

Please open here  Please open here

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

 **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.



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

Product of Ireland.

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

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

©Novartis

 **Sandostatin®**
octreotide acetate INJECTION
50 mcg/mL

120 x 105 x 20.5
2235566
US

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

 **Sandostatin®**
octreotide acetate INJECTION
50 mcg/mL

BR/LOT
GTIN: 00300780180016

 **NOVARTIS**

NDC 0078-0180-01
Unit Dose Package

 **Sandostatin®**
octreotide acetate
INJECTION

Rx only

50 mcg/mL

For Subcutaneous or Intravenous Use 10 Ampuls/1 mL size

2235566

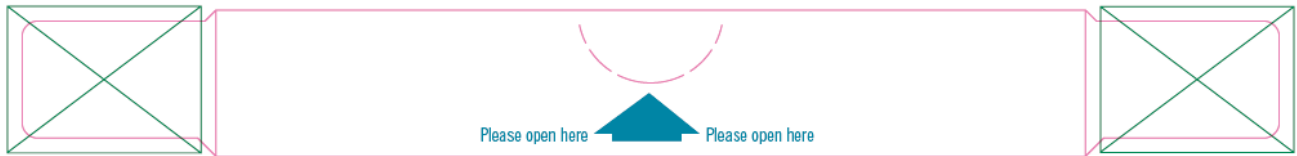


This label may not be the latest approved by FDA.
For the most current labeling information, please visit <https://www.fda.gov>.

NDC 0074-0180-61
Sandostatin[®]
octreotide acetate
Injection (subcutaneous
or intravenous)
1 mL contains
50 mcg octreotide
(as acetate) Rx only
Mfg. by Novartis Pharma
Stein AG ©Novartis 2235565 US

EXPIRES

2235565



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

 **NOVARTIS**

NDC 0078-0181-01
Unit Dose Package

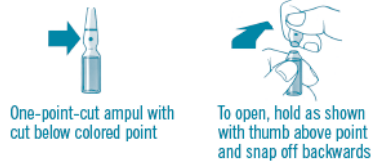
100 mcg/mL; Each mL contains:
octreotide (as acetate) 100 mcg
Inactive ingredients
lactic acid, USP 3.4 mg
mannitol, USP 45 mg
sodium bicarbonate, USP qs to pH 4.2 ± 0.3
water for injection, USP qs to 1 mL

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.**



Instructions for Use



Product of Ireland.

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

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

©Novartis

 **Sandostatin**[®]
octreotide acetate INJECTION
100 mcg/mL

120 x 105 x 205
2235568
US

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

 **Sandostatin**[®]
octreotide acetate INJECTION
100 mcg/mL



 **NOVARTIS**

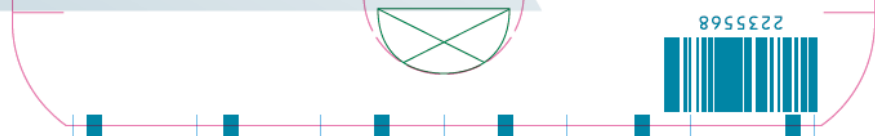
NDC 0078-0181-01
Unit Dose Package

 **Sandostatin**[®]
octreotide acetate
INJECTION

Rx only

100 mcg/mL

For Subcutaneous or Intravenous Use 10 Ampuls/1 mL size



Label may not be the latest approved
information, please visit <https://www>

mc-007-0181-01
Sandostatin[®]
octreotide acetate
injection (subcutaneous or
intravenous)
1 mL contains:
200 mcg octreotide
(as acetate) Rx only
Mfg. by: Novartis Pharma
Stein AG ©Novartis 2235567 US

EXP/LOT

2235567

Please open here  Please open here

NDC 0078-0182-01
Unit Dose Package

10 Ampuls/1 mL size

 **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.



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

Product of Ireland.

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

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

©Novartis

 **Sandostatin®**
 octreotide acetate INJECTION
 500 mcg/mL

 **Sandostatin®**
 octreotide acetate INJECTION
 500 mcg/mL

120 x 105 x 20.5
 2235570
 US

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

EP/LDT
 GTIN: 00300780182010

 **NOVARTIS**

NDC 0078-0182-01
Unit Dose Package

 **Sandostatin®**
octreotide acetate
 INJECTION

Rx only

500 mcg/mL

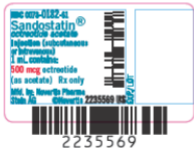
For Subcutaneous or Intravenous Use

10 Ampuls/1 mL size

2235570



I may not be the latest approved
information, please visit <https://www.fda.gov>



2235569-0182-01
Sandostatin®
octreotide acetate
Injection (subcutaneous
or intravenous)
1 mL contains:
500 mcg octreotide
(as acetate) Rx only
Mfg. by: Novartis Pharma
St. Gallen, Switzerland 2235569-0182-01

LOT
6067107

2235569

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/s/

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