

Medtronic

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AUTHOR


DARRYL ZURN

REVISIONS

REV	DESCRIPTION	DATE
A	Initial release	23 April 2021
B	Remove Compatibility Graphic. Remove CE Mark and EC Rep address.	4 June 2021

PART NUMBER	REV	SHEET
B05013	B	1 of 7

1. This specification is a representation of the final Label Design Output generated from a combination of label parts within the associated PDM BOM.
2. These label images are the combinations of the Transform template file, LDS (Label Model Data) record, sample trigger data and representative views of the label die lines.
3. The Serial Number, Use-By Date, Lot Number, Date Of Manufacture, PIN Number, GTIN, GTIN Version (if used), Product Number (REF) and Barcodes are shown for example only. These fields are variable and are to be supplied by Manufacturing at the time of printing. Not all fields will be used for all labels.
4. The Label Set is to be printed using a 600 DPI LaserJet printer or equivalent. The individual labels and PIC are to be printed using a 300 DPI Zebra thermal transfer printer.
5. The Trigger Data shown on the Label Set Image might differ from the printed label.
6. Spacing and text wrapping shown in the image may differ slightly from actual printed label. Content of printed version is accurate.
7. The label printed in Manufacturing must be at an equal or later Model Data and Template revision as shown on the image.
8. This specification is controlled by Neuromodulation Package Engineering.
9. This specification has been assessed for key design characteristics which are defined as
 - Critical KC_C - Identified characteristic is classified as Critical and impacts safety.
 - Major KC_M - Identified characteristic is classified as Major and impacts safety and/or performance.

 MEDTRONIC, MINNEAPOLIS, MINNESOTA USA	TITLE LABEL IMAGE SPEC NEURO SM III 8667-xx				
	IMAGE SPEC TEMPLATE: A86694_c				
MEDTRONIC CONFIDENTIAL This document is the property of Medtronic, and must be accounted for. Information hereon is confidential. Do not reproduce it, reveal it to unauthorized persons or send it outside Medtronic without proper authorization.	DATE 04-Jun-2021	SCALE N/A	ENGINEERING DARRYL ZURN		
	MODEL 8667	Size B	PART NUMBER B05013	REV B	SHEET 2 of 7

Medtronic, Inc., Minneapolis, MN USA
www.medtronic.com
TEMPORARY Implants Device ID Card
PUMP: SYNCHROMED™ III 8667-20 CC: 999
RESERVOIR VOLUME: 20 mL
SERIAL NO.: WWW000000W
IMPLANT DATE: _____ CATHETER: _____
ACCESSORIES: _____
Patient Name: _____
Address: _____
Physician Name: _____
Implanting center: _____
Implanting center address: _____
Phone number: _____

MR Conditional: Please refer to www.medtronic.com/mri for current, up-to-date information regarding MR Conditional labeling and instructions for this device. MR scans performed under different conditions can result in patient injury or damage to the implantable device.

Medtronic 8667-20
SYNCHROMED™ III
SN WWW000000W
Implantation date: _____
For Your Patient Files MR Conditional

Medtronic 8667-20
SYNCHROMED™ III
SN WWW000000W
Implantation date: _____
For Your Patient Files MR Conditional

Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
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Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W

Medtronic SYNCHROMED™ III REF 8667-20

1X Programmable pump
1X Package contents
1X Product documentation

999 Calibration Constant
20 mL Reservoir Volume
Targeted Drug Delivery

SN WWW000000W
YYYY-MM-DD
YYYY-MM-DD

Keep away from magnets

STERILE EO
MD
MR Conditional
www.medtronic.com/manuals
Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA
www.medtronic.com
Country of Origin: USA
M017501C001 A - M017502C001 B
PIN W000WW000W
Rx Only

Medtronic SYNCHROMED™ III REF 8667-20

1X Programmable pump
1X Package contents
1X Product documentation

999 Calibration Constant
20 mL Reservoir Volume
Targeted Drug Delivery

SN WWW000000W
YYYY-MM-DD
YYYY-MM-DD

STERILE EO
Rx Only

Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA
www.medtronic.com
M017501C001 A - M017502C001 B
PIN W000WW000W
STERILE PACK LABEL

