

**STEREOCHEMISTRY ABSTRACTS**

N. M. Allanson, A. H. Davidson, C. D. Floyd, F. M. Martin

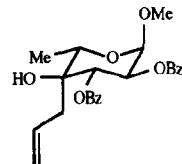
*Tetrahedron: Asymmetry* 1994, 5, 2061

$[\alpha]_D^{24} = -175.2$  ( $c = 0.22$ , DMF)

Source of chirality: natural and stereoselective synthesis

$C_{24}H_{26}O_7$

Methyl 2,3-di-*O*-benzoyl-4-(prop-2'-en-1'-yl)-6-deoxy- $\beta$ -L-glucopyranoside



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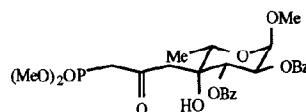
*Tetrahedron: Asymmetry* 1994, 5, 2061

$[\alpha]_D^{24} = -119.0$  ( $c = 0.2$ , DMF)

Source of chirality: natural and stereoselective synthesis

$C_{26}H_{31}O_{11}P$

Methyl 2,3-di-*O*-benzoyl-4-(3'-dimethylphosphono-2'-oxo-propan-1'-yl)-6-deoxy- $\beta$ -L-galactopyranoside



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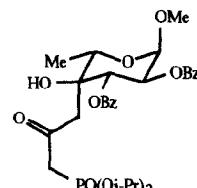
*Tetrahedron: Asymmetry* 1994, 5, 2061

$[\alpha]_D^{24} = -100.8.$  ( $c = 0.4$ , DMF)

Source of chirality: natural and stereoselective synthesis

$C_{30}H_{39}O_{11}P$

Methyl 2,3-di-*O*-benzoyl-4-(3'-diisopropylphosphono-2'-oxo-propan-1'-yl)-6-deoxy- $\beta$ -L-glucopyranoside



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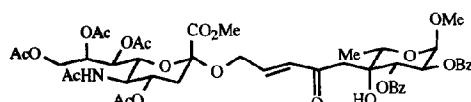
*Tetrahedron: Asymmetry* 1994, 5, 2061

$[\alpha]_D^{24} = -64.7$  ( $c = 0.21$ , DMF)

Source of chirality: natural and stereoselective synthesis

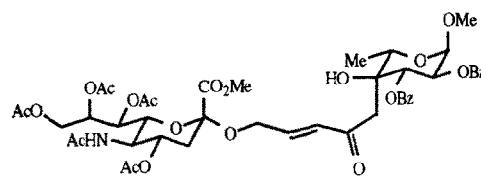
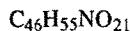
$C_{46}H_{55}NO_{21}$

Methyl 5-acetamido-2-[5''-(methyl 2',3'-di-*O*-benzoyl-6'-deoxy- $\beta$ -L-galactopyranoside-4'-yl)-4''-oxo-pent-2''-en-1-yl]-4,7,8,9-tetra-*O*-acetyl-3,5-dideoxy-D-glycero- $\alpha$ -D-galacto-2-nonulosonate

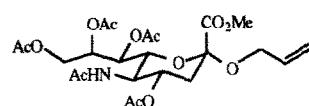


$[\alpha]_D^{24} = -53.3$  ( $c = 0.15$ , DMF)

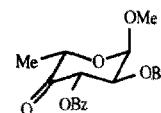
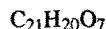
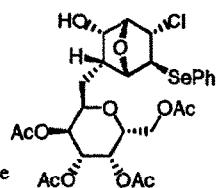
Source of chirality: natural

Methyl 5-acetamido-2-[5''-(methyl 2',3'-di-O-benzoyl-6'-deoxy- $\beta$ -L-glucopyranoside-4'-yl)-4'-oxo-pent-2'-en-1-yl]-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-D-glycero- $\alpha$ -D-galacto-2-nonulosonate $[\alpha]_D^{24} = +9.7$  ( $c = 0.25$ , DMF)

Source of chirality: natural

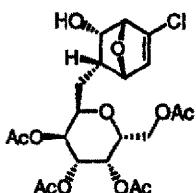
Methyl 2-allyl-5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-D-glycero- $\alpha$ -D-galacto-2-nonulosonate $[\alpha]_D^{24} = -113.1$ . ( $c = 2.8$ , DMF)

Source of chirality: natural

Methyl 2,3-di-O-benzoyl-4-oxo- $\beta$ -L-fucopyranoside

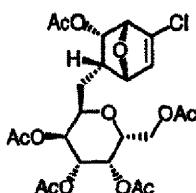
D.e. >98% (by 400 MHz  $^1$ H-NMR)  
 $[\alpha]_D^{27} = +41.2$  ( $c = 0.92$ ,  $CHCl_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1R,2S,3R,4S,5S,6S),  $\alpha$ -D-Gal  
 (by synthesis)

(+)-(1R,2S,3R,4S,5S,6S)-5-exo-Benzeneseleneny1-6-endo-chloro-3-endo-[(2',3',4',6'-tetra-O-acetyl-alpha-D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]heptan-2-endo-ol



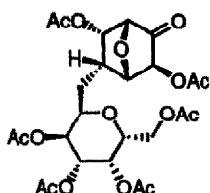
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{27} = +61$  ( $c = 1.3, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1R,2S,3R,4S),  $\alpha$ -D-Gal  
 (by synthesis)

(+)-(1R,2S,3R,4S)-6-Chloro-3-*endo*-[(2',3',4',6'-tetra-O-acetyl- $\alpha$ -D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]hept-5-en-2-*endo*-ol



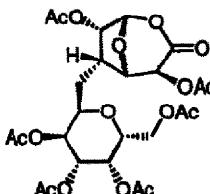
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{24} = +47$  ( $c = 1.07, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1R,2S,3R,4S),  $\alpha$ -D-Gal  
 (by synthesis)

(+)-(1R,2S,3R,4S)-6-Chloro-3-*endo*-[(2',3',4',6'-tetra-O-acetyl- $\alpha$ -D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]hept-5-en-2-*endo*-yl Acetate



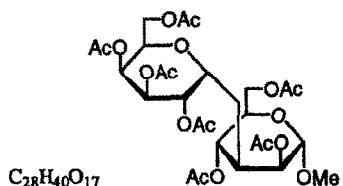
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{24} = +28.3$  ( $c = 1.1, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1R,2S,3S,4S,5S),  $\alpha$ -D-Gal  
 (by synthesis)

(+)-(1R,2S,3S,4S,5S)-6-oxo-3-*endo*-[(2',3',4',6'-tetra-O-acetyl- $\alpha$ -D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]hept-5-en-2-*endo*, 5-*exo*-diyl Diacetate



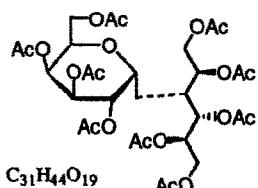
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{24} = -20.3$  ( $c = 1.4, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1S,4S,5S,6S,7S),  $\alpha$ -D-Gal  
 (by synthesis)

(-)-(1S,4S,5S,6S,7S)-3-oxo-6-*endo*-[(2',3',4',6'-tetra-O-acetyl- $\alpha$ -D-galactopyranosyl)methyl]-2,8-dioxabicyclo[3.2.1]octa-4-*exo*, 7-*endo*-diyl Diacetate



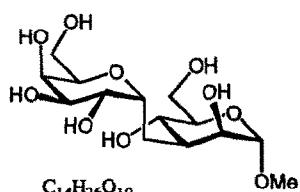
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{23} = +62$  ( $c = 1.35, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration:  $\alpha$ -D-Gal,  $\alpha$ -D-Man  
 (by synthesis)

(+)-Methyl 3-Deoxy-3-[(2',3',4',6'-tetra-O-acetyl- $\alpha$ -D-galactopyranosyl)methyl]-2,4,6-tri-O-acetyl- $\alpha$ -D-mannopyranoside



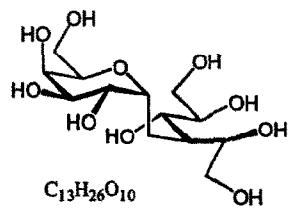
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{24} = +66.5$  ( $c = 1.0, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration:  $\alpha$ -D-Gal, D-Man  
 (by synthesis)

(+)-3-Deoxy-3-[(2',3',4',6'-tetra-O-acetyl- $\alpha$ -D-galactopyranosyl)methyl]-1,2,4,5,6-penta-O-acetyl-D-mannitol



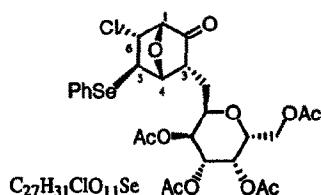
D.e. >98% (by 600 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{23} = +49.6$  ( $c = 0.75, \text{MeOH}$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration  $\alpha$ -D-Galp(1 $\rightarrow$ 3)-CH<sub>2</sub>- $\alpha$ -D-Manp-OMe  
 (assigned by synthesis using D-Gal and (-)-(1S,4S)-7-oxabicyclo[2.2.1]hept-5-en-2-one)

(+)-Methyl 3-Deoxy-3-[( $\alpha$ -D-galactopyranosyl)methyl]- $\alpha$ -D-mannopyranoside



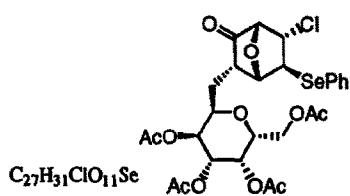
D.e. >98% (by  $^1\text{H-NMR}$  of synthetic precursor)  
 $[\alpha]_D^{23} = +33$  ( $c = 0.9, \text{MeOH}$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: by synthesis

(+)-3-Deoxy-3-[( $\alpha$ -D-galactopyranosyl)methyl]-D-mannitol



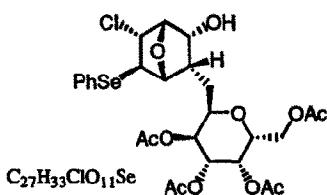
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ ,  $^{13}\text{CH}$  satellites)  
 $[\alpha]_D^{26} = +24$  ( $c = 1.0, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1S,3S,4R,5R,6R),  $\alpha$ -D-Gal  
 (by synthesis starting with (+)-(1R,4R)-7-oxabicyclo[2.2.1]hept-5-en-2-one and D-Gal)

(+)-(1S,3S,4R,5R,6R)-5-exo-Benzeneselenenyl-6-endo-chloro-3-endo-[(2',3',4',6'-tetra-O-acetyl-α-D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]heptan-2-one



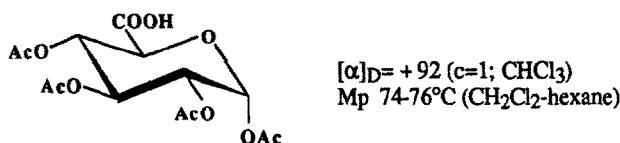
D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{26} = +41$  ( $c = 1.0, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1R,3R,4S,5S,6S),  $\alpha$ -D-Gal  
 (by synthesis)

(+)-(1R,3R,4S,5S,6S)-5-exo-Benzeneselenenyl-6-endo-chloro-3-endo-[(2',3',4',6'-tetra-O-acetyl-α-D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]heptan-2-one

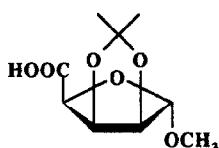


D.e. >98% (by 400 MHz  $^1\text{H-NMR}$ )  
 $[\alpha]_D^{27} = +37.7$  ( $c = 1.2, \text{CHCl}_3$ )  
 Source of chirality: natural (D-Gal)  
 Absolute configuration: (1S,2R,3S,4R,5R,6R),  $\alpha$ -D-Gal  
 (by synthesis)

(+)-(1S,2R,3S,4R,5R,6R)-5-exo-Benzeneselenenyl-6-endo-chloro-3-endo-[(2',3',4',6'-tetra-O-acetyl-α-D-galactopyranosyl)methyl]-7-oxabicyclo[2.2.1]heptan-2-endo-ol

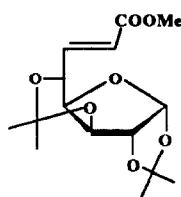


1,2:3,4-Tetra-O-acetyl- $\alpha$ -D-glucopyranuronic acid



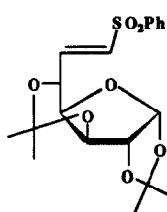
$[\alpha]_D = +19.9$  ( $c = 2.47$ ;  $\text{CHCl}_3$ )

$\text{C}_{29}\text{H}_{14}\text{O}_6$   
Methyl-2,3-O-isopropylidene- $\alpha$ -D-lyxo-furanosiduronic acid



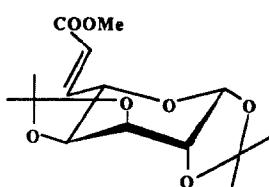
$[\alpha]_D = +27$  ( $c = 1.44$ ;  $\text{CHCl}_3$ )

$\text{C}_{15}\text{H}_{22}\text{O}_7$   
Methyl 6,7-dideoxy-1,2:3,5-di-O-isopropylidene- $\alpha$ -D-gluco-6-ene-octofuranuronate



