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NOVARTIS

NDC 0078-0180-01
Unit Dose Package

50 mcg/mL; Each mL contains:
octreotide (as acetate) . . . . . 50 mcg
Inactive ingredients
lactic acid, USP . . . . . 3.4 mg
mannitol, USP . . . . . 45 mg
sodium bicarbonate, USP . . . . . qs to pH 4.2 ± 0.3
water for injection, USP . . . . . qs to 1 mL

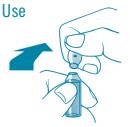
See package insert for dosage and administration information.

Storage: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light. At room temperature, (20°C to 30°C or 70°F to 86°F), Sandostatin is stable for 14 days if protected from light.

N 3 0078-0180-01 6

Instructions for Use

One-point-cut ampul with cut below colored point



To open, hold as shown with thumb above point and snap off backwards

Manufactured by:
Novartis Pharma Stein AG
Schaffhauserstrasse
CH-4332 Stein, Switzerland

Distributed by:
Novartis Pharmaceuticals Corporation
East Hanover, New Jersey 07936

1256727

NDC 0078-0180-01
Unit Dose Package
10 Ampuls/1 mL size

NOVARTIS
Sandostatin®
octreotide acetate INJECTION
50 mcg/mL

120 x 105 x 20.5
1256727
US

EXP.
LOT

NOVARTIS

NDC 0078-0180-01
Unit Dose Package

Sandostatin®

octreotide acetate
INJECTION

Rx only

50 mcg/mL
For Subcutaneous Injection

10 Ampuls/1 mL size

1256727

Please open here Please open here

NOVARTIS

NDC 0078-0181-01
Unit Dose Package

100 mcg/mL; Each mL contains:
octreotide (as acetate) . . . . . 100 mcg
Inactive ingredients
lactic acid, USP . . . . . 3.4 mg
mannitol, USP . . . . . 45 mg
sodium bicarbonate, USP . . . . . qs to pH 4.2 ± 0.3
water for injection, USP . . . . . qs to 1 mL

See package insert for dosage and administration information.

Storage: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light. At room temperature, (20°C to 30°C or 70°F to 86°F), Sandostatin is stable for 14 days if protected from light.

N 3 0078-0181-01 3

Instructions for Use

One-point-cut ampul with cut below colored point

To open, hold as shown with thumb above point and snap off backwards

Manufactured by:
Novartis Pharma Stein AG
Schaffhauserstrasse
CH-4332 Stein, Switzerland

Distributed by:
Novartis Pharmaceuticals Corporation
East Hanover, New Jersey 07936

1256728

NDC 0078-0181-01
Unit Dose Package
10 Ampuls/1 mL size

NOVARTIS
Sandostatin®
octreotide acetate INJECTION
100 mcg/mL

120 x 105 x 20.5
1256728
US

NOVARTIS
Sandostatin®
octreotide acetate INJECTION
100 mcg/mL

NDC 0078-0181-01
Unit Dose Package
10 Ampuls/1 mL size

EXP.
LOT

NOVARTIS

NDC 0078-0181-01
Unit Dose Package

Sandostatin®
octreotide acetate
INJECTION

Rx only

100 mcg/mL
For Subcutaneous Injection

10 Ampuls/1 mL size

1256728

Please open here Please open here

NOVARTIS

NDC 0078-0182-01
Unit Dose Package

500 mcg/mL; Each mL contains:
octreotide (as acetate) . . . . . 500 mcg
Inactive ingredients
lactic acid, USP . . . . . 3.4 mg
mannitol, USP . . . . . 45 mg
sodium bicarbonate, USP . . . . . qs to pH 4.2 ± 0.3
water for injection, USP . . . . . qs to 1 mL

See package insert for dosage and administration information.

Storage: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light. At room temperature, (20°C to 30°C or 70°F to 86°F), Sandostatin is stable for 14 days if protected from light.

