

ZINC SULFATE- zinc sulfate injection, solution
Zydus Lifesciences Limited

ZINC SULFATE INJECTION for intravenous use

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1848-1

Zinc Sulfate Injection, USP

10 mg/10 mL (1 mg/mL) of zinc

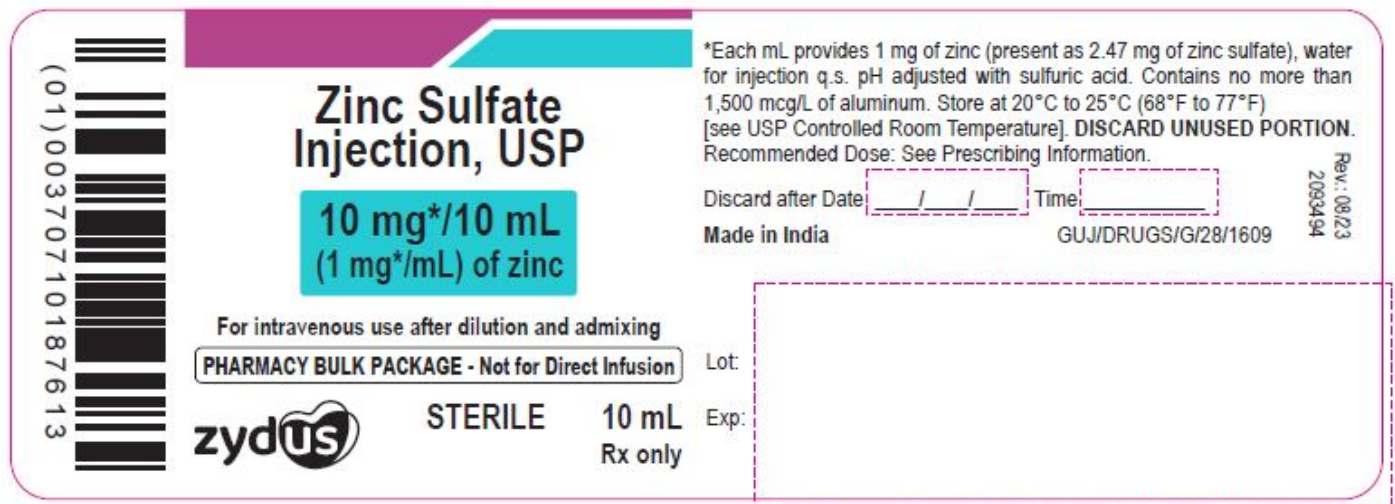
For intravenous use after dilution and admixing

PHARMACY BULK PACKAGE-Not for Direct Infusion

STERILE

10 mL

Rx only



NDC 70771-1848-7

Zinc Sulfate Injection, USP

10 mg/10 mL (1 mg/mL) of zinc

For intravenous use after dilution and admixing

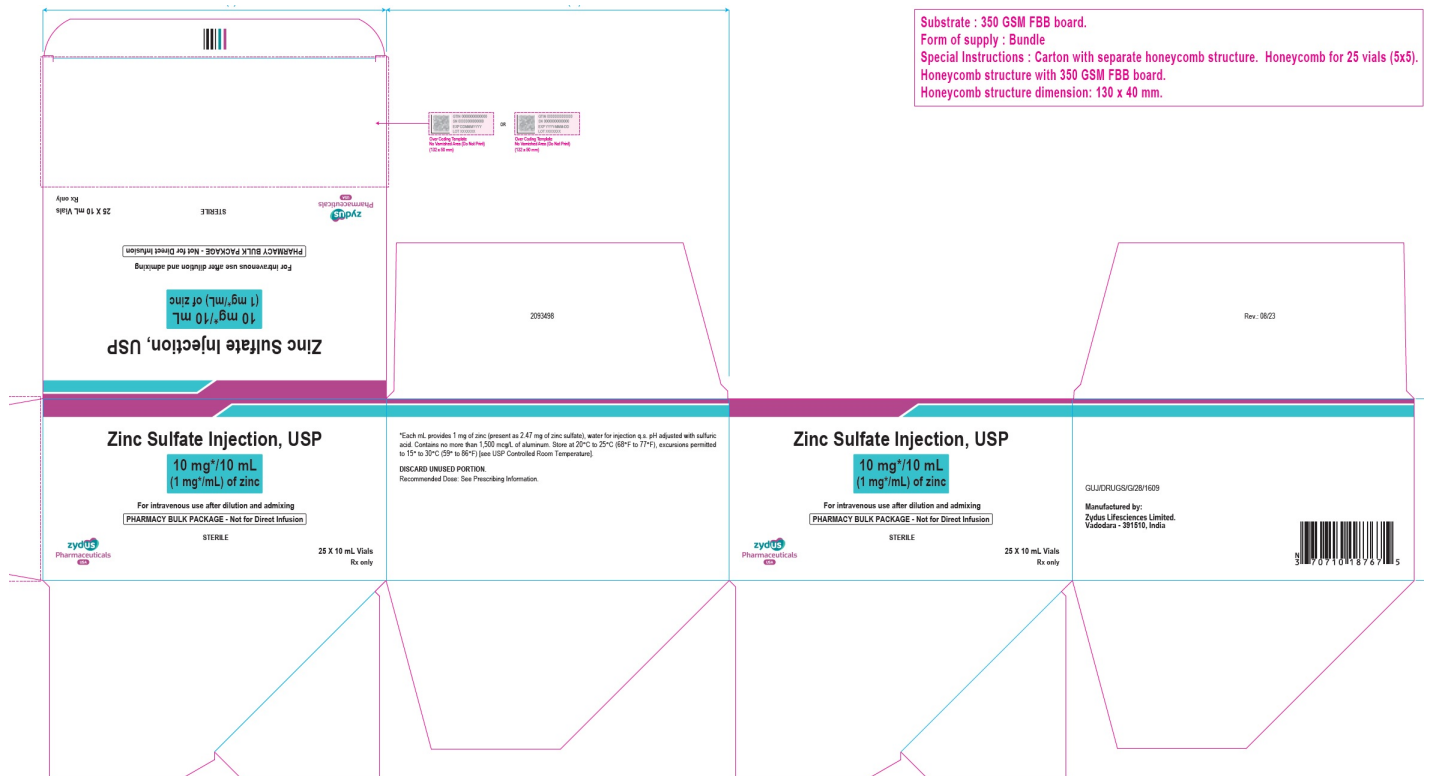
PHARMACY BULK PACKAGE-Not for Direct Infusion