Data P/N
M017502C001 B

Template P/N
M017501C001 A

Medtronic REF 8667-20
SYNCHROMED™ III
SN WWW000000W
YYYY-MM-DD

(01)00643169998193(17)YYMMDD(21)WWW000000W
M017501C001 A - M017502C001 B

END LABEL 1

Medtronic REF 8667-20
M017501C001 A - M017502C001 B

FCC ID: LF58667
This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

END LABEL 2

SHELF BOX LABEL

Trigger Data

```

TemplateNumber: M017501C001_A
ImageSpecID: B05013
ImageSpecRev: B
SpecPgName: B00590 Txt
ImageSpecTemplate: B05013-007
LabelSeqNo: B05013-007
OrientationLaser:
ImageSpecTitle: LABEL IMAGE SPEC NEURO SM III 8667-xx
Engineering: DARRYL GURN
PrintName: Prt_Bmp
PIN: M000WW000W
9876543210
MatrixID:
QuantityOfLabels: 1
DocumentNumber: M017502C
DocumentRevision: B
GTIN: 00643169998193
UDD: YYMMDD
ManufactureDate:
LotSerial: S
LotSerialNumber: WWW000000W
AddressCode: 007
QualificationStatus: 99
BusinessUnit: NRI
LabelPrinter: Prt_Bmp
PICPrinter: Prt_Bmp
ProductNumber: 8667
TemplateType: Neuro_Label_Templates
ReservoirVolume: 20
CalibrationConstant: 999

```

UNIVERSAL LABEL SET (DEVICES) Rev. : E

* A 1 4 6 3 2 0 0 1 * LOT NUMBER

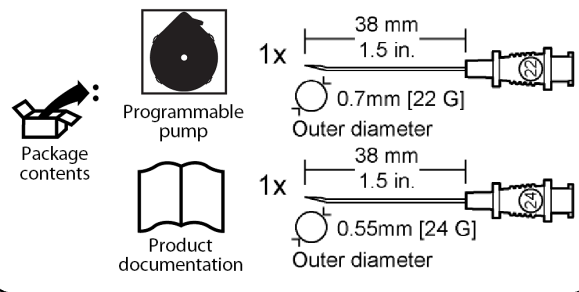
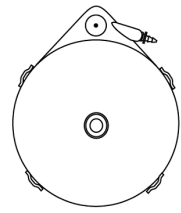
LASER

PART NUMBER	REV	SHEET
B05013	B	3 of 7



SYNCHROMED™ III

REF 8667-20



20 mL

999 Calibration Constant

SN WWW000000W

YYYY-MM-DD

YYYY-MM-DD



MR Conditional



www.medtronic.com/manuals

MD STERILE EO



Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA

Manufactured in:
Juncos, Puerto Rico, USA

Rx Only

www.medtronic.com Country of Origin: USA

PIN W000WWW00W

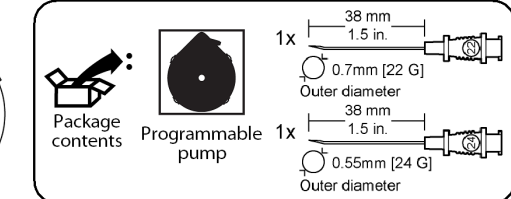
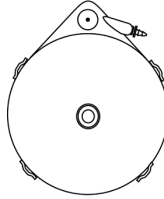
M017501C001 A - M017502C001 B

Shelf Box Label



SYNCHROMED™ III

REF 8667-20



20 mL Reservoir Volume

999 Calibration Constant

SN WWW000000W

YYYY-MM-DD

YYYY-MM-DD



STERILE EO

Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA

www.medtronic.com

Rx Only

PIN W000WWW00W

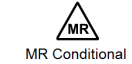
M017501C001 A - M017502C001 B

Sterile Pack Label

M017501C001 A - M017502C001 B



For Your Patient Files



Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
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Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W
Medtronic 8667-20	999 Calibration Constant	SN WWW000000W