N 3 0078-0182-01 0

Instructions for Use

One-point-cut ampul with cut below colored point

To open, hold as shown with thumb above point and snap off backwards

Manufactured by:
Novartis Pharma Stein AG
Schaffhauserstrasse
CH-4332 Stein, Switzerland

Distributed by:
Novartis Pharmaceuticals Corporation
East Hanover, New Jersey 07936

1256731

NDC 0078-0182-01 Unit Dose Package 10 Ampuls/1 mL size

NOVARTIS Sandostatin® octreotide acetate INJECTION 500 mcg/mL

NOVARTIS Sandostatin® octreotide acetate INJECTION 500 mcg/mL

NDC 0078-0182-01 Unit Dose Package 10 Ampuls/1 mL size

120 x 105 x 20.5
1256731
US

EXP.
LOT

NOVARTIS

NDC 0078-0182-01
Unit Dose Package

Sandostatin®

octreotide acetate

INJECTION

Rx only

500 mcg/mL

For Subcutaneous Injection

10 Ampuls/1 mL size

1256731

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g information, please visit https://www.

Sandostatin®
octreotide acetate
INJECTION
1,000 mcg/5 mL (200 mcg/mL)
Total Volume 5 mL Multi-Dose Vial
Each mL of aqueous solution contains: **octreotide (as acetate) 200 mcg**
FOR SUBCUTANEOUS INJECTION
STORAGE: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light.
After initial use, d scard w thin 14 days.
Mfd. by Novartis Pharma Stein AG
NDC 0078-0183-25
Rx only
© Novartis
1256734B US 1256734A
EXP./LOT

Sandostatin®
octreotide acetate
INJECTION
1,000 mcg/5 mL (200 mcg/mL)
Total Volume 5 mL Multi-Dose Vial
Each mL of aqueous solution contains: **octreotide (as acetate) 200 mcg**
FOR SUBCUTANEOUS INJECTION
STORAGE: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light.
After initial use, d scard w thin 14 days.
Mfd. by Novartis Pharma Stein AG
NDC 0078-0183-25
Rx only
© Novartis
1256734B US 256734A
EXP./LOT

Sandostatin®
octreotide acetate
INJECTION
1,000 mcg/5 mL (200 mcg/mL)
Total Volume 5 mL Multi-Dose Vial
Each mL of aqueous solution contains: **octreotide (as acetate) 200 mcg**
FOR SUBCUTANEOUS INJECTION
STORAGE: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light.
After initial use, d scard w thin 14 days.
Mfd. by Novartis Pharma Stein AG
NDC 0078-0183-25
Rx only
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1256734B US 256734A
EXP./LOT

