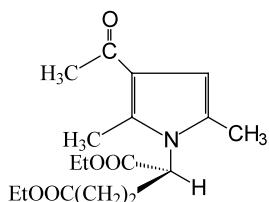
(*S*)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)-3-methylbutyric acid ethyl ester

E.e.=99%

$[\alpha]_D^{20} = -108.7$  (*c* 0.6, CHCl<sub>3</sub>)

Source of chirality: L-valine ethyl ester hydrochloride as starting material

Absolute configuration: *S*



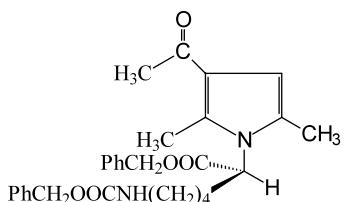
(S)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)pentanedioic acid diethyl ester

E.e.=96%

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=−60.2 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: L-glutamic acid diethyl ester hydrochloride as starting material

Absolute configuration: *S*



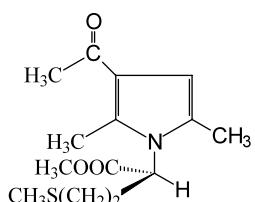
(S)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)-6-phenoxy carbonylaminohexanoic acid benzyl ester

E.e.=99%

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=−29.1 (*c* 0.9, CHCl<sub>3</sub>)

Source of chirality: N<sub>ε</sub>-CBZ-L-lysine benzyl ester hydrochloride as starting material

Absolute configuration: *S*



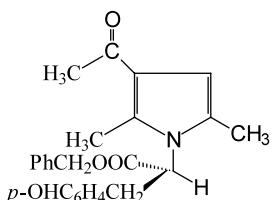
(S)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)-4-methylsulfanylbutyric acid methyl ester

E.e.=96%

[ $\alpha$ ]<sub>D</sub><sup>20</sup>=−77.7 (*c* 0.4, CHCl<sub>3</sub>)

Source of chirality: L-methionine methyl ester hydrochloride as starting material

Absolute configuration: *S*



(S)-2-(3-Acetyl-2,5-dimethylpyrrol-1-yl)-3-(4-hydroxyphenyl)propionic acid phenyl ester

E.e.=98%

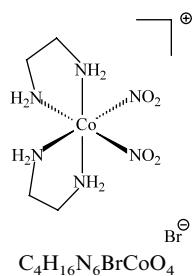
[ $\alpha$ ]<sub>D</sub><sup>20</sup>=−101.7 (*c* 0.6, CHCl<sub>3</sub>)

Source of chirality: L-tyrosine benzyl ester *p*-toluenesulfonate salt as starting material

Absolute configuration: *S*

Remir G. Kostyanovsky,\* Vladimir Yu. Torbeev and Konstantin A. Lyssenko

*Tetrahedron: Asymmetry* 12 (2001) 2721



$\Lambda(-)_{589}$ -*cis*-bis(ethylenediamine)dinitrocobaltbromide

E.e.=56%

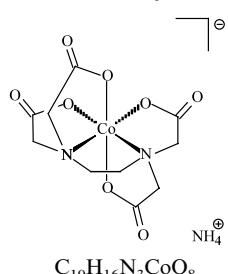
$[\alpha]_{589}^{23} = -24.6$  (*c* 1,  $\text{H}_2\text{O}$ )

Source of chirality: spontaneous resolution

Absolute configuration:  $\Lambda(-)_{589}$

Remir G. Kostyanovsky,\* Vladimir Yu. Torbeev and Konstantin A. Lyssenko

*Tetrahedron: Asymmetry* 12 (2001) 2721



Ammonium  $\Lambda(-)_{546}$ -ethylenediaminetetraacetatocobaltate

E.e.=83%

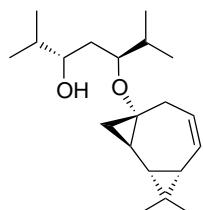
$[\alpha]_{546}^{25} = -1250.9$  (*c* 0.02,  $\text{H}_2\text{O}$ )

Source of chirality: spontaneous resolution

Absolute configuration:  $\Lambda(-)_{546}$

Takahiro Tei, Takashi Sugimura,\* Toshifumi Katagiri, Akira Tai and Tadashi Okuyama