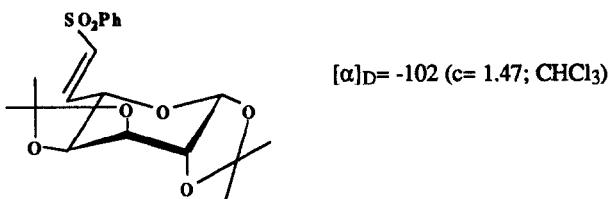
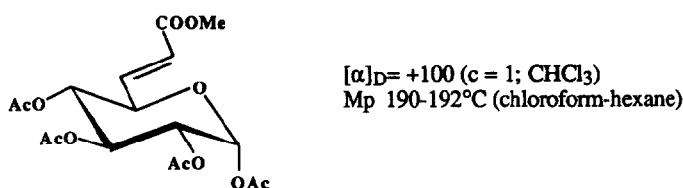
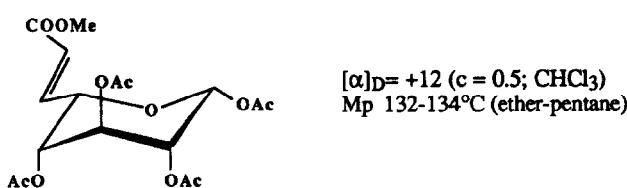
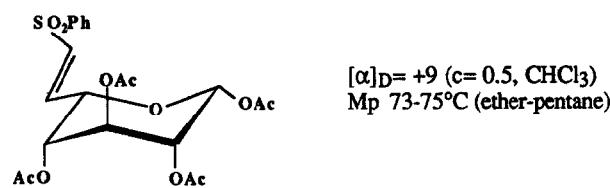
$[\alpha]_D = +42$  ( $c = 0.5$ ;  $\text{CHCl}_3$ ).  
Mp 145–146°C (ether-pentane)

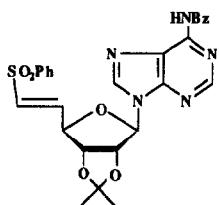
$\text{C}_{29}\text{H}_{24}\text{O}_7\text{S}$   
6,7-Dideoxy-1,2:3,5-di-O-isopropylidene-7-phenylsulphonyl- $\alpha$ -D-gluco-6-ene-heptofuranose



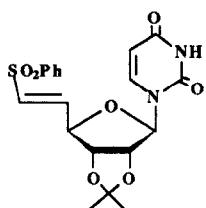
$[\alpha]_D = -93$  ( $c = 2.4$ ;  $\text{CHCl}_3$ )

$\text{C}_{15}\text{H}_{22}\text{O}_7$   
Methyl 6,7-dideoxy-1,2:3,4-di-O-isopropylidene-6-ene- $\alpha$ -L-galacto-octopyranuronate

 $\text{C}_{19}\text{H}_{24}\text{O}_7\text{S}$ 6,7-Dideoxy-1,2:3,4-di-*O*-isopropylidene-7-phenylsulphonyl- $\alpha$ -L-galacto-6-ene-heptopyranose $\text{C}_{17}\text{H}_{22}\text{O}_{11}$ Methyl 6,7-dideoxy-1,2:3,4-tetra-*O*-acetyl-7- $\alpha$ -D-gluco-6-ene-octopyranuronate $\text{C}_{17}\text{H}_{22}\text{O}_{11}$ Methyl 6,7-dideoxy-1,2:3,4-tetra-*O*-acetyl-7- $\alpha$ -L-ido-6-ene-octopyranuronate $\text{C}_{21}\text{H}_{24}\text{NO}_{11}\text{S}$ 6,7-Dideoxy-1,2:3,4-tetra-*O*-acetyl-7-phenylsulphonyl-7- $\alpha$ -L-ido-6-ene-heptopyranose



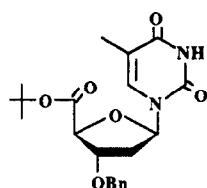
$[\alpha]_D = +94$  ( $c = 0.5$ ; CHCl<sub>3</sub>)  
Mp 115-118°C (CH<sub>2</sub>Cl<sub>2</sub>)

C<sub>27</sub>H<sub>25</sub>N<sub>5</sub>O<sub>6</sub>S(5',6'-Dideoxy-2',3'-O-isopropylidene-6'-phenylsulphonyl-β-D-ribo-5'-ene-hexofuranosyl)-9-N<sup>6</sup>-Benzoyl-adenine

$[\alpha]_D = +64$  ( $c = 0.5$ ; CHCl<sub>3</sub>)  
Mp 111-114°C (CH<sub>2</sub>Cl<sub>2</sub>-hexane)

C<sub>19</sub>H<sub>20</sub>N<sub>2</sub>O<sub>7</sub>S

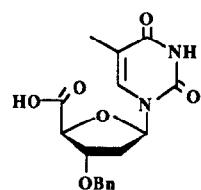
(5',6'-Dideoxy-2',3'-O-isopropylidene-6'-phenylsulphonyl-β-D-ribo-6'-ene-hexofuranosyl)-1-Uracil



$[\alpha]_D = +38$  ( $c = 1$ ; CHCl<sub>3</sub>)  
Mp 156-158°C (CH<sub>2</sub>Cl<sub>2</sub>-hexane)

C<sub>21</sub>H<sub>26</sub>N<sub>2</sub>O<sub>6</sub>

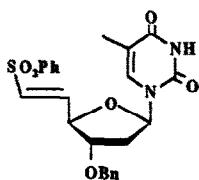
(tert-Butyl-3'-O-benzyl-3'-deoxy-β-D-ribo-furanuronate)-1-thymine



$[\alpha]_D = +28$  ( $c = 1$ ; DMF)  
Mp 238-240°C (methanol)

C<sub>17</sub>H<sub>18</sub>N<sub>2</sub>O<sub>6</sub>

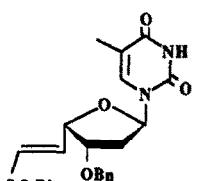
(3'-O-Benzyl-3'-deoxy-β-D-ribo-furanuronic acid)-1-thymine



$[\alpha]_D = +81$  ( $c = 0.5$ ;  $\text{CHCl}_3$ )  
Mp 161-163°C ( $\text{CH}_2\text{Cl}_2$ -ether)

 $\text{C}_{24}\text{H}_{24}\text{N}_2\text{O}_6\text{S}$ 

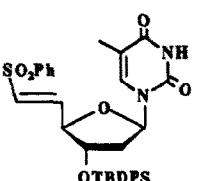
(3'-O-Benzyl-6'-phenylsulphonyl-2',5',6'-trideoxy-beta-D-ribo-5'-ene-hexofuranosyl)-1-thymine



$[\alpha]_D = +15$  ( $c = 0.5$ ; DMF)  
Mp 84-86°C ( $\text{CH}_2\text{Cl}_2$ -pentane)

 $\text{C}_{24}\text{H}_{24}\text{N}_2\text{O}_6\text{S}$ 

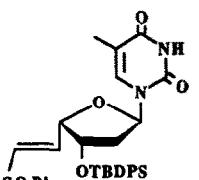
(3'-O-Benzyl-6'-phenylsulphonyl-2',5',6'-trideoxy-beta-L-ribo-5'-ene-hexofuranosyl)-1-thymine



$[\alpha]_D = +44$  ( $c = 0.5$ ;  $\text{CHCl}_3$ )  
Mp 92-96°C (ether-pentane)

 $\text{C}_{33}\text{H}_{36}\text{N}_2\text{O}_6\text{S Si}$ 

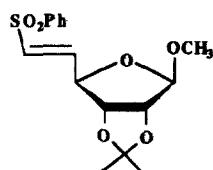
(3'-O-tert-Butyldiphenylsilyl-6'-phenylsulphonyl-2',5',6'-trideoxy-beta-D-ribo-5'-ene-hexofuranosyl)-1-thymine



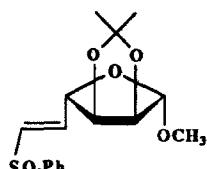
$[\alpha]_D = +17$  ( $c = 1$ ;  $\text{CHCl}_3$ )  
Mp 118-120°C ( $\text{CH}_2\text{Cl}_2$ -hexane)

 $\text{C}_{33}\text{H}_{36}\text{N}_2\text{O}_6\text{S Si}$ 

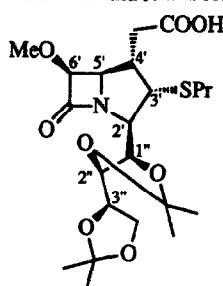
(3'-O-tert-Butyldiphenylsilyl-6'-phenylsulphonyl-2',5',6'-trideoxy-beta-L-ribo-5'-ene-hexofuranosyl)-1-thymine



$[\alpha]_D = +2.4$  ( $c = 0.5$ ;  $\text{CHCl}_3$ )  
Mp 118-120°C (ether-pentane)

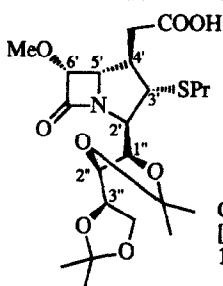
 $\text{C}_{16}\text{H}_{20}\text{O}_6\text{S}$ .Methyl 5,6-dideoxy-2,3-isopropylidene-6-phenylsulphonyl- $\beta$ -D-ribo-5-ene-hexo-furanoside

$[\alpha]_D = -2.4$  ( $c = 0.5$ ;  $\text{CHCl}_3$ )  
Mp 118-120°C (ether-pentane)

 $\text{C}_{16}\text{H}_{20}\text{O}_6\text{S}$ .Methyl 5,6-dideoxy-2,3-isopropylidene-6-phenylsulphonyl- $\alpha$ -L-lyxo-5-ene-hexo-furanoside

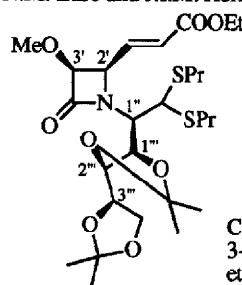
$[\alpha]_D = -$  ( $\text{CHCl}_3$ )  
Source of chirality: D-Glucosamine.  
Absolute configuration: 2'R, 3'S, 4'S, 5'R, 6'S, 1''R, 2''S, 3''R

$\text{C}_{22}\text{H}_{35}\text{NO}_8\text{S}$   
[6-Methoxy-7-oxo-3-propylsulfanyl-2-(1,2;3,4-di-O-isopropylidene-1,2,3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]hept-4-yl] acetic acid.



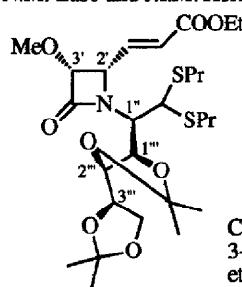
$[\alpha]_D = +$  ( $\text{CHCl}_3$ )  
Source of chirality: D-Glucosamine.  
Absolute configuration: 2'R, 3'S, 4'R, 5'S, 6'R, 1''R, 2''S, 3''R

$\text{C}_{22}\text{H}_{35}\text{NO}_8\text{S}$   
[6-Methoxy-7-oxo-3-propylsulfanyl-2-(1,2;3,4-di-O-isopropylidene-1,2,3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]hept-4-yl] acetic acid.



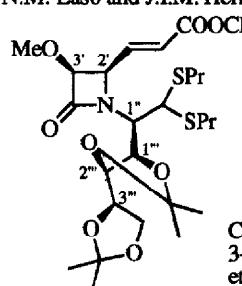
$[\alpha]_D = -94$  (c 1, CHCl<sub>3</sub>)  
Source of chirality: *D*-Glucosamine.  
Absolute configuration: 2'R, 3'S, 1"R, 1'''R, 2'''S, 3'''R

C<sub>27</sub>H<sub>45</sub>NO<sub>8</sub>S<sub>2</sub>  
3-{[2,2-Bis-propylsulfanyl]-1-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-ethyl}-3-methoxy-4-oxo-azetidin-2-yl acrylic acid ethyl ester.



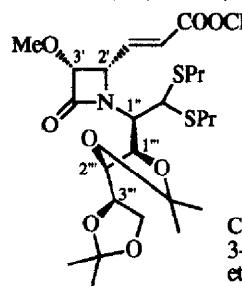
$[\alpha]_D = +17$  (c 1, CHCl<sub>3</sub>)  
Source of chirality: *D*-Glucosamine.  
Absolute configuration: 2'S, 3'R, 1"R, 1'''R, 2'''S, 3'''R

C<sub>27</sub>H<sub>45</sub>NO<sub>8</sub>S<sub>2</sub>  
3-{[2,2-Bis-propylsulfanyl]-1-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-ethyl}-3-methoxy-4-oxo-azetidin-2-yl acrylic acid ethyl ester.



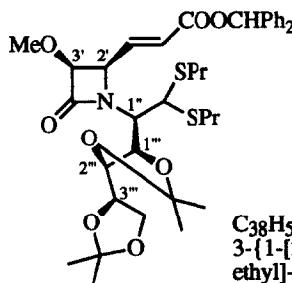
$[\alpha]_D = -102$  (c 1, CHCl<sub>3</sub>)  
Source of chirality: *D*-Glucosamine.  
Absolute configuration: 2'R, 3'S, 1"R, 1'''R, 2'''S, 3'''R

C<sub>32</sub>H<sub>47</sub>NO<sub>8</sub>S<sub>2</sub>  
3-{[2,2-Bis-propylsulfanyl]-1-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-ethyl}-3-methoxy-4-oxo-azetidin-2-yl acrylic acid benzyl ester.



$[\alpha]_D = +15$  (c 1, CHCl<sub>3</sub>)  
Source of chirality: *D*-Glucosamine.  
Absolute configuration: 2'S, 3'R, 1"R, 1'''R, 2'''S, 3'''R

C<sub>32</sub>H<sub>47</sub>NO<sub>8</sub>S<sub>2</sub>  
3-{[2,2-Bis-propylsulfanyl]-1-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-ethyl}-3-methoxy-4-oxo-azetidin-2-yl acrylic acid benzyl ester.