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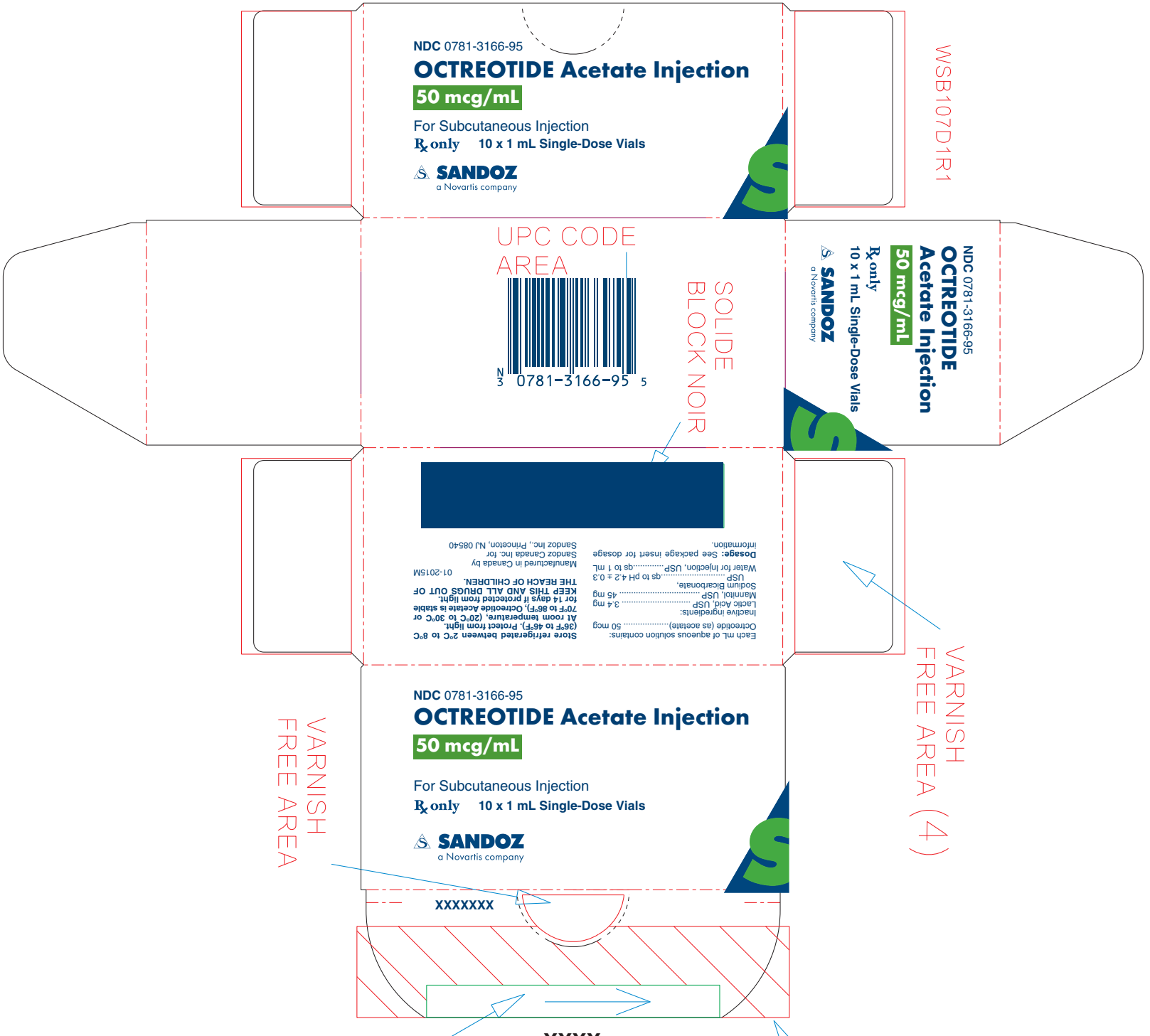