*Tetrahedron: Asymmetry* 12 (2001) 2727



9,9-Dimethyl-4-(2,6-dimethyl-5-hydroxyheptyl-3-oxy)tricyclo[6.1.0.0<sup>2,4</sup>]non-6-ene

E.e.>99%

$[\alpha]_D^{20} = -29.3$  (*c* 1.2, methanol)

Source of chirality: (3*S*,5*S*)-2,6-dimethyl-3,5-heptane-diol

Absolute configuration: 1*S*,2*R*,4*R*,8*S*

Cristiana Fava, Roberta Galeazzi, Giovanna Mobbili and Mario Orena\*

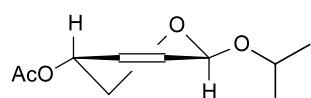
*Tetrahedron: Asymmetry* 12 (2001) 2731

Ee >99%

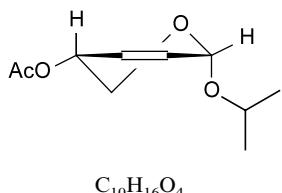
$[\alpha]_D +120.4$  (*c* 0.5,  $\text{CHCl}_3$ )

Source of chirality: D-xylose

Absolute configuration: 1*S*,4*S*

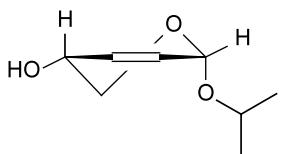


Isopropyl  $\beta$ -D-4-*O*-acetyl-2,3-dideoxypent-2-enoglycopyranoside



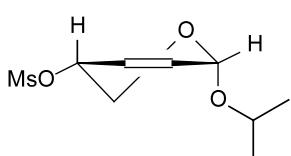
Isopropyl  $\alpha$ -D-4-O-acetyl-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +97.1$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*S*



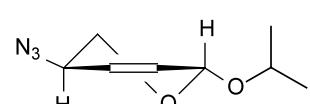
Isopropyl  $\alpha$ -D-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +37.9$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*S*



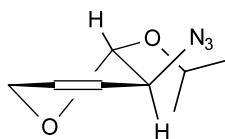
Isopropyl  $\alpha$ -D-4-O-methanesulphonyl-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +98.2$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*S*



Isopropyl  $\alpha$ -L-4-azido-2,3,4-trideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D -53.6$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*R*



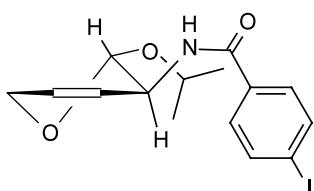
Isopropyl  $\alpha$ -2-azido-2,3,4-trideoxypent-3-enopyranoside

Ee >99%

$[\alpha]_D -71.4$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

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



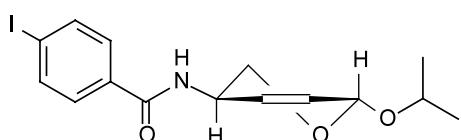
Isopropyl  $\alpha$ -2-(*p*-iodobenzamido)-2,3,4-trideoxypent-3-enopyranoside

Ee >99%

$[\alpha]_D -61.6$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

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



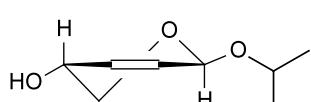
Isopropyl  $\alpha$ -L-4-(*p*-iodobenzamido)-2,3,4-trideoxypent-2-enoglyceropyranoside

Ee >99%

$[\alpha]_D -48.2$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1*R*,4*R*



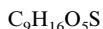
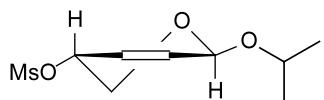
Isopropyl  $\beta$ -D-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%

$[\alpha]_D +101.4$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1*S*,4*S*



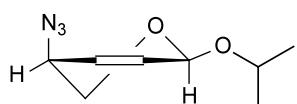
Isopropyl  $\beta$ -D-4-O-methanesulphonyl-2,3-dideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D +102.0$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1S,4S



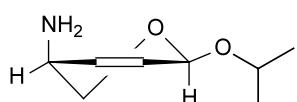
Isopropyl  $\beta$ -L-4-azido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D -132.2$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1S,4R