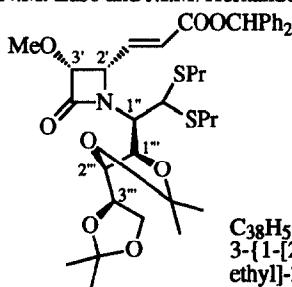


$[\alpha]_D = -83$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2'*R*, 3'*S*, 1''*R*, 1'''*R*, 2''*S*, 3'''*R*.

C<sub>38</sub>H<sub>51</sub>NO<sub>8</sub>S<sub>2</sub>  
3-{1-[2,2-Bis-propylsulfanyl-1-(1,2;3,4-di-*O*-isopropylidene-1,2;3,4-tetrahydroxybutyl)-ethyl]-3-methoxy-4-oxo-azetidin-2-yl}acrylic acid benzhydryl ester.

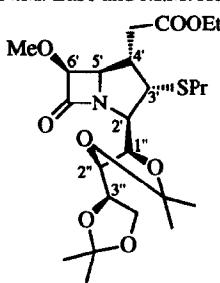


$[\alpha]_D = +28$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2'S, 3'R, 1''R, 1'''R, 2''S, 3'''R.

C<sub>38</sub>H<sub>51</sub>NO<sub>8</sub>S<sub>2</sub>  
3-{1-[2,2-Bis-propylsulfanyl-1-(1,2;3,4-di-*O*-isopropylidene-1,2;3,4-tetrahydroxybutyl)-ethyl]-3-methoxy-4-oxo-azetidin-2-yl}acrylic acid benzhydryl ester.

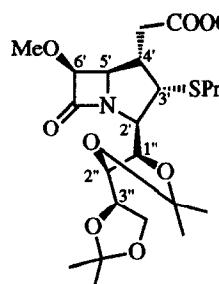


$[\alpha]_D = -16$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2'R, 3'S, 4'S, 5'R, 6'S, 1''R, 2''S, 3'''R.

C<sub>24</sub>H<sub>39</sub>NO<sub>8</sub>S  
[6-Methoxy-7-oxo-3-propylsulfanyl-2-(1,2;3,4-di-*O*-isopropylidene-1,2;3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]hept-4-yl] acetic acid ethyl ester.

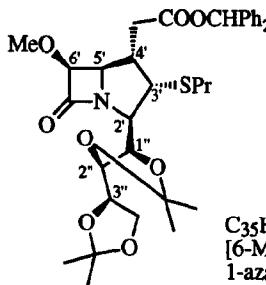


$[\alpha]_D = -8$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2'R, 3'S, 4'S, 5'R, 6'S, 1''R, 2''S, 3'''R.

C<sub>29</sub>H<sub>41</sub>NO<sub>8</sub>S  
[6-Methoxy-7-oxo-3-propylsulfanyl-2-(1,2;3,4-di-*O*-isopropylidene-1,2;3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]hept-4-yl] acetic acid benzyl ester.



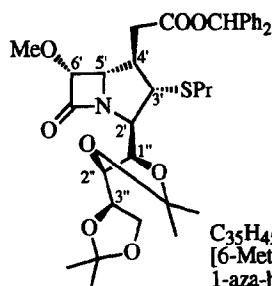
$[\alpha]_D = -11$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2'R, 3'S, 4'S, 5'R, 6'S, 1''R, 2''S, 3''R

C<sub>35</sub>H<sub>45</sub>NO<sub>8</sub>S

[6-Methoxy-7-oxo-3-propylsulfanyl-2-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]hept-4-yl] acetic acid benzhydryl ester.



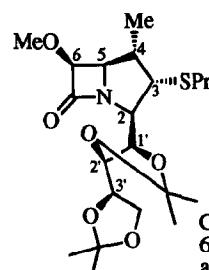
$[\alpha]_D = +30$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2'R, 3'S, 4'R, 5'S, 6'R, 1''R, 2''S, 3''R

C<sub>35</sub>H<sub>45</sub>NO<sub>8</sub>S

[6-Methoxy-7-oxo-3-propylsulfanyl-2-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]hept-4-yl] acetic acid benzhydryl ester.



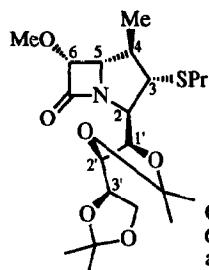
$[\alpha]_D = -35$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2R, 3S, 4S, 5R, 6S, 1'R, 2'S, 3'R

C<sub>21</sub>H<sub>35</sub>NO<sub>6</sub>S

6-Methoxy-4-methyl-3-propylsulfanyl-2-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]heptan-7-one.



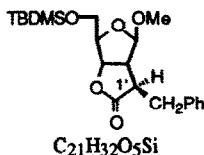
$[\alpha]_D = +41$  (c 1, CHCl<sub>3</sub>)

Source of chirality: *D*-Glucosamine.

Absolute configuration: 2R, 3S, 4R, 5S, 6R, 1'R, 2'S, 3'R

C<sub>21</sub>H<sub>35</sub>NO<sub>6</sub>S

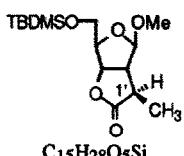
6-Methoxy-4-methyl-3-propylsulfanyl-2-(1,2;3,4-di-O-isopropylidene-1,2;3,4-tetrahydroxybutyl)-1-aza-bicyclo[3.2.0]heptan-7-one.


 $[\alpha]_D^{25} = -2.0 \text{ (c } 1, \text{ CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 5-O-(*t*-butylidemethylsilyl)-3-O-cinnamoyl-2-O-(imidazol-1-yl)thiocarbonyl- $\beta$ -D-ribofuranoside

**Absolute configuration:** 1' R

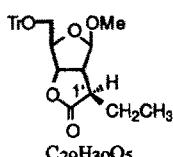
Methyl 5-O-(*t*-butylidemethylsilyl)-2-C-[{(R)carboxybenzylmethyl}-]2-deoxy-3,2-gamma-lactone-beta-D-ribofuranoside


 $[\alpha]_D^{25} = -30.0 \text{ (c } 1, \text{ CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 3-O-acryloyl-5-O-(*t*-butylidemethylsilyl)-2-O-(imidazol-1-yl)thiocarbonyl- $\beta$ -D-ribofuranoside

**Absolute configuration:** 1' R

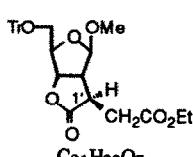
Methyl 5-O-(*t*-butylidemethylsilyl)-2-C-[{(R)carboxymethylmethyl}-]2-deoxy-3,2-gamma-lactone-beta-D-ribofuranoside


 $[\alpha]_D^{25} = -11.8 \text{ (c } 1, \text{ CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 3-O-crotonyl-2-O-(imidazol-1-yl)thiocarbonyl-5-O-trityl- $\beta$ -D-ribofuranoside

Methyl 2-C-[(R)carboxyethylmethyl]-2-deoxy-5-O-trityl-3,2-gamma-lactone-beta-D-ribofuranoside

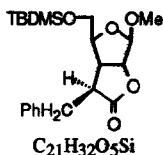
**Absolute configuration:** 1' R


 $[\alpha]_D^{25} = -1.4 \text{ (c } 1, \text{ CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 3-O-ethylfumaroyl-2-O-(imidazol-1-yl)thiocarbonyl-5-O-trityl- $\beta$ -D-ribofuranoside

**Absolute configuration:** 1' R

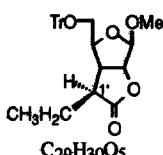
Methyl 2-C-[(R)carboxy(ethoxycarbonylmethyl)-]2-deoxy-5-O-trityl-3,2-gamma-lactone-beta-D-ribofuranoside


 $[\alpha]_D^{25} = -85.2 \text{ (c 1, CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 5-O-(*t*-butyldimethylsilyl)-2-O-cinnamoyl-3-O-(imidazol-1-yl)thiocarbonyl- $\beta$ -D-ribofuranoside

Methyl 5-O-(*t*-butyldimethylsilyl)-3-C-[{(S)carboxybenzylmethyl}-]3-deoxy-5-O-trityl-2,3- $\gamma$ -lactone- $\beta$ -D-ribofuranoside

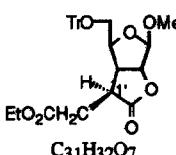
**Absolute configuration:** 1' S


 $[\alpha]_D^{25} = -49.4 \text{ (c 1, CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 2-O-crotonyl-3-O-(imidazol-1-yl)thiocarbonyl-5-O-trityl- $\beta$ -D-ribofuranoside

Methyl 3-C-[{(S)carboxyethylmethyl}-]3-deoxy-5-O-trityl-2,3- $\gamma$ -lactone- $\beta$ -D-ribofuranoside

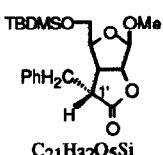
**Absolute configuration:** 1' S


 $[\alpha]_D^{25} = -54.5 \text{ (c 1, CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 2-O-ethylfumaroyl-3-O-(imidazol-1-yl)thiocarbonyl-5-O-trityl- $\beta$ -D-ribofuranoside

Methyl 3-C-[{(S)carboxy(etoxycarbonylmethyl)methyl}-]3-deoxy-5-O-trityl-2,3- $\gamma$ -lactone- $\beta$ -D-ribofuranoside

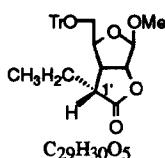
**Absolute configuration:** 1' S


 $[\alpha]_D^{25} = -105.6 \text{ (c 1, CHCl}_3\text{)}$ 

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 5-O-(*t*-butyldimethylsilyl)-2-O-cinnamoyl-3-O-(imidazol-1-yl)thiocarbonyl- $\beta$ -D-ribofuranoside

Methyl 5-O-(*t*-butyldimethylsilyl)-3-C-[{(R)carboxybenzylmethyl}-]3-deoxy-5-O-trityl-2,3- $\gamma$ -lactone- $\beta$ -D-ribofuranoside

**Absolute configuration:** 1' R

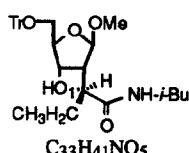


$[\alpha]_D^{25} = -40.4$  (c 1, CHCl<sub>3</sub>)

**Source of Chirality:** Asymmetric synthesis (radical cyclization) from methyl 2-O-crotonyl-3-O-(imidazol-1-yl)thiocarbonyl-5-O-trityl-beta-D-ribofuranoside

Methyl 3-C-[R]carboxyethylmethyl]-3-deoxy-5-O-trityl-2,3-gamma-lactone-beta-D-ribofuranoside

Absolute configuration: 1'R

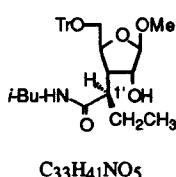


$[\alpha]_D^{25} = -17.2$  (c 1, CHCl<sub>3</sub>)

**Source of Chirality:** gamma-lactone-ring opening of methyl 2-C-[(R)carboxyethylmethyl]-2-deoxy-5-O-trityl-3,2-gamma-lactone-beta-D-ribofuranoside

Methyl 2-deoxy-2-C-[1'-(N-isobutyl)carbamoyl-1'(R)propyl]-5-O-trityl-beta-D-ribofuranoside

Absolute configuration: 1'R

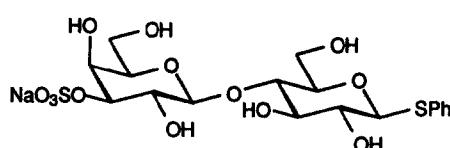


$[\alpha]_D^{25} = -0.4$  (c 1, CHCl<sub>3</sub>)

**Source of Chirality:** gamma-lactone-ring opening of methyl 3-C-[(S)carboxyethylmethyl]-3-deoxy-5-O-trityl-2,3-gamma-lactone-beta-D-ribofuranoside

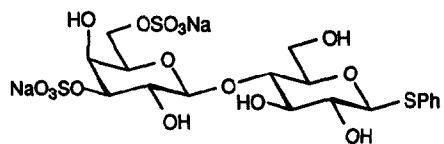
Methyl 3-deoxy-3-C-[1'-(N-isobutyl)carbamoyl-1'(S)-propyl]-5-O-trityl-beta-D-ribofuranoside

Absolute configuration: 1'S



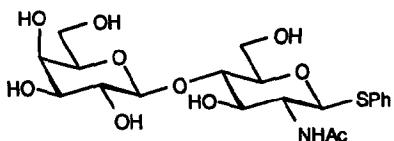
Phenyl 1-deoxy-4-O-(3'-O-sulfo-beta-D-galactopyranosyl)-1-thio-beta-D-glucopyranoside, sodium salt

$[\alpha]^{24}_D -26.2$  (c 4.8 in MeOH)



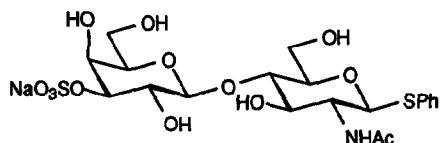
Phenyl 1-deoxy-4-O-(3',6'-di-O-sulfo- $\beta$ -D-galactopyranosyl)-1-thio- $\beta$ -D-glucopyranoside, disodium salt

$[\alpha]^{24}_D -29.9$  (*c* 1.5 in MeOH)



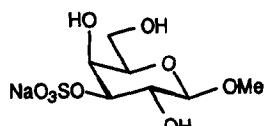
Phenyl 2-acetamido-1,2-di-deoxy-4-O-( $\beta$ -D-galactopyranosyl)-1-thio- $\beta$ -D-glucopyranoside

$[\alpha]^{23}_D +8.3$  (*c* 0.9 in H<sub>2</sub>O)



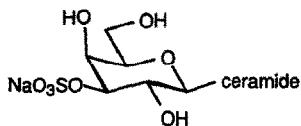
Phenyl 2-acetamido-1,2-di-deoxy-4-O-(3'-O-sulfo- $\beta$ -D-galactopyranosyl)-1-thio- $\beta$ -D-glucopyranoside, sodium salt