25 x 10 mL Vials

STERILE

10 mL

Rx only



NDC 70771-1849-1

Zinc Sulfate Injection, USP

30 mg/10 mL (3 mg/mL) of zinc

For intravenous use after dilution and admixing

PHARMACY BULK PACKAGE-Not for Direct Infusion

STERILE

10 mL

Rx only

 (01)00370710187712	Zinc Sulfate Injection, USP 30 mg*/10 mL (3 mg*/mL) of zinc		*Each mL provides 3 mg of zinc (present as 7.41 mg of zinc sulfate), water for injection q.s. pH adjusted with sulfuric acid. Contains no more than 2,500 mcg/L of aluminum. Store at 20°C to 25°C (68°F to 77°F) [see USP Controlled Room Temperature]. DISCARD UNUSED PORTION. Recommended Dose: See Prescribing Information.		Rev.: 08/23 2093495
	For intravenous use after dilution and admixing PHARMACY BULK PACKAGE - Not for Direct Infusion		Discard after Date: / / Time:		
	zydus		STERILE		
	10 mL Rx only		Lot: Exp:		

NDC 70771-1849-7

Zinc Sulfate Injection, USP

30 mg/10 mL (3 mg/mL) of zinc

For intravenous use after dilution and admixing

PHARMACY BULK PACKAGE-Not for Direct Infusion

25 X 10 mL Vials

STERILE

10 mL

Rx only

NDC 70771-1850-7

Zinc Sulfate Injection, USP

25 mg/5 mL (5 mg/mL) of zinc

For intravenous use after dilution and admixing

PHARMACY BULK PACKAGE-Not for Direct Infusion

25 X 5 mL Vials

STERILE

5 mL

Rx only

 (01)00370710187811	Zinc Sulfate Injection, USP	*Each mL provides 5 mg of zinc (present as 12.34 mg of zinc sulfate), water for injection q.s. pH adjusted with sulfuric acid. Contains no more than 2,500 mcg/L of aluminum. Store at 20°C to 25°C (68°F to 77°F) [see USP Controlled Room Temperature]. DISCARD UNUSED PORTION. Recommended Dose: See Prescribing Information. Discard after Date / / Time: Made in India
	25 mg*/5 mL (5 mg*/mL) of zinc For intravenous use after dilution and admixing PHARMACY BULK PACKAGE - Not for Direct Infusion zydus STERILE 5 mL Rx only	
		Rev: 08/23 203493 GUJ/DRUGS/G/28/1609 Lot: Exp:

ZINC SULFATE

zinc sulfate injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1848
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SULFURIC ACID (UNII: O40UQP6WCF)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1848-7	25 in 1 CARTON	12/07/2023	
1	NDC:70771-1848-1	10 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
ANDA	ANDA217074	12/07/2023	

ZINC SULFATE			
zinc sulfate injection, solution			

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1849
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SULFURIC ACID (UNII: O40UQP6WCF)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1849-7	25 in 1 CARTON	12/07/2023	
1	NDC:70771-1849-1	10 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

ANDA	ANDA217074	12/07/2023	
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ZINC SULFATE

zinc sulfate injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1850
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SULFURIC ACID (UNII: O40UQP6WCF)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1850-7	25 in 1 CARTON	12/07/2023	
1	NDC:70771-1850-1	5 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
ANDA	ANDA217074	12/07/2023	

Labeler - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		873671928	MANUFACTURE(70771-1848, 70771-1849, 70771-1850) , ANALYSIS(70771-1848, 70771-1849, 70771-1850)