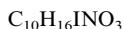
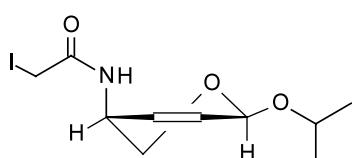
Isopropyl  $\beta$ -L-4-amino-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D -83.2$  (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1S,4R



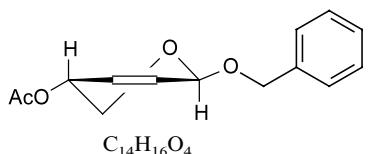
Isopropyl  $\beta$ -L-4-iodoacetamido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D -81.3$  (*c* 0.5, CHCl<sub>3</sub>)

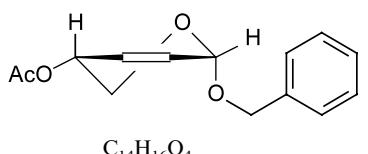
Source of chirality: D-xylose

Absolute configuration: 1S,4R



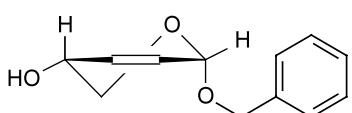
Benzyl  $\beta$ -D-4-O-acetyl-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +137.7$  ( $c$  0.5,  $\text{CHCl}_3$ )  
Source of chirality: D-xylose  
Absolute configuration: 1R,4S



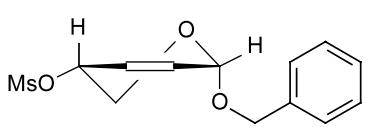
Benzyl  $\alpha$ -D-4-O-acetyl-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +77.1$  ( $c$  0.5,  $\text{CHCl}_3$ )  
Source of chirality: D-xylose  
Absolute configuration: 1S,4S



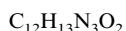
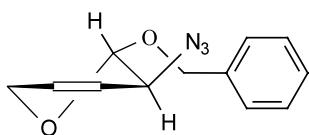
Benzyl  $\alpha$ -D-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +73.3$  ( $c$  0.5,  $\text{CHCl}_3$ )  
Source of chirality: D-xylose  
Absolute configuration: 1S,4S



Benzyl  $\alpha$ -D-4-O-methanesulphonyl-2,3-dideoxypent-2-enoglyceropyranoside

Ee >99%  
 $[\alpha]_D +84.2$  ( $c$  0.5,  $\text{CHCl}_3$ )  
Source of chirality: D-xylose  
Absolute configuration: 1S,4S



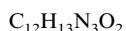
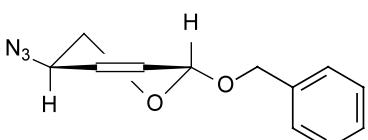
Benzyl  $\alpha$ -D-azido-2,3,4-trideoxypent-3-enopyranoside

Ee >99%

[ $\alpha$ ]<sub>D</sub> +68.8 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

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



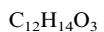
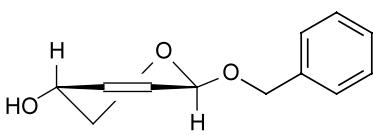
Benzyl  $\alpha$ -L-4-azido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

[ $\alpha$ ]<sub>D</sub> -66.8 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

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



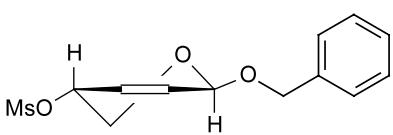
Benzyl  $\beta$ -D-2,3-dideoxypent-2-enoglycero pyranoside

Ee >99%

[ $\alpha$ ]<sub>D</sub> +99.2 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

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



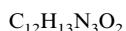
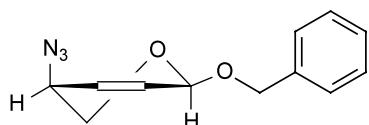
Benzyl  $\beta$ -D-4-O-methanesulphonyl-2,3-dideoxypent-2-enoglycero pyranoside

Ee >99%

[ $\alpha$ ]<sub>D</sub> +81.4 (*c* 0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

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



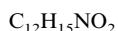
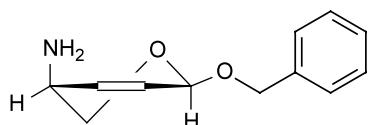
Benzyl  $\beta$ -L-4-azido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D -140.4$  ( $c$  0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1R,4R