$[\alpha]^{24}_D -13.0$  (*c* 2.9 in MeOH)



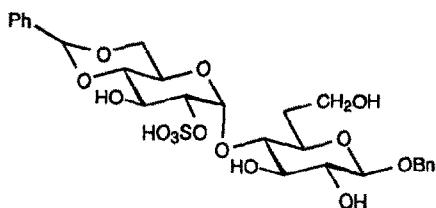
Methyl 3-O-sulfo- $\beta$ -D-galactopyranoside, sodium salt

$[\alpha]^{23}_D +8.3$  (*c* 3.6 in MeOH)



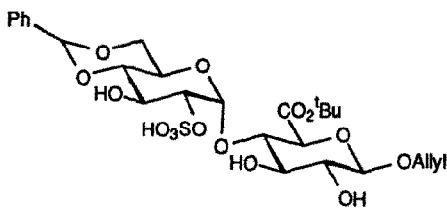
3-O-Sulfo- $\beta$ -D-galactosylceramide, sodium salt

$[\alpha]^{23}_D +2.6$  (*c* 1.0 in MeOH)



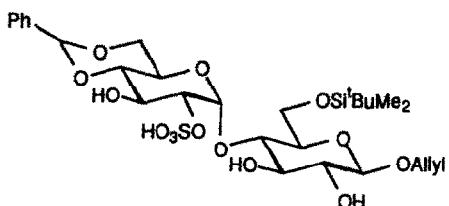
Benzyl 4-O-(4',6'-O-benzylidene-2'-O-sulfo- $\alpha$ -D-glucopyranosyl)- $\beta$ -D-glucopyranoside

$[\alpha]^{23}_D +26.0$  (*c* 1.0 in MeOH)



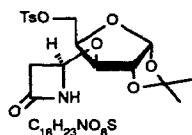
tert Butyl [allyl 4-O-(4',6'-O-benzylidene-2'-O-sulfo- $\alpha$ -D-glucopyranosyl)- $\beta$ -D-glucopyranosid]uronate

$[\alpha]^{23}_D +29.2$  (*c* 1.17 in methanol)



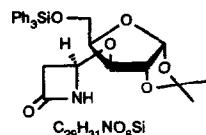
Allyl 4-O-(4',6'-O-benzylidene-2'-O-sulfo- $\alpha$ -D-glucopyranosyl)-6-O-tert-butyldimethylsilyl- $\beta$ -D-glucopyranoside

$[\alpha]^{25}_D +32.2$  (*c* 1.03 in MeOH)



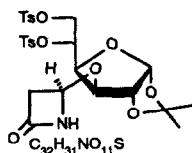
$[\alpha]_D = -24.5$  (*c* 0.3,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

(R)-3-O-(azetidin-2'-onyl-4')-1,2-O-isopropylidene-5-O-tosyl- $\alpha$ -D-xylofuranose.



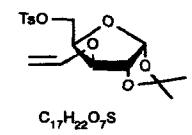
D.e. > 99%  
 $[\alpha]_D = -25.6$  (*c* 1,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

(R)-3-O-(azetidin-2'-onyl-4')-1,2-O-isopropylidene-5-O-triphenylsilyl- $\alpha$ -D-xylofuranose.



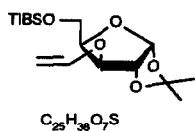
D.e. > 99%  
 $[\alpha]_D = -22.9$  (*c* 0.6,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

(R)-3-O-(azetidin-2'-onyl-4')-1,2-O-isopropylidene-5,6-di-O-tosyl- $\alpha$ -D-glucofuranose.



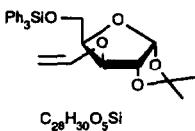
$[\alpha]_D = -19.9$  (*c* 1.2,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

1,2-O-isopropylidene-5-O-tosyl-3-O-vinyl- $\alpha$ -D-xylofuranose.



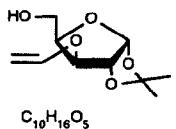
$[\alpha]_D = -11.6$  (*c* 2.2,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

1,2-O-isopropylidene-5-O-triisopropylbenzenesulfonyl-3-O-vinyl- $\alpha$ -D-xylofuranose.



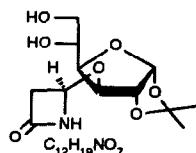
$[\alpha]_D = -42.5$  (*c* 0.1,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

1,2-O-isopropylidene-5-O-triphenylsilyl-3-O-vinyl- $\alpha$ -D-xylofuranose.



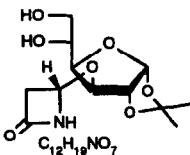
$[\alpha]_D = -47.0$  (*c* 0.3,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

1,2-O-isopropylidene-3-O-vinyl- $\alpha$ -D-xylofuranose.

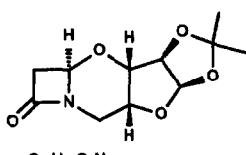


D.e. 66.7%  
 $[\alpha]_D = -5.7$  (*c* 1,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

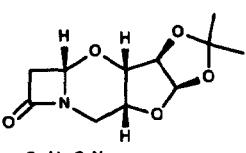
(4'R)-3-O-(azetidin-2'-onyl-4')-1,2-O-isopropylidene- $\alpha$ -D-glucofuranose.



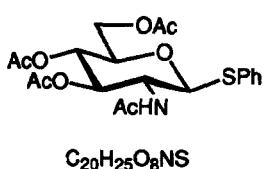
$[\alpha]_D = -13.6$  (*c* 0.14,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

(4'S) 3-O-(azetidin-2'-onyl-4')-1,2-O-isopropylidene- $\alpha$ -D-glucofuranose.

D.e. 3.3%  
 $[\alpha]_D = 121.1$  (*c* 0.9,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

(4'R) 5-amino-5-deoxy-1,2-O-isopropylidene-3-O:5-N-(azetidin-2'-onyl-4')- $\alpha$ -D-xylofuranose.

$[\alpha]_D = -29.8$  (*c* 0.8,  $\text{CH}_2\text{Cl}_2$ )  
Source of chirality: D-glucose

(4'S) 5-amino-5-deoxy-1,2-O-isopropylidene-3-O:5-N-(azetidin-2'-onyl-4')- $\alpha$ -D-xylofuranose.

$[\alpha]_D = -24$  (*c* 1.0,  $\text{CHCl}_3$ )

m.p. = 210-212°C

Source of chirality: 2-Amino-2-deoxy-D-glucose

Phenyl 2-Acetamido-3,4,6-tri-O-acetyl-2-deoxy-1-thio- $\beta$ -D-glucopyranoside

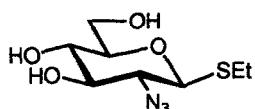
 $[\alpha]_D = -42$  (c 1.0, CHCl<sub>3</sub>)

m.p = 194-196°C

Source of chirality: 2-Amino-2-deoxy-D-glucose

C<sub>16</sub>H<sub>25</sub>O<sub>8</sub>NS

Ethyl 2-Acetamido-3,4,6-tri-O-acetyl-2-deoxy-1-thio-beta-D-glucopyranoside

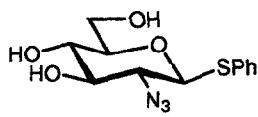
 $[\alpha]_D = -68$  (c 1.0, MeOH)

m.p = 103-104°C

Source of chirality: 2-Amino-2-deoxy-D-glucose

C<sub>8</sub>H<sub>15</sub>O<sub>4</sub>N<sub>3</sub>S

Ethyl 2-Azido-2-deoxy-1-thio-beta-D-glucopyranoside

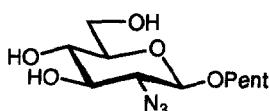
 $[\alpha]_D = -29$  (c 1.0, MeOH)

m.p = 112-114°C

Source of chirality: 2-Amino-2-deoxy-D-glucose

C<sub>12</sub>H<sub>15</sub>O<sub>4</sub>N<sub>3</sub>S

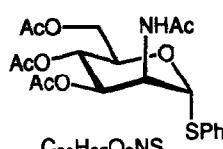
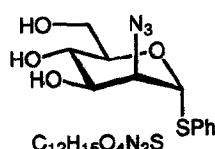
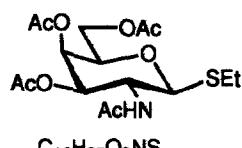
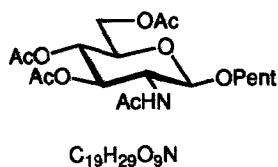
Phenyl 2-Azido-2-deoxy-1-thio-beta-D-glucopyranoside

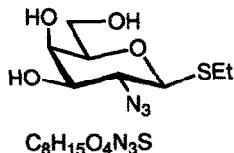
 $[\alpha]_D = -10$  (c 1.0, MeOH)

Source of chirality: 2-Amino-2-deoxy-D-glucose

C<sub>19</sub>H<sub>29</sub>O<sub>9</sub>N

4-Pentenyl 2-Azido-2-deoxy-beta-D-glucopyranoside

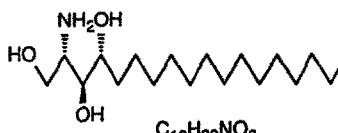




$[\alpha]_D = -15$  (c 1.0, MeOH)

Source of chirality: 2-Amino-2-deoxy-D-galactose

Ethyl 2-Azido-2-deoxy-β-D-galactopyranoside



(2S,3S,4R)-2-Amino-1,3,4-octadecanetriol

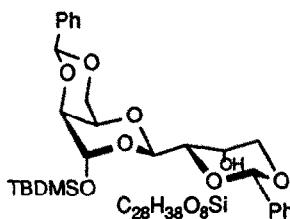
E.e. = 100 %

$[\alpha]_D^{20} = +7.7$  (c = 1, pyridine)

Source of chirality: natural (D-galactose), inversion (C-2), asymm. synth. (Grignard or reduction)

Absolute configuration: 2S,3S,4R

(identical with natural phytosphingosine)



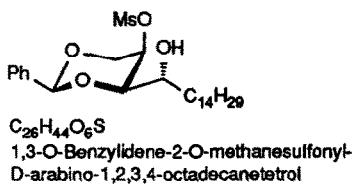
1-O-tert-Butyldimethylsilyl-bis-(2,4-O-benzylidene-D-threose)-1,1':3,1'-acetal

E.e. = 100 %

$[\alpha]_D^{23} = +25$  (c = 1, CHCl<sub>3</sub>)

Source of chirality: natural (D-galactose), asymm. induction (acetalization)

Absolute configuration: 1R,2S,3R,1'R,2'S,3'R  
(natural configuration; assigned by NMR-spectroscopy)



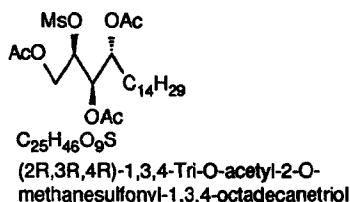
1,3-O-Benzylidene-2-O-methanesulfonyl-D-arabino-1,2,3,4-octadecanetetrol

E.e. = 100 %

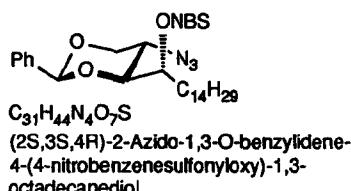
Source of chirality: natural (D-galactose), asymm. synth. (Grignard or reduction)

Absolute configuration: 2R,3R,4R

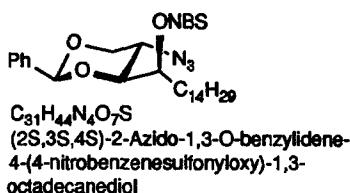
(2R,3R: natural configuration; 4R assigned by NMR-spectroscopy)



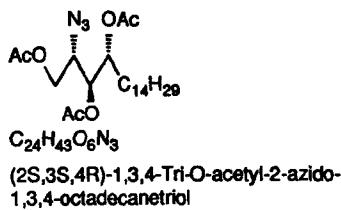
E.e. = 100 %  
 $[\alpha]_D^{20} = +25$  ( $c = 1, \text{CHCl}_3$ )  
 Source of chirality: natural (D-galactose), asymm. synth. (Grignard or reduction)  
 Absolute configuration: 2R,3R,4R  
 (2R,3R: natural configuration; 4R: assigned by NMR-spectroscopy)



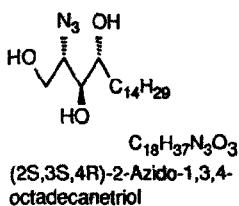
E.e. = 100 %  
 $[\alpha]_D^{20} = +31.8$  ( $c = 2, \text{CHCl}_3$ )  
 Source of chirality: natural (D-galactose), inversion (C-2), asymm. synth. (Grignard or reduction)  
 Absolute configuration: 2S,3S,4R  
 (2S: inverted natural configuration; 3S natural configuration, 4R: assigned by NMR-spectroscopy of derivatives)



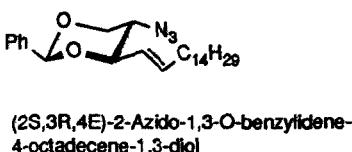
E.e. = 100 %  
 $[\alpha]_D^{20} = +2.1$  ( $c = 0.76, \text{CHCl}_3$ )  
 Source of chirality: natural (D-galactose), inversion (C-2), asymm. synth. (Grignard )  
 Absolute configuration: 2S,3S,4S  
 (2S: inverted natural configuration; 3S natural configuration, 4S: assigned by NMR-spectroscopy of derivatives)



