

**ISOTONIC SODIUM CHLORIDE- biofleks isotonic sodium chloride
sterile injection
HAVER FARMA ILAC ANONIM SIRKETI**

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ADMINISTERED INTRAVENOUSLY ASEPTICALLY. Do not use if the solution is not clear or the bag is damaged. Please read the package leaflet. Consult your doctor right away if you have any unexpected effect. It is disposable. Partially used solutions should not be stored. Keep out of reach and sight of the children and store in the original package. Store at room temperature below 25°C. Sold with prescription only.

biofleks
100 mL
0.9 % ISOTONIC
SODIUM CHLORIDE SOLUTION
STERILE

Per 100 mL
Sodium Chloride
Water for Injection

Electrolyte Concentrations (mEq/L)

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Phone: (201) 676-3232 / E-Mail: info@vintradingllc.com Address: 203 Lemoine Ave #118, Fort Lee, NJ 07024

OSEL İlaç San. ve Tic. A.Ş.
Beykoz / İstanbul / TURKEY

100mL Bag

60 mm

biofleks® **100 mL**

**0.9 % ISOTONIC
SODIUM CHLORIDE SOLUTION
STERILE**

25 **25**

Per 100 mL
Sodium Chloride 0.9 g
Water for Injection q.s.

Electrolyte Concentrations (mEq/L)
Sodium 154 Chloride 154

50 **50** **80 mm**

75 **75**

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OSEL İlaç San. ve Tic. A.Ş.
Beykoz / İstanbul / TURKEY

 Batch No: Exp. Date: 

Amerika Product Code: PVC - 254163/1

Graphic Designer: Bahar ATABAY

Regulatory Affairs Department	Regulatory Affairs Department	Production	Quality Control Laboratory	Quality Control Laboratory
Date:	Date:	Date:	Date:	Date:

250mL Bag

biofleks®

250 mL

**0.9 % ISOTONIC
SODIUM CHLORIDE SOLUTION
STERILE**

50

Per 100 mL

Sodium Chloride
Water for Injection

0.9 g
q.s.