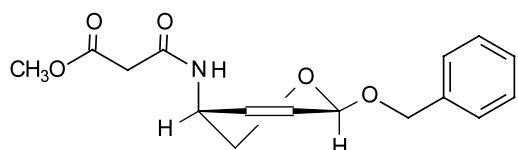
Benzyl  $\beta$ -L-4-amino-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D -63.6$  ( $c$  0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1R,4R



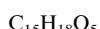
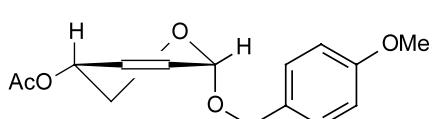
Benzyl  $\beta$ -L-4-methoxycarbonylacetamido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D -38.8$  ( $c$  0.5, CHCl<sub>3</sub>)

Source of chirality: D-xylose

Absolute configuration: 1R,4R



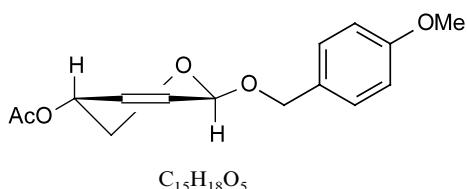
p-Methoxybenzyl  $\alpha$ -D-4-O-acetyl-2,3-dideoxypent-2-enoglycero pyranoside

Ee >99%

$[\alpha]_D +63.4$  ( $c$  0.5, CHCl<sub>3</sub>)

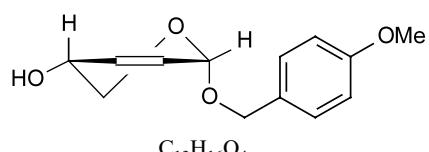
Source of chirality: D-xylose

Absolute configuration: 1S,4S



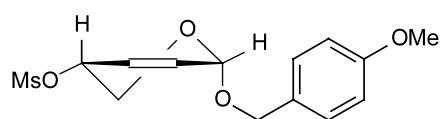
*p*-Methoxybenzyl  $\beta$ -D-4-O-acetyl-2,3-dideoxypent-2-enoglyceroxyranoside

Ee >99%  
 $[\alpha]_D +58.3$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*S*



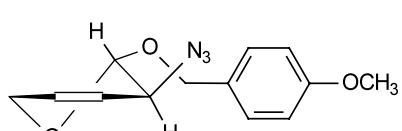
*p*-Methoxybenzyl  $\alpha$ -D-2,3-dideoxypent-2-enoglyceroxyranoside

Ee >99%  
 $[\alpha]_D +31.0$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*S*,4*S*



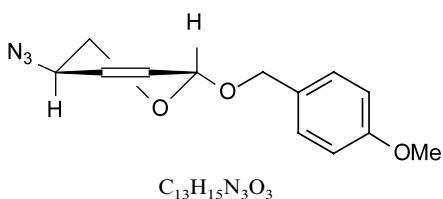
*p*-Methoxybenzyl  $\alpha$ -D-4-O-methanesulphonyl-2,3-dideoxypent-2-enoglyceroxyranoside

Ee >99%  
 $[\alpha]_D +61.2$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*S*,4*S*



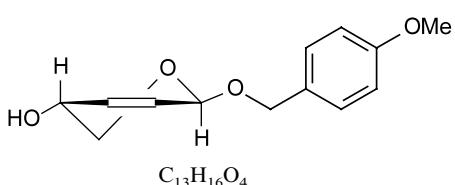
*p*-Methoxybenzyl  $\alpha$ -2-azido-2,3,4-trideoxypent-3-enopyranoside

Ee >99%  
 $[\alpha]_D +181.3$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,2*S*



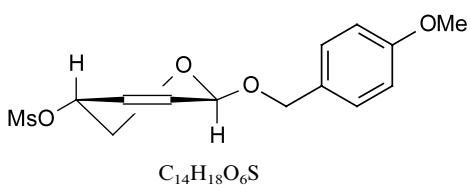
*p*-Methoxybenzyl  $\alpha$ -L-4-azido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%  
 $[\alpha]_D -136.1$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*S*,4*R*



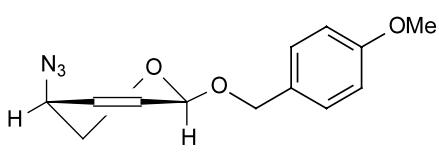
*p*-Methoxybenzyl  $\beta$ -D-2,3-dideoxypent-2-enoglycero pyranoside