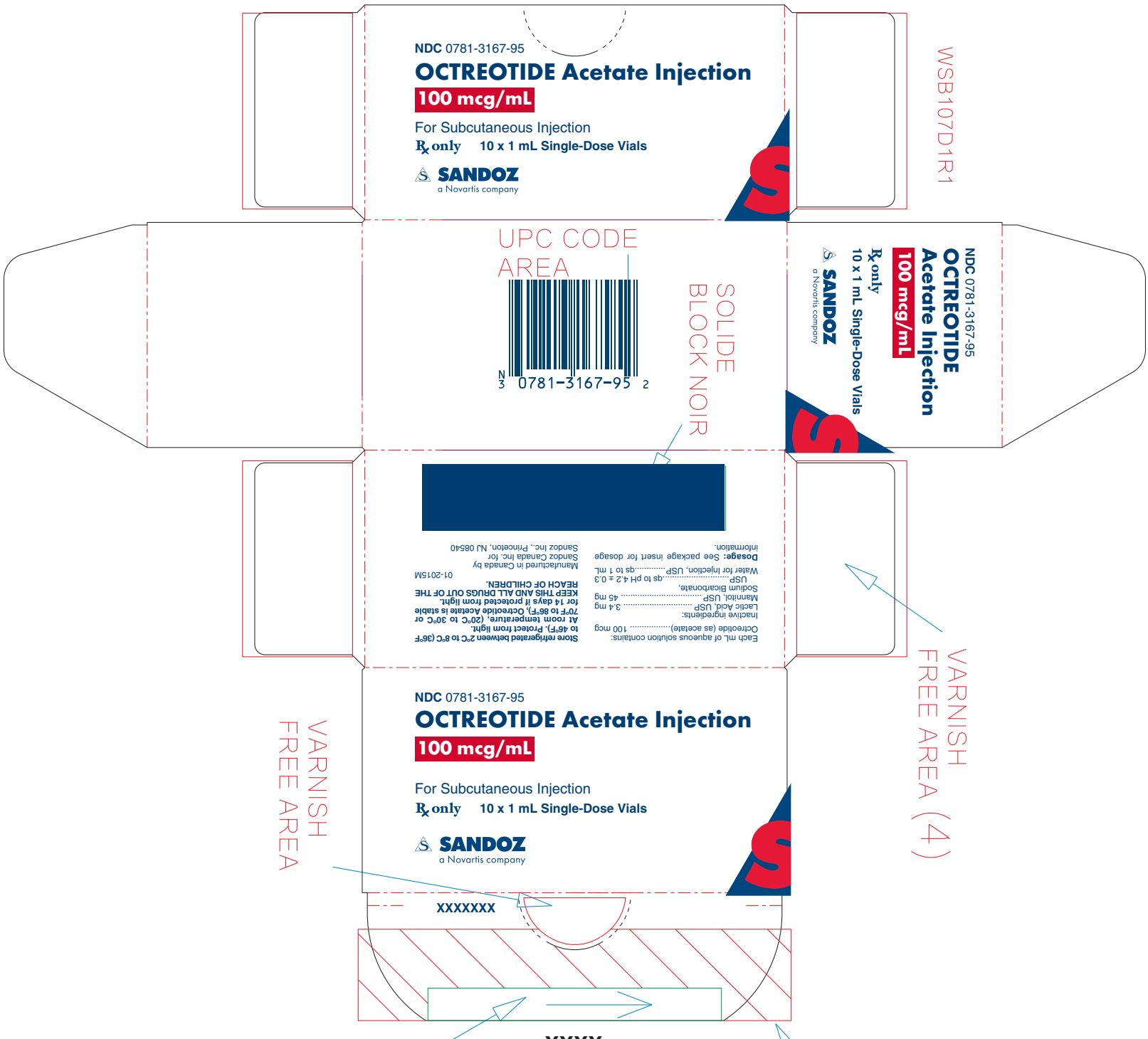
g information, please visit https://www.