FYR Labels

Medtronic 8667-20

SYNCHROMED™ III

YYYY-MM-D Implantation date

REF 8667-20



SN WWW000000W



For Your Patient Files



Medtronic 8667-20

SYNCHROMED™ III

SN WWW000000W Implantation date



For Your Patient Files



Medtronic 8667-20

SYNCHROMED™ III

SN WWW000000W Implantation date



For Your Patient Files



SYNCHROMED™ III

REF 8667-20

SN WWW000000W

YYYY-MM-DD



(01)00643169998193(17)YYMMDD(21)WWW000000W

M017501C001 A - M017502C001 B



End Label 1



M017501C001 A - M017502C001 B

REF 8667-20

FCC ID: LF58667

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation

End Label 2

ZEBRA

NOTE: IMAGES NOT TO SCALE

PART NUMBER	REV	SHEET
B05013	B	4 of 7

Medtronic, Inc., Minneapolis, MN USA
www.medtronic.com
TEMPORARY Implants Device ID Card
PUMP: SYNCHROMED™ III 8667-40 CC: 999
RESERVOIR VOLUME: 40 mL
SERIAL NO.: WWW000000W
IMPLANT DATE: _____ CATHETER: _____
ACCESSORIES: _____
Patient Name: _____
Address: _____
Physician Name: _____
Implanting center: _____
Implanting center address: _____
Phone number: _____

MR Conditional: Please refer to www.medtronic.com/mri for current, up-to-date information regarding MR Conditional labeling and instructions for this device. MR scans performed under different conditions can result in patient injury or damage to the implantable device.

Medtronic 8667-40
SYNCHROMED™ III

SN WWW000000W Implantation date: _____

For Your Patient Files MR Conditional

Medtronic 8667-40
SYNCHROMED™ III

SN WWW000000W Implantation date: _____

For Your Patient Files MR Conditional

Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
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Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W

Medtronic SYNCHROMED™ III REF 8667-40

1X 38 mm 1.5 in. Outer diameter
1X 0.7mm [22 G] Outer diameter
1X 38 mm 1.5 in. Outer diameter
1X 0.55mm [24 G] Outer diameter

Programmable pump
Package contents
Product documentation

40 mL Reservoir Volume
999 Calibration Constant
Targeted Drug Delivery

SN WWW000000W
YYYY-MM-DD
YYYY-MM-DD

MD STERILE EO
MR Conditional
www.medtronic.com/manuals

Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA

www.medtronic.com Country of Origin: USA
PIN W000WW000W

Manufactured in: Juncos, Puerto Rico, USA
Rx Only

Medtronic SYNCHROMED™ III REF 8667-40

1X 38 mm 1.5 in. Outer diameter
1X 0.7mm [22 G] Outer diameter
1X 38 mm 1.5 in. Outer diameter
1X 0.55mm [24 G] Outer diameter

Programmable pump
Package contents

40 mL Reservoir Volume
999 Calibration Constant
Targeted Drug Delivery

SN WWW000000W
YYYY-MM-DD
YYYY-MM-DD

STERILE EO
MR Conditional
www.medtronic.com

Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA

www.medtronic.com Country of Origin: USA
PIN W000WW000W

Manufactured in: Juncos, Puerto Rico, USA
Rx Only

Data P/N
M017502C001 B

Template P/N
M017501C001 A

STERILE PACK LABEL

Trigger Data

```

TemplateNumber: M017501C001_A
ImageSpecID: 40B05013
ImageSpecRev: B
SpecPgName: B00590 Txt
ImageSpecTemplate: A866947c
LabelSeqNo: B05013-007
OrientationLaser:
ImageSpecTitle: LABEL IMAGE SPEC NEURO SM III 8667-xx
Engineering: DARRYL ZURN
PrintName: Prt_Bmp
PIN: W000WW000W
9876543210
MatrixID:
QuantityOfLabels: 1
DocumentNumber: M017502C
DocumentRevision: B
GTIN: 00643169998193
UDD: YYYMMDD
ManufactureDate:
LotSerial: S
LotSerialNumber: WW000000W
AddressCode: 007
QualificationStatus: 99
BusinessUnit: NRI
LabelPrinter: Prt_Bmp
PICPrinter: Prt_Bmp
ProductNumber: 8667
TemplateType: Neuro_Label_Templates
ReservoirVolume: 40
CalibrationConstant: 999
  
```

UNIVERSAL LABEL SET (DEVICES) Rev. : E

* A 1 4 6 3 2 0 0 1 *

LOT NUMBER

Medtronic REF 8667-40

SYNCHROMED™ III SN WWW000000W

YYYY-MM-DD

(01)00643169998193(17)YYMMDD(21)WWW000000W

M017501C001 A - M017502C001 B

END LABEL 1

Medtronic REF 8667-40

M017501C001 A - M017502C001 B

FCC ID: LF58667

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation

END LABEL 2

SHELF BOX LABEL

Blank area for shelf box label.

Blank area for shelf box label.

