EPINEPHRINE PROFESSIONAL- epinephrine and isopropyl alcohol
Focus Health Group

Epinephrine Professional Convenience Kit

For treating anaphylaxis:

Adults and Children 30 kg (66 lbs) or more: 0.3 to 0.5 mg (0.3 to 0.5 ml) intramuscularly or subcutaneously into anterolateral aspect of the thigh every 5 to 10 minutes as necessary.

Children 30 kg (66 lbs) or less: 0.01 mg/kg (0.01 ml/kg), up to 0.3 mg (0.3 ml), intramuscularly or subcutaneously into anterolateral aspect of the thigh every 5 to 10 minutes as necessary.

Principal Display Panel

EPINEPHRINE PROFESSIONAL

epinephrine and isopropyl alcohol kit

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:24357-011

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24357-011-13	1 in 1 CARTON; Type 0: Not a Combination Product	07/01/2019	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 VIAL	1 mL
Part 2	4 PACKET	4 mL

Part 1 of 2

ADRENALIN

epinephrine injection

Product Information	
Item Code (Source)	NDC:42023-159
Route of Administration	SUBCUTANEOUS, INTRAMUSCULAR

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
EPINEPHRINE (UNII: YKH834O4BH) (EPINEPHRINE - UNII:YKH834O4BH)	EPINEPHRINE	1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
TARTARIC ACID (UNII: W4888I119H)	2.25 mg in 1 mL
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	1 mg in 1 mL
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	0.2 mg in 1 mL
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	0.457 mg in 1 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	7.3 mg in 1 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42023-159-25	1 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA204200	07/01/2013	

Part 2 of 2

MCKESSON ALCOHOL PREP PAD

isopropyl alcohol swab

Product Information	
Item Code (Source)	NDC:68599-5804
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68599-5804-1	1 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/09/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA204200	07/01/2019	

Labeler - Focus Health Group (826939949)

Establishment

Name	Address	ID/FEI	Business Operations
119548429		119548429	manufacture(42023-159)

Establishment

Name	Address	ID/FEI	Business Operations
Focus Health Group		826939949	label(24357-011)

Revised: 2/2026 Focus Health Group