50

100

Electrolyte Concentrations (mEq/L) Sodium 154 Chloride 154

100

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150

150

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200

200

OSEL İlaç San. ve Tic. A.Ş.
Beykoz / İstanbul / TURKEY



Batch No: Exp. Date



80mm

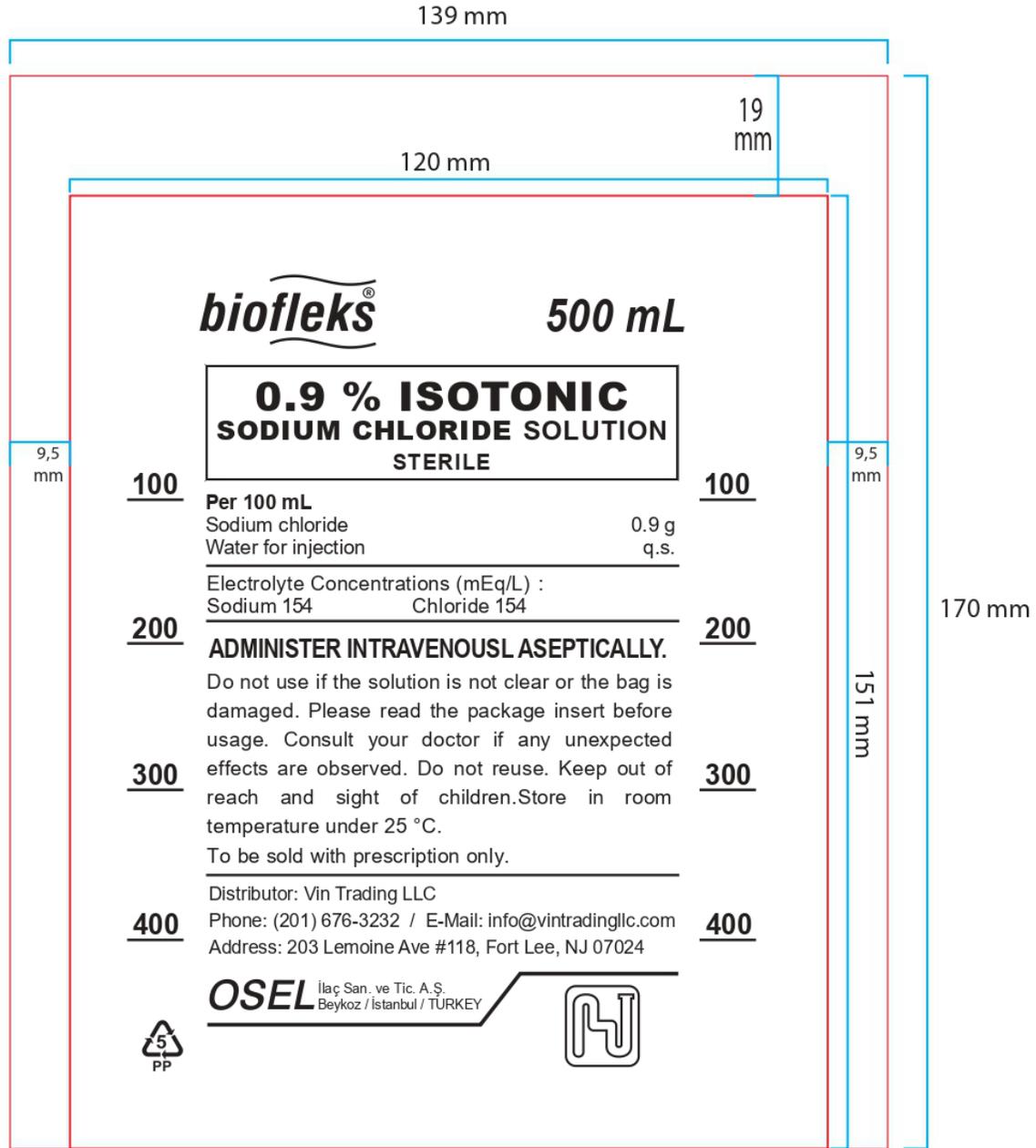
100mm

Amerika Product Code: 254164/1

Graphic Designer: Bahar ATABAY

Regulatory Affairs Department	Regulatory Affairs Department	Production	Quality Control Laboratory	Quality Control Laboratory
Date:	Date:	Date:	Date:	Date:

500 mL Bag



Amerika Product Code: PP - XXXXXX/0

Graphic Designer: Bahar ATABAY

Regulatory Affairs Department	Regulatory Affairs Department	Production	Quality Control Laboratory	Quality Control Laboratory
Date:	Date:	Date:	Date:	Date:

1000mL Bag

100mm

220mm

biofleks[®]

1000 mL

0.9 % ISOTONIC SODIUM CHLORIDE SOLUTION STERILE
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100

100

Per 100 mL

Sodium Chloride 0.9 g

200

Water for Injection q.s. **200**

Electrolyte Concentrations (mEq/L)

300

Sodium 154 Chloride 154 **300**

**ADMINISTERED INTRAVENOUSLY
ASEPTICALLY.**

400

Do not use if the solution is not clear or the bag is damaged.

400

Please read the package leaflet.

500

Consult your doctor right away if you have any unexpected effect. It

500

is disposable. Partially used solutions

600

should not be stored. Keep out of reach and sight of the children and

600

store in the original package. Store

700

at room temperature below 25°C.

700

Sold with prescription only.

800

Distributor: Vin Trading LLC

Phone: (201) 676-3232

E-Mail: info@vintredinalla.com

800

E-Mail: info@vitradingilc.com

Address: 203 Lemoine Ave #118,
Fort Lee, NJ 07024

900

OSEL İlaç San. ve Tic. A.Ş.
Beykoz / İstanbul / TURKEY

900



Batch No:

Exp. Date:



Amerika Product Code: 254166/1

Graphic Designer: Bahar ATABAY

Regulatory Affairs Department	Regulatory Affairs Department	Production	Quality Control Laboratory	Quality Control Laboratory
Date:	Date:	Date:	Date:	Date:

ISOTONIC SODIUM CHLORIDE

biofleks isotonic sodium chloride sterile injection

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:85184-0001
Route of Administration	INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS, IRRIGATION		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	0.9 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:85184-0001-1	100 mL in 1 BAG; Type 0: Not a Combination Product	01/01/2025	
2	NDC:85184-0001-2	250 mL in 1 BAG; Type 0: Not a Combination Product	01/01/2025	
3	NDC:85184-0001-3	500 mL in 1 BAG; Type 0: Not a Combination Product	01/01/2025	
4	NDC:85184-0001-4	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2025	

Labeler - HAVER FARMA ILAC ANONIM SIRKETI (365593526)

Registrant - HAVER FARMA ILAC ANONIM SIRKETI (365593526)

Establishment

Name	Address	ID/FEI	Business Operations
HAVER FARMA ILAC ANONIM SIRKETI		365593526	manufacture(85184-0001)

Revised: 5/2025

HAVER FARMA ILAC ANONIM SIRKETI