Ee >99%  
 $[\alpha]_D +68.4$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*S*



*p*-Methoxybenzyl  $\beta$ -D-4-O-methanesulphonyl-2,3-dideoxypent-2-enoglycero pyranoside

Ee >99%  
 $[\alpha]_D +77.6$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*S*

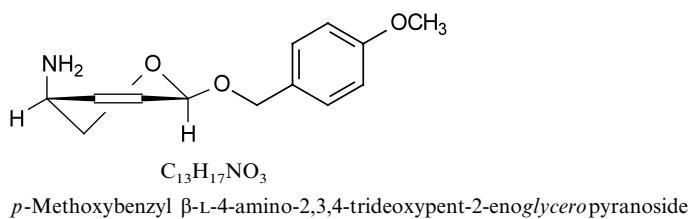


*p*-Methoxybenzyl  $\beta$ -L-4-azido-2,3,4-trideoxypent-2-enoglycero pyranoside

Ee >99%  
 $[\alpha]_D -137.5$  (*c* 0.5, CHCl<sub>3</sub>)  
Source of chirality: D-xylose  
Absolute configuration: 1*R*,4*R*

Cristiana Fava, Roberta Galeazzi, Giovanna Mobbili  
and Mario Orena\*

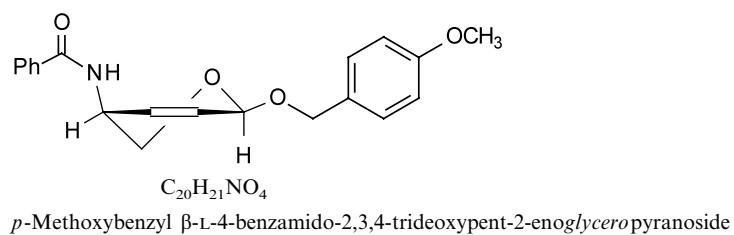
*Tetrahedron: Asymmetry* 12 (2001) 2731



Ee >99%  
 $[\alpha]_D^{25} +77.4$  (*c* 0.5,  $\text{CHCl}_3$ )  
 Source of chirality: D-xylose  
 Absolute configuration: 1*R*,4*R*

Cristiana Fava, Roberta Galeazzi, Giovanna Mobbili  
and Mario Orena\*

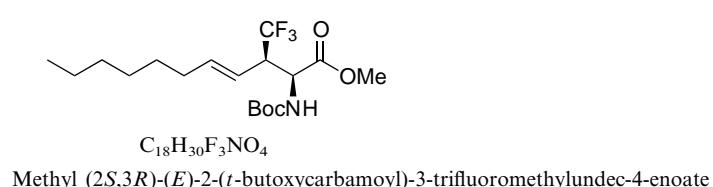
*Tetrahedron: Asymmetry* 12 (2001) 2731



Ee >99%  
 $[\alpha]_D^{25} -80.0$  (*c* 0.5,  $\text{CHCl}_3$ )  
 Source of chirality: D-xylose  
 Absolute configuration: 1*R*,4*R*

Tsutomu Konno,\* Takeshi Daitoh, Takashi Ishihara  
and Hiroki Yamanaka

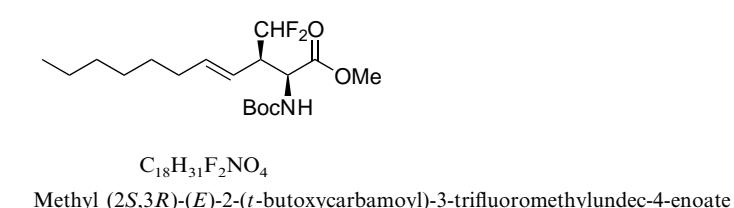
*Tetrahedron: Asymmetry* 12 (2001) 2743



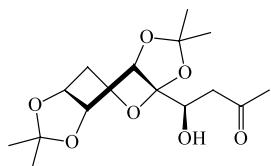
E.e. = 92%  
 $[\alpha]_D^{25} = +12.5$  (*c* 0.5,  $\text{CHCl}_3$ )  
 Source of chirality: asymmetric synthesis  
 Absolute configuration: 2*S*,3*R*