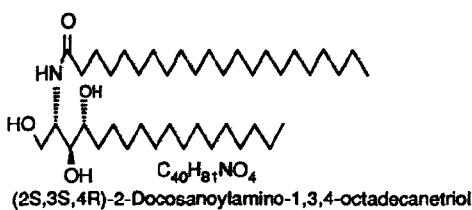
E.e. = 100 %  
 $[\alpha]_D^{20} = +16$  ( $c = 1, \text{CHCl}_3$ )  
 Source of chirality: natural (D-galactose), inversion (C-2), asymm. synth. (Grignard or reduction)  
 Absolute configuration: 2S,3S,4R  
 (2S: inverted natural configuration; 3S natural configuration, 4R: assigned by NMR-spectroscopy of derivatives)



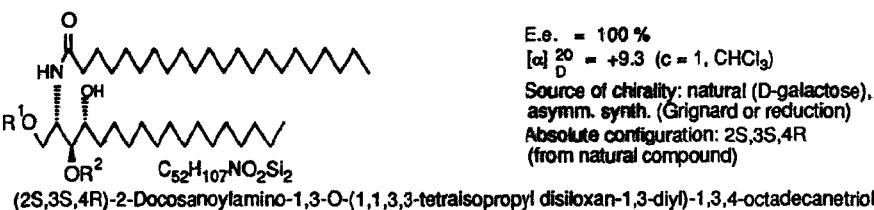
E.e. = 100 %  
 $[\alpha]_D^{20} = +17$  ( $c = 0.25$ ,  $\text{CHCl}_3/\text{MeOH}$  1:1)  
 Source of chirality: natural (D-galactose), inversion (C-2),  
 asymm. synth. (Grignard or reduction)  
 Absolute configuration: 2S,3S,4R  
 (2S: inverted natural configuration; 3S natural configuration,  
 4R: assigned by NMR-spectroscopy of derivatives)

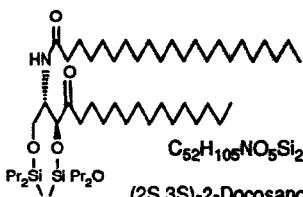


E.e. = 100 %  
 $[\alpha]_D^{20} = +11.7$  ( $c = 3$ ,  $\text{CHCl}_3$ )  
 Source of chirality: natural (D-galactose), inversion (C-2),  
 Absolute configuration: 2S,3R  
 (2S: inverted natural configuration; 3R natural configuration,



E.e. = 100 %  
 $[\alpha]_D^{20} = +7.2$  ( $c = 0.25$ ,  $\text{CHCl}_3/\text{MeOH}$  1:1)  
 Source of chirality: natural (D-galactose), inversion (C-2),  
 asymm. synth. (Grignard or reduction)  
 Absolute configuration: 2S,3S,4R  
 (from natural compound)





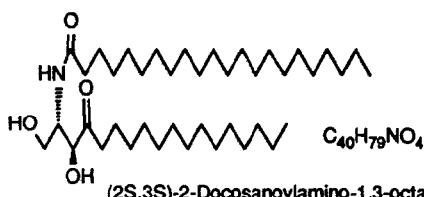
E.e. = 100 %

 $[\alpha]_D^{20} = +9$  ( $c = 1, CHCl_3$ )

Source of chirality: natural (D-galactose), inversion (C-2),

Absolute configuration: 2S,3S  
(from natural compound)

(2S,3S)-2-Docosanoylamino-1,3-O-(1,1,3,3-tetraisopropylidilsiloxan-1,3-diy)-1,3-octadecanediol-4-one



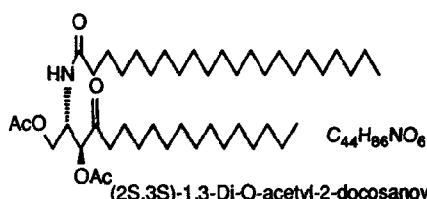
E.e. = 100 %

 $[\alpha]_D^{20} = +18$  ( $c = 0.25, CHCl_3/MeOH 1:1$ )

Source of chirality: natural (D-galactose), inversion (C-2),

Absolute configuration: 2S,3S  
(from natural compound)

(2S,3S)-2-Docosanoylamino-1,3-octadecanediol-4-one



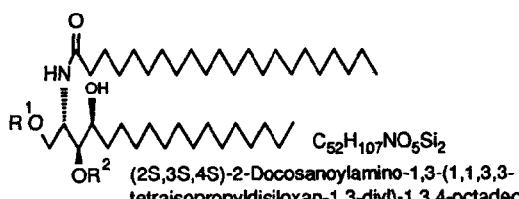
E.e. = 100 %

 $[\alpha]_D^{20} = +3.4$  ( $c = 0.5, CHCl_3$ )

Source of chirality: natural (D-galactose), inversion (C-2),

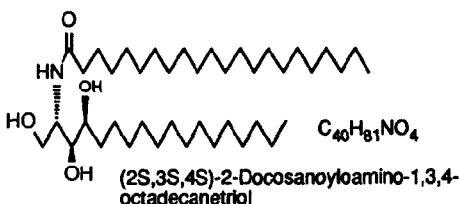
Absolute configuration: 2S,3S  
(from natural compound)

(2S,3S)-1,3-Di-O-acetyl-2-docosanoylamino-1,3-octadecanediol-4-one



E.e. = 100 %

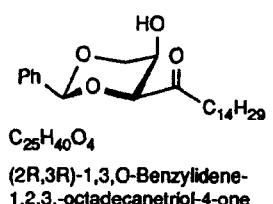
 $[\alpha]_D^{20} = -4$  ( $c = 1, CHCl_3$ )Source of chirality: natural (D-galactose), inversion (C-2),  
asymm. synth. (reduction)Absolute configuration: 2S,3S,4S  
(2S,3S: from natural compound, 4S: assigned by  
NMR-spectroscopy of derivatives)



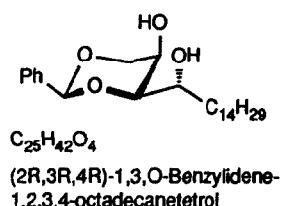
E.e. = 100 %  
 $[\alpha]_D^{20} = +7.6$  ( $c = 0.25$ ,  $\text{CHCl}_3/\text{MeOH}$  1:1)

Source of chirality: natural (D-galactose), inversion (C-2),  
 asymm. synth. (reduction)

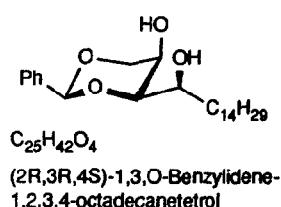
Absolute configuration: 2S,3S,4S  
 (2S,3S: from natural compound, 4S: assigned by  
 NMR-spectroscopy of derivatives)



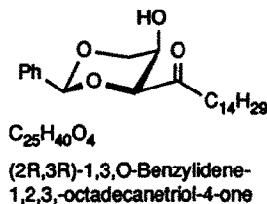
E.e. = 100 %  
 $[\alpha]_D^{20} = -55$  ( $c = 2$ ,  $\text{CHCl}_3$ )  
 Source of chirality: natural (D-galactose)  
 Absolute configuration 2R,3R  
 (natural configuration)



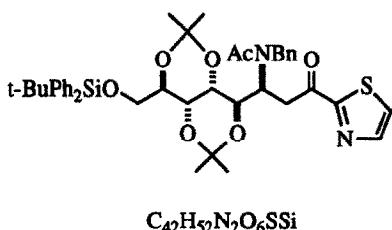
E.e. = 100 %  
 Source of chirality: natural (D-galactose)  
 Absolute configuration 2R,3R,4R  
 (2R,3R: natural configuration, 4R: assigned by NMR-spectroscopy of derivatives)



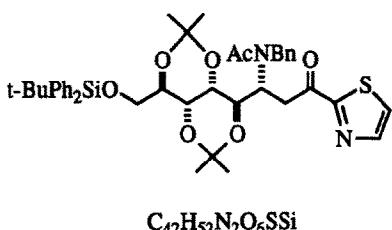
E.e. = 100 %  
 Source of chirality: natural (D-galactose)  
 Absolute configuration 2R,3R,4S  
 (2R,3R: natural configuration, 4S: assigned by NMR-spectroscopy of derivatives)



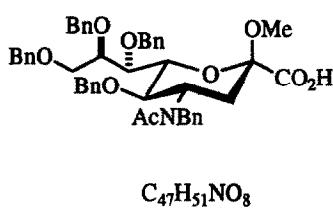
E.e. = 100 %  
 $[\alpha]_D^{20} = -55 (c = 2, \text{CHCl}_3)$   
Source of chirality: natural (D-galactose)  
Absolute configuration 2R,3R  
(natural configuration)



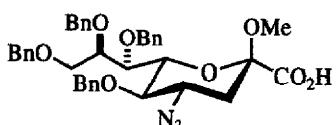
$[\alpha]_D^{20} = -60 (c = 0.7, \text{CHCl}_3)$   
Source of chirality: D-mannose  
Absolute configuration: 3S,4R,5R,6R,7R



$[\alpha]_D^{20} = +46 (c = 0.6, \text{CHCl}_3)$   
Source of chirality: D-mannose  
Absolute configuration: 3R,4R,5R,6R,7R

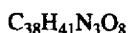


$[\alpha]_D^{20} = -26 (c = 0.5, \text{CHCl}_3)$   
Source of chirality: D-mannose  
Absolute configuration: 2S,4S,5R,6R,7R,8R

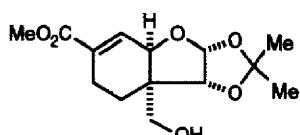
 $[\alpha]_D^{20} = -52$  ( $c = 1.3$ , CHCl<sub>3</sub>)

Source of chirality: D-mannose

Absolute configuration: 2S,4S,5R,6P,7R,8R

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217

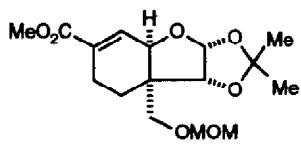
 $[\alpha]^{24}_D +62.4$  ( $c 1.33$ , CHCl<sub>3</sub>)

Source of chirality : D-glucose

Absolute configuration : 1R, 3R, 8R, 9R

8-(Hydroxymethyl)-5-(methoxycarbonyl)-11,11-dimethyl-2,10,12-trioxatricyclo[7.3.0.0<sup>3,8</sup>]dodec-4-eneJun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217



mp 123.0-124.0 °C

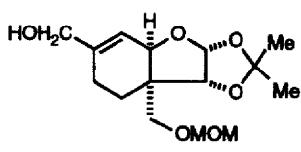
 $[\alpha]^{29}_D +43.7$  ( $c 1.10$ , CHCl<sub>3</sub>)

Source of chirality : D-glucose

Absolute configuration : 1R, 3R, 8R, 9R

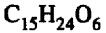
5-(Methoxycarbonyl)-8-[(methoxymethoxy)methyl]-11,11-dimethyl-2,10,12-trioxatricyclo[7.3.0.0<sup>3,8</sup>]dodec-4-eneJun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

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 $[\alpha]^{24}_D +55.7$  ( $c 1.11$ , CHCl<sub>3</sub>)

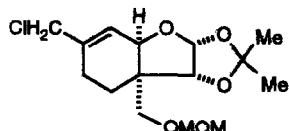
Source of chirality : D-glucose

Absolute configuration : 1R, 3R, 8R, 9R

5-(Hydroxymethyl)-8-[(methoxymethoxy)methyl]-11,11-dimethyl-2,10,12-trioxatricyclo[7.3.0.0<sup>3,8</sup>]dodec-4-ene

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217



5-(Chloromethyl)-8-[(methoxymethoxy)methyl]-11,11-dimethyl-2,10,12-trioxatricyclo[7.3.0.0^3,8]dodec-4-ene

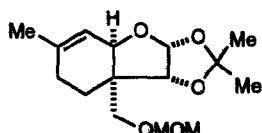
$[\alpha]^{24}_D +49.8$  (*c* 1.03,  $CHCl_3$ )

Source of chirality : D-glucose

Absolute configuration : 1*R*, 3*R*, 8*R*, 9*R*

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
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8-[(Methoxymethoxy)methyl]-5,11,11-trimethyl-2,10,12-trioxatricyclo[7.3.0.0^3,8]dodec-4-ene

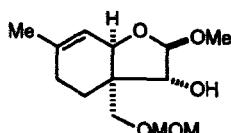
$[\alpha]^{24}_D +28.1$  (*c* 0.67,  $CHCl_3$ )

Source of chirality : D-glucose

Absolute configuration : 1*R*, 3*R*, 8*R*, 9*R*

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7-Hydroxy-8-methoxy-6-[(methoxymethoxy)methyl]-3-methyl-9-oxabicyclo[4.3.0]non-2-ene

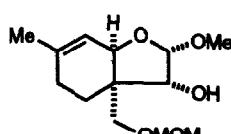
$[\alpha]^{21}_D -71.2$  (*c* 1.33,  $CHCl_3$ )

Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*, 8*R*

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217



7-Hydroxy-8-methoxy-6-[(methoxymethoxy)methyl]-3-methyl-9-oxabicyclo[4.3.0]non-2-ene

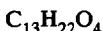
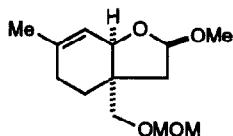
$[\alpha]^{20}_D +65.6$  (*c* 1.47,  $CHCl_3$ )

Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*, 8*S*

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217



8-Methoxy-6-[(methoxymethoxy)methyl]-3-methyl-9-oxabicyclo[4.3.0]non-2-ene

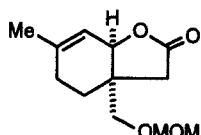
[α]<sub>D</sub><sup>23</sup> -90.6 (c 0.57, CHCl<sub>3</sub>)

Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 8*R*

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6-[(Methoxymethoxy)methyl]-3-methyl-9-oxabicyclo[4.3.0]non-2-en-8-one

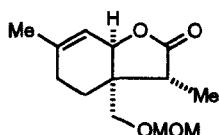
[α]<sub>D</sub><sup>23</sup> -10.6 (c 1.19, CHCl<sub>3</sub>)

Source of chirality : D-glucose

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

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and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217



6-[(Methoxymethoxy)methyl]-3,7-dimethyl-9-oxabicyclo[4.3.0]non-2-en-8-one

[α]<sub>D</sub><sup>23</sup> +15.9 (c 1.02, CHCl<sub>3</sub>)

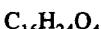
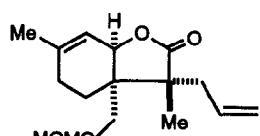
(containing ca. 6% of the 7*S* isomer)

Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

Tetrahedron: Asymmetry 1994, 5, 2217



7-Allyl-6-[(methoxymethoxy)methyl]-3,7-dimethyl-9-oxabicyclo[4.3.0]non-2-en-8-one

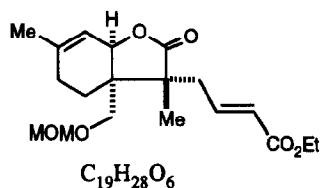
[α]<sub>D</sub><sup>21</sup> +2.5 (c 0.61, CHCl<sub>3</sub>)

Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

*Tetrahedron: Asymmetry* 1994, 5, 2217



[α]<sub>D</sub><sup>26</sup> +25.8 (c 0.72, CHCl<sub>3</sub>)

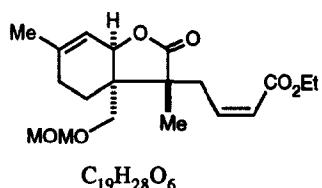
Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*

7-[(2*E*)-3-(Ethoxycarbonyl)-2-propenyl]-6-[(methoxymethoxy)-methyl]-3,7-methyl-9-oxabicyclo[4.3.0]non-2-en-8-one

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

*Tetrahedron: Asymmetry* 1994, 5, 2217



[α]<sub>D</sub><sup>26</sup> +3.6 (c 0.77, CHCl<sub>3</sub>)

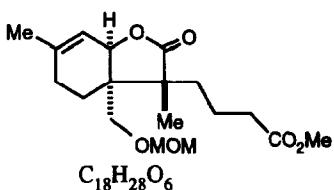
Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*

7-[(2*Z*)-3-(Ethoxycarbonyl)-2-propenyl]-6-[(methoxymethoxy)-methyl]-3,7-methyl-9-oxabicyclo[4.3.0]non-2-en-8-one

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,\*  
and Seiichiro Ogawa

*Tetrahedron: Asymmetry* 1994, 5, 2217



[α]<sub>D</sub><sup>26</sup> -4.5 (c 0.54, CHCl<sub>3</sub>)

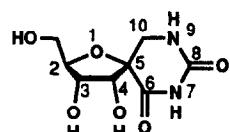
Source of chirality : D-glucose

Absolute configuration : 1*R*, 6*R*, 7*R*

7-[3-(Methoxycarbonyl)propyl]-6-[(methoxymethoxy)-methyl]-3,7-methyl-9-oxabicyclo[4.3.0]non-2-en-8-one

Hiromi Sano, Shigeru Mio, Junko Kitagawa, and Soji Sugai

*Tetrahedron: Asymmetry* 1994, 5, 2233



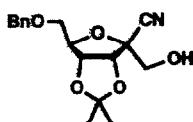
[α]<sub>D</sub><sup>25</sup> +63.8 (c 0.98, MeOH)

Source of chirality: natural and diastereoselective  
C-glycosidation

C<sub>8</sub>H<sub>12</sub>N<sub>2</sub>O<sub>6</sub>

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

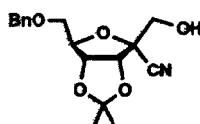
(2*R*,3*R*,4*R*,5*S*)-3,4-dihydroxymethyl-2-hydroxymethyl-1-oxa-7,9-diazaspiro[4.5]decane-6,8-dione

 $[\alpha]_D^{25} -29.7$  (c 1.72, CHCl<sub>3</sub>)

Source of chirality: natural and diastereoselective  
C-glycosidation

C&lt;sub&gt;17&lt;/sub&gt;H&lt;sub&gt;21&lt;/sub&gt;NO&lt;sub&gt;5&lt;/sub&gt;

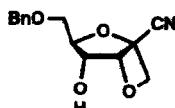
Absolute configuration 2R, 3R, 4R, 5R

6-O-Benzyl-3,4-O-isopropylidene-  
beta-D-psicofuranosyl cyanide $[\alpha]_D^{25} -13.1$  (c 1.22, MeOH)

Source of chirality: natural and diastereoselective  
C-glycosidation

C&lt;sub&gt;17&lt;/sub&gt;H&lt;sub&gt;21&lt;/sub&gt;NO&lt;sub&gt;5&lt;/sub&gt;

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

6-O-Benzyl-3,4-O-isopropylidene-  
alpha-D-psicofuranosyl cyanide $[\alpha]_D^{25} +74.5$  (c 1.10, MeOH)

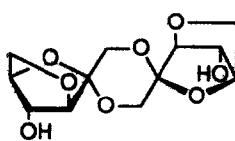
Source of chirality: natural and diastereoselective  
C-glycosidation

C&lt;sub&gt;14&lt;/sub&gt;H&lt;sub&gt;15&lt;/sub&gt;NO&lt;sub&gt;4&lt;/sub&gt;

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

1,3-Anhydro-6-O-benzyl-beta-D-psicofuranose  
cyanide

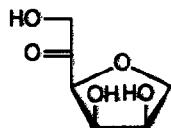
E.e. = 100%

 $[\alpha]_D = +77$  (c 1, H<sub>2</sub>O)

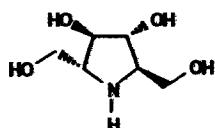
3,6-Anhydro-alpha-D-fructofuranose 3,6-Anhydro-beta-D-fructofuranose  
1,2':2,1'-dianhydride  
C<sub>12</sub>H<sub>16</sub>O<sub>8</sub>

Source of chirality: D-fructose as starting material

E.e. = 100%

 $[\alpha]_D = -49$  (c 1.1, H<sub>2</sub>O)3,6-Anhydro-keto-D-fructose  
C<sub>6</sub>H<sub>10</sub>O<sub>5</sub>

Source of chirality: D-fructose as starting material

 $[\alpha]_D +56.4$  (c, 7.0, H<sub>2</sub>O)

Source of chirality: natural and asymm. synth

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

C<sub>6</sub>H<sub>13</sub>NO<sub>4</sub>

2R, 5R-Dihydroxymethyl-3R, 4R-dihydroxypyrrolidine

E.e. = 100%

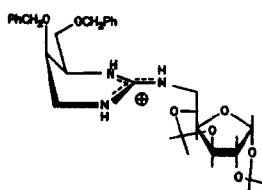
 $[\alpha]_D^{25} = +74.8$  (c, 1.0 in MeOH)(4R,5S)-2-[N-(6-Deoxy-1:2:3,5-di-O-isopropylidene- $\alpha$ -D-glucofuranosyl)amino]-benzylidene-5-oxy-4-oxymethyl-1,4,5,6-tetrahydropyrimidine  
C<sub>24</sub>H<sub>33</sub>N<sub>3</sub>O<sub>7</sub>

Source of chirality: D-glucose and L-arabinose as starting material

E.e. = 100%

 $[\alpha]_D^{25} = +38.7$  (c, 1.60 in H<sub>2</sub>O)(4R,5S)-2-[N-(6-Deoxy- $\alpha$ / $\beta$ -D-glucopyranosyl)amino]-5-hydroxy-4-hydroxymethyl-1,4,5,6-tetrahydropyrimidine  
C<sub>11</sub>H<sub>22</sub>N<sub>3</sub>O<sub>7</sub>

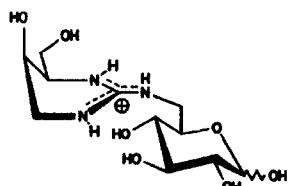
Source of chirality: D-glucose and L-arabinose as starting material



E.e. = 100%  
 $[\alpha]_D^{23} = -5.0(c, 2.00 \text{ in EtOH})$

(4R,5R)-2-[N-(6-Deoxy-1,2:3,5-di-O-isopropylidene- $\alpha$ -D-glucofuranosyl)amino]-5-O-benzyl-4-O-benzylmethyl-1,4,5,6-tetrahydropyrimidine  
 $C_{31}H_{44}N_3O_2$

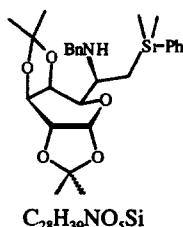
Source of chirality: D-glucose as starting material



E.e. = 100%  
 $[\alpha]_D^{23} = +7.9(c, 2.28 \text{ in H}_2\text{O})$

(4R,5R)-2-[N-(6-Deoxy- $\alpha$ -D-glucopyranosyl)amino]-5-hydroxy-4-hydroxymethyl-1,4,5,6-tetrahydropyrimidine  
 $C_{11}H_{22}N_3O_4$

Source of chirality: D-glucose as starting material

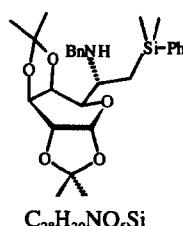


D.e. = >97% (by 300 MHz  $^1\text{H}$  NMR spectroscopy)

$[\alpha]_D^{20} -50.2$  (c 1.0,  $\text{CHCl}_3$ )

Source of chirality: natural and asymm. synth. (nucleophilic addition)

Absolute configuration: 2R, 3S, 4S, 5R, 6R  
 (assigned via chemical conversion)

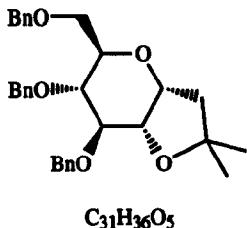


D.e. = >97% (by 300 MHz  $^1\text{H}$  NMR spectroscopy)

$[\alpha]_D^{20} -34.0$  (c 1.0,  $\text{CHCl}_3$ )

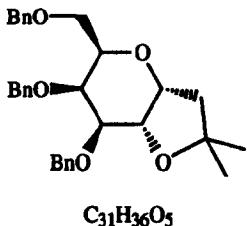
Source of chirality: natural and asymm. synth. (nucleophilic addition)

Absolute configuration: 2R, 3S, 4S, 5R, 6S  
 (assigned via chemical conversion)

E.e. = 100% (by  $^1H$  NMR) $[\alpha]_D = +46.8$  ( $c = 5.49$ ,  $CHCl_3$ )

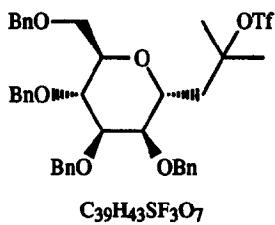
Source of chirality: D-glucose

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

E.e. = 100% (by  $^1H$  NMR) $[\alpha]_D = +24.6$  ( $c = 5.95$ ,  $CHCl_3$ )

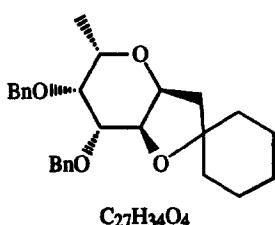
Source of chirality: D-galactose

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

E.e. = 100% (by  $^1H$  NMR) $[\alpha]_D = +21.3$  ( $c = 6.16$ ,  $CHCl_3$ )

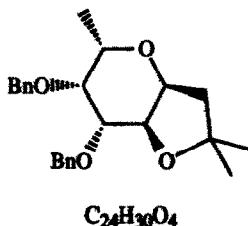
Source of chirality: D-mannose

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

E.e. = 100% (by  $^1H$  NMR) $[\alpha]_D = -22.6$  ( $c = 7.20$ ,  $CHCl_3$ )

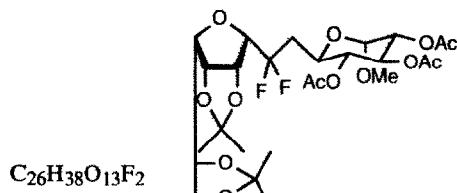
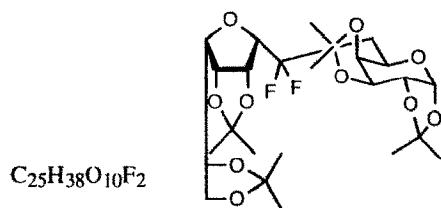
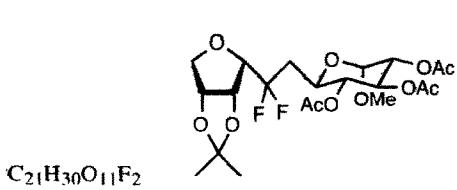
Source of chirality: L-fucose

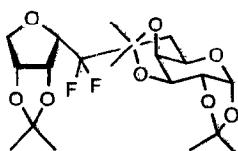
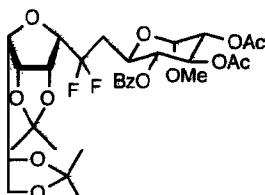
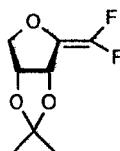
Absolute configuration: 1S,2R,3R,4R,5S

E.e. = 100% (by  $^1H$  NMR) $[\alpha]_D^{20} = -16.8$  ( $c = 6.48$ ,  $CHCl_3$ )

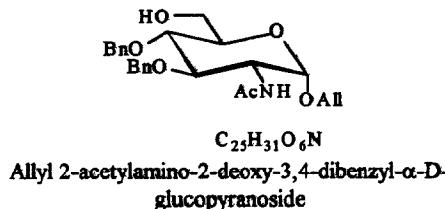
Source of chirality: L-fucose

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

 $[\alpha]_D^{20} = +72.7$  ( $c = 0.22$ ,  $CHCl_3$ )Source of chirality: D-galactofuranose,  $\alpha$ -D-glucopyranoside, stereoselective radical addition.Absolute configuration 1S,2R,3S,4R,5R,8S,9R,10R,11S,12R  
(assigned by  $^1H$  NMR coupling)Methyl 2,3,4-tri-O-acetyl-8,11-anhydro-6,7-dideoxy-9,10,12,13-di-O-isopropylidene-7,7-difluoro-D-glycero-L-galacto- $\alpha$ -D-glucopyranoside. $[\alpha]_D^{20} = -43.8$  ( $c = 0.14$ ,  $CHCl_3$ )Source of chirality: D-galactofuranose,  $\alpha$ -D-galactopyranose, stereoselective radical addition.Absolute configuration 1R,2R,3S,4S,5R,8S,9R,10R,11S,12R  
(assigned by  $^1H$  NMR coupling)1,2,3,4;9,10;12,13-Tetra-O-isopropylidene-6,7-dideoxy-7,7-difluoro-8,11-anhydro-D-glycero-L-galacto- $\alpha$ -D-galacto-tridecose糖衍生物 $[\alpha]_D^{20} = +71.4$  ( $c = 0.7$ ,  $CHCl_3$ )Source of chirality: D-erythofuranose,  $\alpha$ -D-glucopyranoside, stereoselective radical addition.Absolute configuration 1S,2R,3S,4R,5R,8S,9R,10R  
(assigned by  $^1H$  NMR coupling)Methyl 2,3,4-tri-O-acetyl-9,10-O-isopropylidene-6,7-dideoxy-7,7-difluoro-8,11-anhydro-D-arabino- $\alpha$ -D-glucouradose糖衍生物.