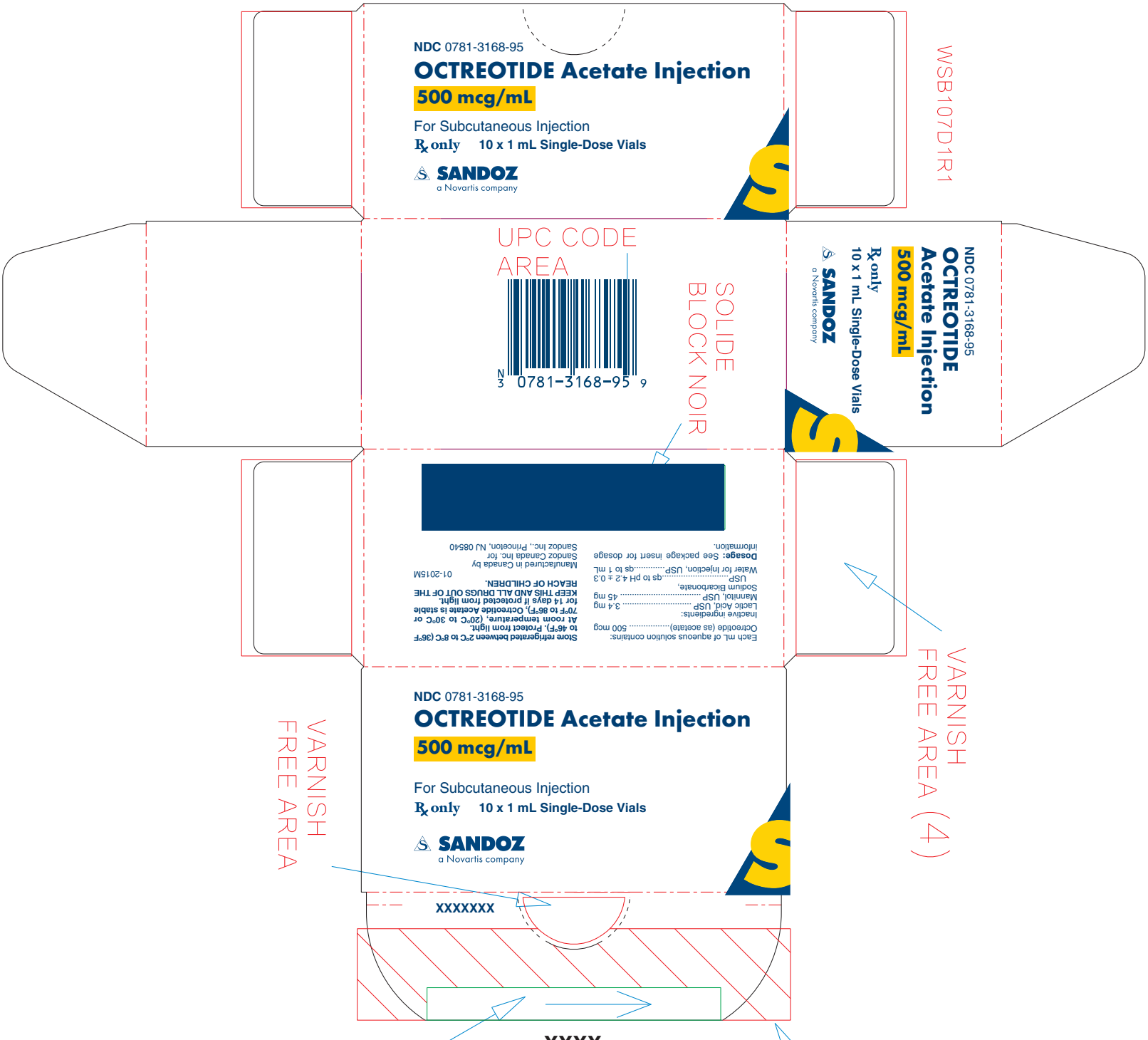
Sandostatin®
octreotide acetate
INJECTION
5,000 mcg/5 mL (1,000 mcg/mL)
Total Volume 5 mL Multi-Dose Vial
Each mL of aqueous solution contains: **octreotide (as acetate) 1,000 mcg**
FOR SUBCUTANEOUS INJECTION
STORAGE: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light.
After initial use, discard with n 14 days.
Mfd. by Novartis Pharma Stein AG
NDC 0078-0184-25
Rx only
©Novartis
1256740A **US** 1256740B 1256740C
EXP./LOT