Tsutomu Konno,\* Takeshi Daitoh, Takashi Ishihara  
and Hiroki Yamanaka

*Tetrahedron: Asymmetry* 12 (2001) 2743



E.e. = 84%  
 $[\alpha]_D^{25} = +5.3$  (*c* 1.1,  $\text{CHCl}_3$ )  
 Source of chirality: asymmetric synthesis  
 Absolute configuration: 2*S*,3*R*



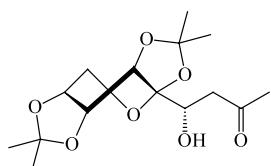
C<sub>15</sub>H<sub>24</sub>O<sub>7</sub>

1,3-Dideoxy-5,6:7,8-di-O-isopropylidene-β-D-manno-non-2,5-diulo-5,9-pyranose

[α]<sub>D</sub> +1.7, [α]<sub>405</sub> +12 (c 1.2, chloroform)

Source of chirality: D-fructose and stereoselective synthesis

Absolute configuration: 4R,5S,6S,7R,8R (assigned by chemical correlation)



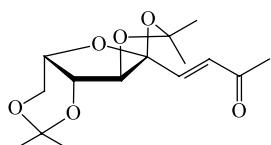
C<sub>15</sub>H<sub>24</sub>O<sub>7</sub>

1,3-Dideoxy-5,6:7,8-di-O-isopropylidene-β-D-gluco-non-2,5-diulo-5,9-pyranose

[α]<sub>D</sub> -18, [α]<sub>405</sub> -32 (c 1.7, chloroform)

Source of chirality: D-fructose and stereoselective synthesis

Absolute configuration: 4S,5S,6S,7R,8R (assigned by chemical correlation)



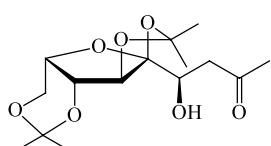
C<sub>15</sub>H<sub>22</sub>O<sub>6</sub>

(E)-1,3,4-Trideoxy-5,6:7,9-di-O-isopropylidene-α-L-xylo-non-3-ene-2,5-diulo-5,8-furanose

[α]<sub>D</sub><sup>26</sup> +24 (c 1, chloroform)

Source of chirality: L-sorbose and stereoselective synthesis

Absolute configuration: 3E,5S,6S,7S,8S (assigned by spectroscopic data)



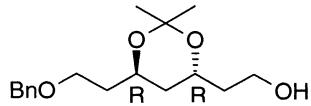
C<sub>15</sub>H<sub>24</sub>O<sub>7</sub>

1,3-Dideoxy-5,6:7,9-di-O-isopropylidene-α-L-gulo-non-2,5-diulo-5,8-furanose

[α]<sub>D</sub><sup>27</sup> +15 (c 1.1, chloroform)

Source of chirality: L-sorbose and stereoselective synthesis

Absolute configuration: 4R,5S,6S,7S,8S (assigned by chemical correlation)

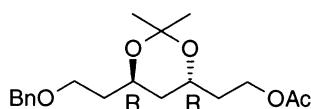


$C_{17}H_{28}O_4$   
(-)-(3*R*,5*R*)-7-Benzylxyloxy-3,5-*O*-isopropylidene-heptane-1,3,5-triol

E.e.=98%

 $[\alpha]_D = -9.1$  ( $c = 1.0$ , CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

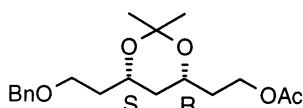
Absolute configuration: 3*R*,5*R*

$C_{19}H_{30}O_5$   
(+)-(3*R*,5*R*)-1-Acetoxy-7-benzylxyloxy-3,5-dihydroxy-3,5-*O*-isopropylidene-heptane

E.e.=98%

 $[\alpha]_D = +1.5$  ( $c = 0.9$ , CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 3*R*,5*R*

$C_{19}H_{30}O_5$   
(-)-(3*R*,5*S*)-1-Acetoxy-7-benzylxyloxy-3,5-dihydroxy-3,5-*O*-isopropylidene-heptane

E.e.=98%

 $[\alpha]_D = -1.3$  ( $c = 1.1$ , CHCl<sub>3</sub>)

Source of chirality: enzymatic resolution

Absolute configuration: 3*R*,5*S*