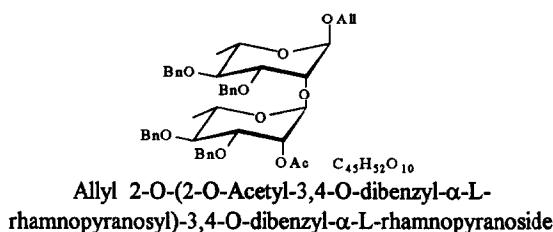
C<sub>20</sub>H<sub>30</sub>O<sub>8</sub>F<sub>2</sub>[ $\alpha$ ]<sub>D</sub><sup>20</sup>=-62.8 (c=0.18, CHCl<sub>3</sub>)Source of chirality: D-erythrofuranose,  $\alpha$ -D-galactopyranose, stereoselective radical addition.Absolute configuration 1R,2R,3S,4S,5R,8S,9R,10R  
(assigned by <sup>1</sup>H NMR coupling)1,2;3,4;9,10-Tri-O-isopropylidene-6,7-dideoxy-7,7-difluoro-8,11-anhydro-D-*arabino*- $\alpha$ -D-*galacto*-undecopyranose.C<sub>31</sub>H<sub>40</sub>F<sub>2</sub>O<sub>13</sub>[ $\alpha$ ]<sub>D</sub><sup>20</sup>=+56.8 (c=1.30, CHCl<sub>3</sub>)Source of chirality: D-galactofuranose,  $\alpha$ -D-glucopyranose, stereoselective radical addition.Absolute configuration 1S,2R,3S,4R,5R,8S,9R,10R,11S,12R  
(assigned by <sup>1</sup>H NMR coupling)Methyl 2,3-di-O-acetyl-8,11-anhydro-4-O-benzoyl-6,7-dideoxy-9,10,12,13-di-O-isopropylidene-7,7-difluoro-D-glycero-1-galacto- $\alpha$ -D-glucoside.C<sub>8</sub>H<sub>10</sub>F<sub>2</sub>O<sub>3</sub>[ $\alpha$ ]<sub>D</sub><sup>20</sup>=-206.0 (c=2.15, CH<sub>2</sub>Cl<sub>2</sub>)

Source of chirality: D-erythrofuranose

Absolute configuration 3R,4R  
(assigned by <sup>1</sup>H NMR coupling)2,5-Anhydro-1-deoxy-1,1-difluoro-3,4-O-isopropylidene-D-*erythro*-penta-1-enitol.e.e = 100% (<sup>1</sup>H NMR)[ $\alpha$ ]<sub>D</sub><sup>20</sup> +56.7 (c 1.3, CHCl<sub>3</sub>)

Source of chirality: asymm. synth.

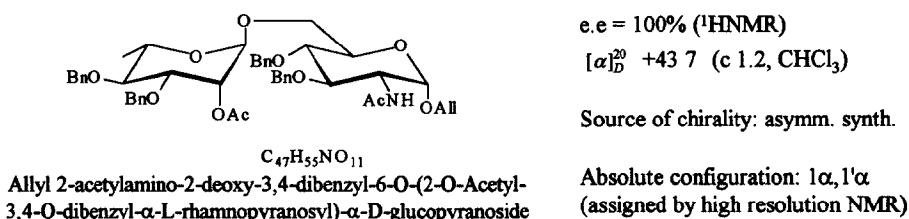
Absolute configuration: 1 $\alpha$   
(assigned by high resolution NMR)



e.e = 100% ( $^1\text{H}$ NMR)  
 $[\alpha]_D^{20} -19.3$  (c 1.2,  $\text{CHCl}_3$ )

Source of chirality: asymm. synth.

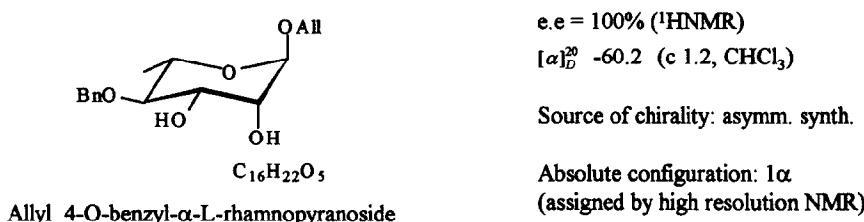
Absolute configuration:  $1\alpha, 1'\alpha$   
 (assigned by high resolution NMR)



e.e = 100% ( $^1\text{H}$ NMR)  
 $[\alpha]_D^{20} +43.7$  (c 1.2,  $\text{CHCl}_3$ )

Source of chirality: asymm. synth.

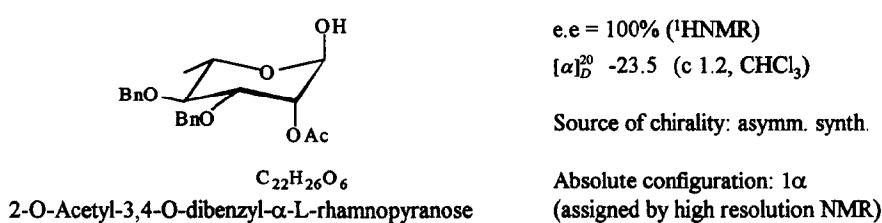
Absolute configuration:  $1\alpha, 1'\alpha$   
 (assigned by high resolution NMR)



e.e = 100% ( $^1\text{H}$ NMR)  
 $[\alpha]_D^{20} -60.2$  (c 1.2,  $\text{CHCl}_3$ )

Source of chirality: asymm. synth.

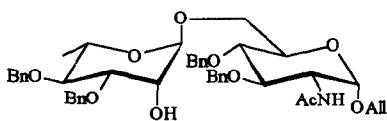
Absolute configuration:  $1\alpha$   
 (assigned by high resolution NMR)



e.e = 100% ( $^1\text{H}$ NMR)  
 $[\alpha]_D^{20} -23.5$  (c 1.2,  $\text{CHCl}_3$ )

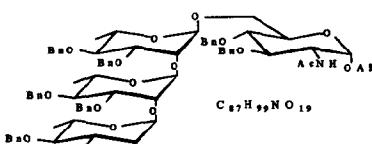
Source of chirality: asymm. synth.

Absolute configuration:  $1\alpha$   
 (assigned by high resolution NMR)

e.e = 100% (<sup>1</sup>H NMR)[α]<sub>D</sub><sup>20</sup> +34.7 (c 1.2, CHCl<sub>3</sub>)

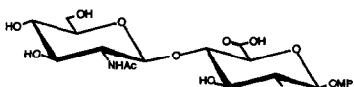
Source of chirality: asymm. synth.

Allyl 2-acetylamino-2-deoxy-3,4-dibenzyl-6-O-(3,4-O-dibenzyl-alpha-L-rhamnopyranosyl)-alpha-D-glucopyranoside  
C<sub>45</sub>H<sub>53</sub>NO<sub>10</sub>

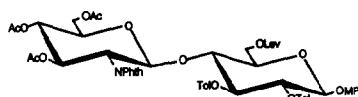
Absolute configuration: 1 $\alpha$ , 1' $\alpha$   
(assigned by high resolution NMR)e.e = 100% (<sup>1</sup>H NMR)[α]<sub>D</sub><sup>20</sup> +34.7 (c 1.4, CHCl<sub>3</sub>)

Source of chirality: asymm. synth.

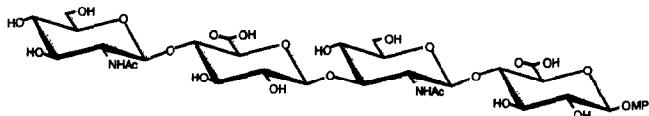
Allyl 2-acetylamino-2-deoxy-3,4-dibenzyl-6-O-[2-O-(2-O-acetyl-3,4-O-dibenzyl-alpha-L-rhamnopyranosyl)-3,4-O-dibenzyl-alpha-L-rhamnopyranosyl]-3,4-O-dibenzyl-alpha-L-rhamnopyranosyl-alpha-D-glucopyranoside

Absolute configuration: 1 $\alpha$ , 1' $\alpha$ , 1" $\alpha$ , 1" $\alpha$   
(assigned by high resolution NMR)C<sub>21</sub>H<sub>29</sub>NO<sub>13</sub>[α]<sub>D</sub> -40° (c 1, H<sub>2</sub>O)source of chirality: D-glucose  
D-glucosamine

4-Methoxyphenyl O-(2-acetamido-2-deoxy-β-D-glucopyranosyl)-(1→4)-β-D-glucopyranosyluronic acid

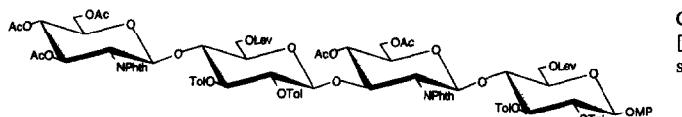
C<sub>54</sub>H<sub>55</sub>NO<sub>20</sub>[α]<sub>D</sub> +57° (c 1, CH<sub>2</sub>Cl<sub>2</sub>)source of chirality: D-glucose  
D-glucosamine

4-Methoxyphenyl O-(3,4,6-tri-O-acetyl-2-deoxy-2-phthalimido-beta-D-glucopyranosyl)-(1→4)-6-O-levulinoyl-2,3-di-O-p-toluoyle-beta-D-glucopyranoside



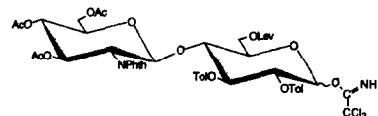
$C_{35}H_{50}N_2O_{24}$   
 $[\alpha]_D -45^\circ$  (c 1,  $H_2O$ )  
 source of chirality: D-glucose  
 D-glucosamine

4-Methoxyphenyl  $O$ -(2-acetamido-2-deoxy- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)- $O$ -( $\beta$ -D-glucopyranosyluronic acid)-(1 $\rightarrow$ 3)- $O$ -(2-acetamido-2-deoxy- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)- $\beta$ -D-glucopyranosyluronic acid



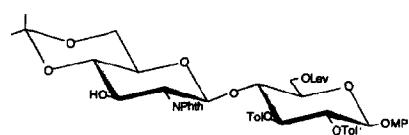
$C_{99}H_{100}N_2O_{37}$   
 $[\alpha]_D +38^\circ$  (c 1,  $CH_2Cl_2$ )  
 source of chirality: D-glucose  
 D-glucosamine

4-Methoxyphenyl  $O$ -(3,4,6-tri- $O$ -acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)- $O$ -(6- $O$ -levulinoyl-2,3-di- $O$ - $p$ -toluoyl- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 3)- $O$ -(4,6-di- $O$ -acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-6- $O$ -levulinoyl-2,3-di- $O$ - $p$ -toluoyl- $\beta$ -D-glucopyranoside



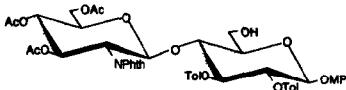
$C_{49}H_{49}Cl_3N_2O_{19}$   
 $[\alpha]_D +80^\circ$  (c 1,  $CH_2Cl_2$ ) ( $\alpha:\beta$  3:2)  
 source of chirality: D-glucose  
 D-glucosamine

$O$ -(3,4,6-Tri- $O$ -acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-6- $O$ -levulinoyl-2,3-di- $O$ - $p$ -toluoyl- $\alpha/\beta$ -D-glucopyranosyl trichloroacetimidate



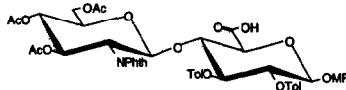
$C_{51}H_{53}NO_{17}$   
 $[\alpha]_D +31^\circ$  (c 1,  $CH_2Cl_2$ )  
 source of chirality: D-glucose  
 D-glucosamine