Sandostatin®
octreotide acetate
INJECTION
5,000 mcg/5 mL (1,000 mcg/mL)
Total Volume 5 mL Multi-Dose Vial
Each mL of aqueous so ution contains: **octreotide (as acetate) 1,000 mcg**
FOR SUBCUTANEOUS INJECTION
STORAGE: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from ight.
After nitial use, discard within 14 days.
Mfd. by Novartis Pharma Stein AG
NDC 0078-0184-25
Rx only
©Novartis
256740A **US** 1256740B 1256740C
EXP./LOT

Sandostatin®
octreotide acetate
INJECTION
5,000 mcg/5 mL (1,000 mcg/mL)
Total Volume 5 mL Multi-Dose Vial
Each mL of aqueous so ution contains: **octreotide (as acetate) 1,000 mcg**
FOR SUBCUTANEOUS INJECTION
STORAGE: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from ight.
After nitial use, discard within 14 days.
Mfd. by Novartis Pharma Stein AG
NDC 0078-0184-25
Rx only
©Novartis
256740A **US** 1256740B 1256740C
EXP./LOT







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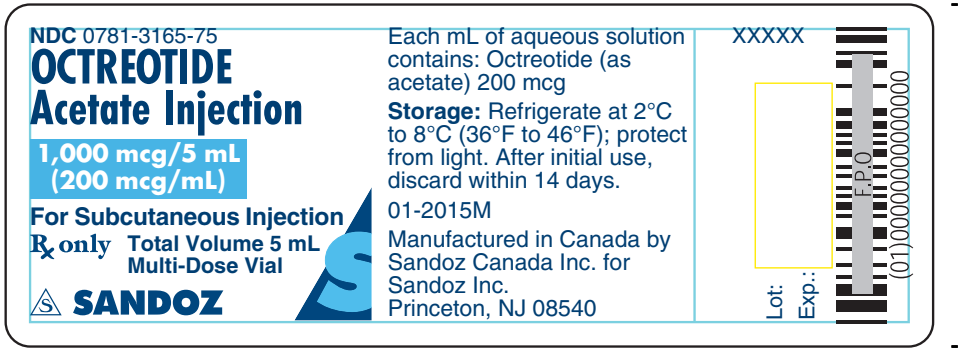

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(b) (4)

NDC 0781-3165-75
OCTREOTIDE Acetate Injection
1,000 mcg/5 mL (200 mcg/mL)
For Subcutaneous Injection
Rx only
Total Volume 5 mL Multi-Dose Vial
SANDOZ a Novartis company

Each mL of aqueous solution contains:
Octreotide (as acetate) .....200 mcg

Inactive Ingredients:
Lactic Acid, USP ..... 3.4 mg
Mannitol, USP ..... 45 mg
Phenol, USP .....5.0 mg
Sodium Bicarbonate, USP .....qs to pH 4.2 ± 0.3
Water for Injection, USP .....qs to 1 mL

DOSAGE: See package insert for dosage information.

Manufactured in Canada by
Sandoz Canada Inc. for
Sandoz Inc.
Princeton, NJ 08540

01-2015M

SANDOZ a Novartis company

Storage: Refrigerate at 2°C to 8°C (36°F to 46°F); protect from light. After initial use, discard within 14 days.

At room temperature, (20°C to 30°C or 70°F to 86°F), Octreotide Acetate is stable for 14 days if protected from light.

KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

NDC 0781-3165-75
OCTREOTIDE Acetate Injection
1,000 mcg/5 mL (200 mcg/mL)
For Subcutaneous Injection
Rx only
Total Volume 5 mL Multi-Dose Vial
SANDOZ a Novartis company

UPC CODE AREA

128 CODE AREA

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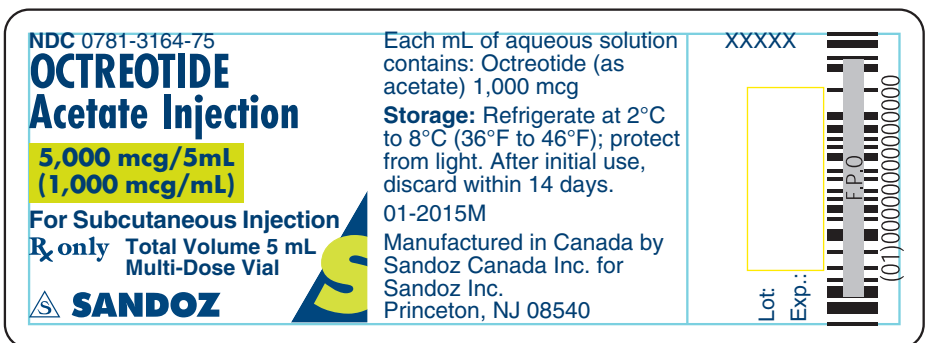

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