LASER

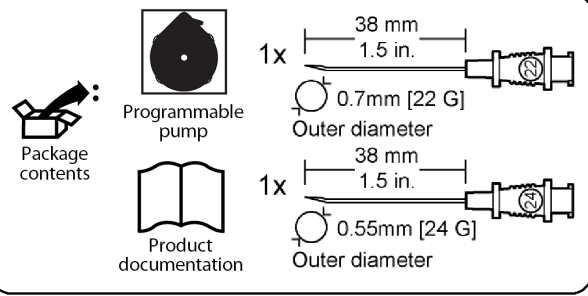
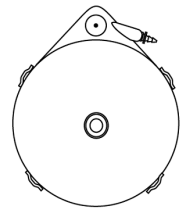
NOTE: IMAGES NOT TO SCALE

PART NUMBER	REV	SHEET
B05013	B	5 of 7



SYNCHROMED™ III

REF 8667-40



40 mL

999 Calibration Constant

SN WWW000000W
YYYY-MM-DD
YYYY-MM-DD



MR Conditional



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MD STERILE EO



Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA

Manufactured in:
Juncos, Puerto Rico, USA

Rx Only

www.medtronic.com Country of Origin: USA

PIN W000WWW00W

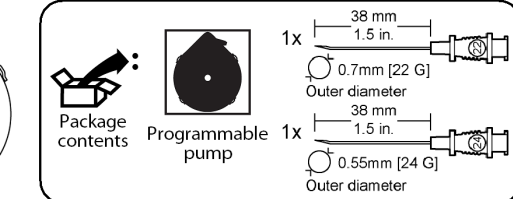
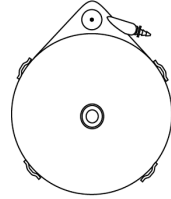
M017501C001 A - M017502C001 B

Shelf Box Label



SYNCHROMED™ III

REF 8667-40



40 mL Reservoir Volume

999 Calibration Constant

SN WWW000000W
YYYY-MM-DD
YYYY-MM-DD



STERILE EO

Medtronic, Inc., 710 Medtronic Parkway
Minneapolis, MN 55432 USA

www.medtronic.com

Rx Only

PIN W000WWW00W

M017501C001 A - M017502C001 B

Sterile Pack Label

M017501C001 A - M017502C001 B



For Your Patient Files



MR Conditional

Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
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Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W
Medtronic 8667-40	999 Calibration Constant	SN WWW000000W

FYR Labels

Medtronic 8667-40

SYNCHROMED™ III

YYYY-MM-D

REF 8667-40

Implantation date



SN WWW000000W



For Your Patient Files

MR Conditional

Medtronic 8667-40

SYNCHROMED™ III

SN WWW000000W

Implantation date



For Your Patient Files

MR Conditional

Medtronic 8667-40

SYNCHROMED™ III

SN WWW000000W

Implantation date



For Your Patient Files

MR Conditional



SYNCHROMED™ III

REF 8667-40

SN WWW000000W

YYYY-MM-DD



(01)00643169998193(17)YYMMDD(21)WWW000000W

M017501C001 A - M017502C001 B



End Label 1



M017501C001 A - M017502C001 B

REF 8667-40

FCC ID: LF58667

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation

End Label 2

ZEBRA

NOTE: IMAGES NOT TO SCALE

PART NUMBER	REV	SHEET
B05013	B	6 of 7

en Pump / **fr** Pompe / **de** Pumpe /
es Bomba / **nl** Pomp / **it** Pompa / **no** Pumpe /
sv Pump / **el** Αντλία / **da** Pumpe / **pt** Bomba /
pt-BR Bomba / **hu** Pumpa / **pl** Pompa / **cs** Pumpa /
sk Pumpa / **tr** Pompa / **ro** Pompă / **bg** Помпа / **is** Dæla /
sl Črpalka / **hr** Pumpa / **et** Pump / **lt** Pompa / **lv** Sūkņis /
fi Pumppu / **uk** Помпа / **ru** Насос /
mk Пумпа / **vi** Máy bơm / **sr** Pumpa

MD 8667 SYNCHROMED™ III

UDI



UDI-DI: (01)00643169998193



SN WWW000000W

MR Conditional



Medtronic, Inc., 710 Medtronic Parkway
 Minneapolis, MN 55432 USA, www.medtronic.com

M017501C001 A
 M017502C001 B

PIC

NOTE: IMAGES NOT TO SCALE

PART NUMBER	REV	SHEET
B05013	B	7 of 7