4-Methoxyphenyl  $O$ -(2-deoxy-4,6- $O$ -isopropylidene-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-6- $O$ -levulinoyl-2,3-di- $O$ - $p$ -toluoyl- $\beta$ -D-glucopyranoside



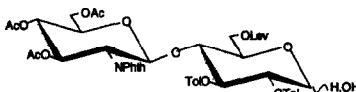
C<sub>49</sub>H<sub>49</sub>NO<sub>18</sub>  
[α]<sub>D</sub> +86° (c 1, CH<sub>2</sub>Cl<sub>2</sub>)  
source of chirality: D-glucose  
D-glucosamine

4-Methoxyphenyl O-(3,4,6-tri-O-acetyl-2-deoxy-2-phthalimido-β-D-glucopyranosyl)-(1→4)-2,3-di-O-p-toloyl-β-D-glucopyranoside



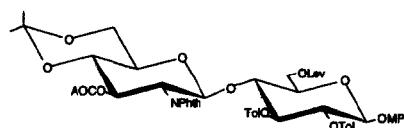
C<sub>49</sub>H<sub>47</sub>NO<sub>19</sub>  
[α]<sub>D</sub> +4° (c 1, CH<sub>2</sub>Cl<sub>2</sub>)  
source of chirality: D-glucose  
D-glucosamine

4-Methoxyphenyl O-(3,4,6-tri-O-acetyl-2-deoxy-2-phthalimido-β-D-glucopyranosyl)-(1→4)-2,3-di-O-p-toloyl-β-D-glucopyranosyluronic acid



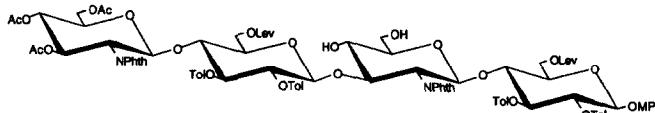
C<sub>47</sub>H<sub>49</sub>NO<sub>19</sub>  
[α]<sub>D</sub> +93° (c 1, CH<sub>2</sub>Cl<sub>2</sub>) (α:β 2.7:1)  
source of chirality: D-glucose  
D-glucosamine

O-(3,4,6-Tri-O-acetyl-2-deoxy-2-phthalimido-β-D-glucopyranosyl)-(1→4)-6-O-levulinoyl-2,3-di-O-p-toloyl-α/β-D-glucopyranose



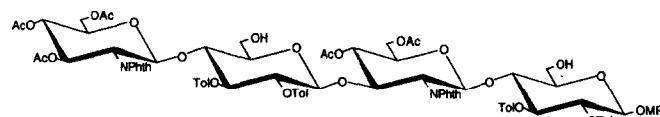
C<sub>55</sub>H<sub>57</sub>NO<sub>19</sub>  
[α]<sub>D</sub> +25° (c 1, CH<sub>2</sub>Cl<sub>2</sub>)  
source of chirality: D-glucose  
D-glucosamine

4-Methoxyphenyl O-(3-O-allyloxycarbonyl-2-deoxy-4,6-O-isopropylidene-2-phthalimido-β-D-glucopyranosyl)-(1→4)-6-O-levulinoyl-2,3-di-O-p-toloyl-β-D-glucopyranoside



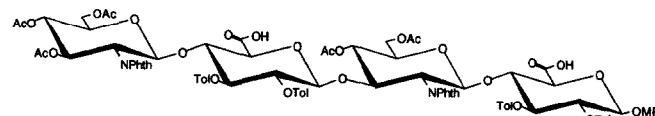
C<sub>95</sub>H<sub>96</sub>N<sub>2</sub>O<sub>35</sub>  
 $[\alpha]_D^{20} +70^\circ$  (c 1, CH<sub>2</sub>Cl<sub>2</sub>)  
 source of chirality: D-glucose  
 D-glucosamine

4-Methoxyphenyl *O*-(3,4,6-tri-*O*-acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-*O*-(6-*O*-levulinoyl-2,3-di-*O*-*p*-toluoyl- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 3)-*O*-(2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-6-*O*-levulinoyl-2,3-di-*O*-*p*-toluoyl- $\beta$ -D-glucopyranoside



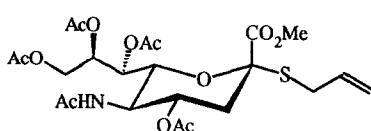
C<sub>89</sub>H<sub>88</sub>N<sub>2</sub>O<sub>33</sub>  
 $[\alpha]_D^{20} +59^\circ$  (c 1, CH<sub>2</sub>Cl<sub>2</sub>)  
 source of chirality: D-glucose  
 D-glucosamine

4-Methoxyphenyl *O*-(3,4,6-tri-*O*-acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-*O*-(2,3-di-*O*-*p*-toluoyl- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 3)-*O*-(4,6-di-*O*-acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-2,3-di-*O*-*p*-toluoyl- $\beta$ -D-glucopyranoside



C<sub>89</sub>H<sub>84</sub>N<sub>2</sub>O<sub>35</sub>  
 $[\alpha]_D^{20} +6^\circ$  (c 1, CH<sub>2</sub>Cl<sub>2</sub>)  
 source of chirality: D-glucose  
 D-glucosamine

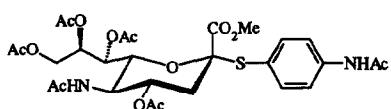
4-Methoxyphenyl *O*-(3,4,6-tri-*O*-acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-*O*-(2,3-di-*O*-*p*-toluoyl- $\beta$ -D-glucopyranosyluronic acid)-(1 $\rightarrow$ 3)-*O*-(4,6-di-*O*-acetyl-2-deoxy-2-phthalimido- $\beta$ -D-glucopyranosyl)-(1 $\rightarrow$ 4)-2,3-di-*O*-*p*-toluoyl- $\beta$ -D-glucopyranosyluronic acid



$[\alpha]_D^{23} = +38.9$  (c 1.0, CHCl<sub>3</sub>)  
 m.p. 108.5-111 °C  
 Source of chirality: N-acetylneuraminic acid as starting material

C<sub>23</sub>H<sub>33</sub>NO<sub>12</sub>S

Allyl (methyl 5-acetamido-4,7,8,9-tetra-*O*-acetyl-3,5-dideoxy-2-thio-D-glycero- $\alpha$ -D-galacto-2-nonulopyranosid) onate



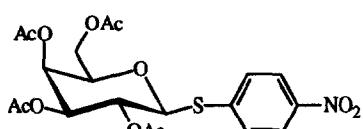
$[\alpha]_D^{23} = +33.9$  (c 1.86, CHCl<sub>3</sub>)

m.p. 97-98 °C

Source of chirality: N-acetylneuraminic acid as starting material

C<sub>28</sub>H<sub>36</sub>N<sub>2</sub>O<sub>13</sub>S

4-N-acetamidophenyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-alpha-D-galacto-2-nonulopyranosid) onate



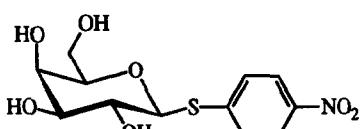
$[\alpha]_D^{23} = -8.3$  (c 1.0, CHCl<sub>3</sub>)

m.p. 168.7-169 °C

Source of chirality: D-galactose as starting material

C<sub>13</sub>H<sub>23</sub>NO<sub>11</sub>S

4-Nitrophenyl 2,3,4,6-tetra-O-acetyl-1-thio-beta-D-galactopyranoside



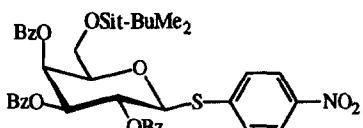
$[\alpha]_D^{23} = -100.8$  (c 1.0, CH<sub>3</sub>OH)

m.p. 160.2-161.5 °C

Source of chirality: D-galactose as starting material

C<sub>12</sub>H<sub>15</sub>NO<sub>7</sub>S

4-Nitrophenyl 1-thio-beta-D-galactopyranoside



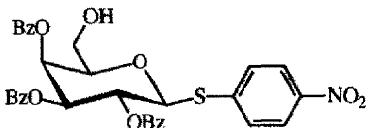
$[\alpha]_D^{23} = +84.4$  (c 1.0, CHCl<sub>3</sub>)

m.p. 159-160 °C

Source of chirality: D-galactose as starting material

C<sub>39</sub>H<sub>41</sub>NO<sub>10</sub>SSI

4-Nitrophenyl 2,3,4-tri-O-benzoyl-6-O-t-butyldimethylsilyl-1-thio-beta-D-galactopyranoside

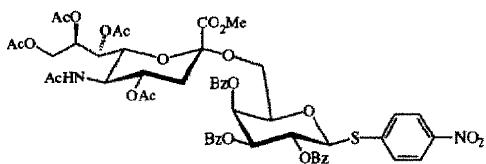


$[\alpha]_D^{23} = +69.9^\circ$  (c 1.0, CHCl<sub>3</sub>)

m.p. 106.5-107.7 °C

Source of chirality: D-galactose as starting material

C<sub>33</sub>H<sub>27</sub>NO<sub>10</sub>S  
4-Nitrophenyl 2,3,4-tri-O-benzoyl-1-thio-β-D-galactopyranoside

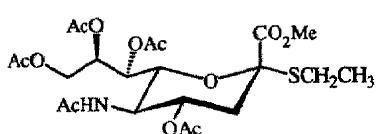


$[\alpha]_D^{23} = +45.2^\circ$  (c 1.1, CHCl<sub>3</sub>)

m.p. 117.4-118.5 °C

Source of chirality: N-acetylneuraminic acid and D-galactose as starting material

C<sub>53</sub>H<sub>34</sub>N<sub>2</sub>O<sub>21</sub>S  
4-Nitrophenyl O-(methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-D-glycero-α-D-galacto-2-nonulopyranosylonate)-(2→6)-2,3,4-tri-O-benzoyl-1-thio-β-D-galactopyranoside

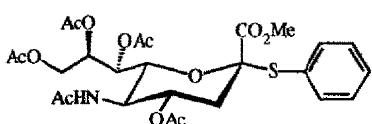


$[\alpha]_D^{23} = +31.4^\circ$  (c 1.0, CHCl<sub>3</sub>)

m.p. 83-84 °C

Source of chirality: N-acetylneuraminic acid as starting material

C<sub>22</sub>H<sub>33</sub>NO<sub>12</sub>S  
Ethyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-α-D-galacto-2-nonulopyranosid) onate

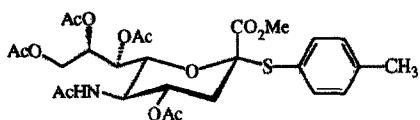


$[\alpha]_D^{23} = +20.9^\circ$  (c 1.0, CHCl<sub>3</sub>)

m.p. 141-142 °C

Source of chirality: N-acetylneuraminic acid as starting material

C<sub>26</sub>H<sub>33</sub>NO<sub>12</sub>S  
Phenyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-α-D-galacto-2-nonulopyranosid) onate

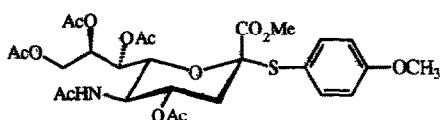
 $[\alpha]_D^{23} = +24.0$  (c 1.0, CHCl<sub>3</sub>)

m.p. 114-115 °C

Source of chirality: N-acetyleneuraminic acid as starting material

C<sub>27</sub>H<sub>35</sub>NO<sub>12</sub>S

4-Methylphenyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-alpha-D-galacto-2-nonulopyranosid) onate

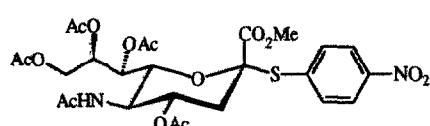
 $[\alpha]_D^{23} = +20.7$  (c 1.0, CHCl<sub>3</sub>)

m.p. 136-137 °C

Source of chirality: N-acetyleneuraminic acid as starting material

C<sub>27</sub>H<sub>35</sub>NO<sub>13</sub>S

4-Methoxyphenyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-alpha-D-galacto-2-nonulopyranosid) onate

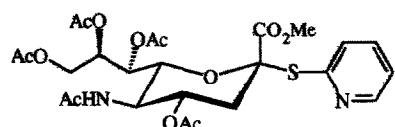
 $[\alpha]_D^{23} = +18.2$  (c 1.0, CHCl<sub>3</sub>)

m.p. 170-171 °C

Source of chirality: N-acetyleneuraminic acid as starting material

C<sub>26</sub>H<sub>32</sub>N<sub>2</sub>O<sub>14</sub>S

4-Nitrophenyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-alpha-D-galacto-2-nonulopyranosid) onate

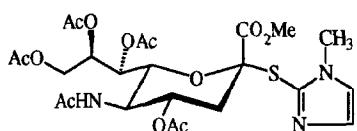
 $[\alpha]_D^{23} = +29.0$  (c 1.0, CHCl<sub>3</sub>)

m.p. 150-152 °C

Source of chirality: N-acetyleneuraminic acid as starting material

C<sub>25</sub>H<sub>32</sub>N<sub>2</sub>O<sub>12</sub>S

2-Pyridyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero-alpha-D-galacto-2-nonulopyranosid) onate



$[\alpha]_D^{23} = +17.7$  (c 1.0, CHCl<sub>3</sub>)

m.p. 140-142 °C

Source of chirality: N-acetylneurameric acid as starting material

C<sub>24</sub>H<sub>33</sub>N<sub>3</sub>O<sub>12</sub>S

2-N<sup>1</sup>-methylimidazolyl (methyl 5-acetamido-4,7,8,9-tetra-O-acetyl-3,5-dideoxy-2-thio-D-glycero- $\alpha$ -D-galacto-2-nonulopyranosid) onate