

specific B₁₂ binding proteins, transcobalamin I and II to the various tissues. The liver is the main organ for vitamin B₁₂ storage.

Within 48 hours after injection of 100 or 1,000 mcg of vitamin B₁₂, 50 to 98% of the injected dose may appear in the urine. The major portion is excreted within the first eight hours. Intravenous administration results in even more rapid excretion with little opportunity for liver storage.

Gastrointestinal absorption of vitamin B₁₂ depends on the presence of sufficient intrinsic factor and calcium ions. Intrinsic factor deficiency causes pernicious anemia, which may be associated with subacute combined degeneration of the spinal cord. Prompt parenteral administration of vitamin B₁₂ prevents progression of neurologic damage.

The average diet supplies about 5 to 15 mcg/day of vitamin B₁₂ in a protein-bound form that is available for absorption after normal digestion. Vitamin B₁₂ is not present in foods of plant origin, but is abundant in foods of animal origin. In people with normal absorption, deficiencies have been reported only in strict vegetarians who consume no products of animal origin (including no milk products or eggs).

Vitamin B₁₂ is bound to intrinsic factor during transit through the stomach; separation occurs in the terminal ileum in the presence of calcium, and vitamin B₁₂ enters the mucosal cell for absorption. It is then transported by the transcobalamin binding proteins. A small amount (approximately 1% of the total amount ingested) is absorbed by simple diffusion, but this mechanism is adequate only with very large doses. Oral absorption is considered too undependable to rely on in patients with pernicious anemia or other conditions resulting in malabsorption of vitamin B₁₂.

Cyanocobalamin is the most widely used form of vitamin B₁₂, and has hematopoietic activity apparently identical to that of the antianemia factor in purified liver extract. Hydroxycobalamin is equally as effective as cyanocobalamin, and they share the cobalamin molecular structure.

INDICATIONS AND USAGE

Cyanocobalamin is indicated for vitamin B₁₂ deficiencies due to malabsorption which may be associated with the following conditions:

- Addisonian (pernicious) anemia
- Gastrointestinal pathology, dysfunction, or surgery, including gluten enteropathy or sprue, small bowel bacterial overgrowth, total or partial gastrectomy
- Fish tapeworm infestation
- Malignancy of pancreas or bowel
- Folic acid deficiency

It may be possible to treat the underlying disease by surgical correction of anatomic lesions leading to small bowel bacterial overgrowth, expulsion of fish tapeworm, discontinuation of drugs leading to vitamin malabsorption (see **Drug/Laboratory Test Interactions**), use of a gluten-free diet in nontropical sprue, or administration of antibiotics in tropical sprue. Such measures remove the need for long-term administration of cyanocobalamin.

Requirements of vitamin B₁₂ in excess of normal (due to pregnancy, thyrotoxicosis, hemolytic anemia, hemorrhage, malignancy, hepatic and renal disease) can usually be

met with oral supplementation.

Cyanocobalamin Injection is also suitable for the vitamin B₁₂ absorption test (**Schilling Test**).

CONTRAINDICATIONS

Sensitivity to cobalt and/or vitamin B₁₂ is a contraindication.

WARNINGS

Patients with early Leber's disease (hereditary optic nerve atrophy) who were treated with cyanocobalamin suffered severe and swift optic atrophy.

Hypokalemia and sudden death may occur in severe megaloblastic anemia which is treated intensely.

Anaphylactic shock and death have been reported after parenteral vitamin B₁₂ administration. An intradermal test dose is recommended before cyanocobalamin injection is administered to patients suspected of being sensitive to this drug.

This product contains benzyl alcohol. Benzyl alcohol has been reported to be associated with a fatal "Gaspings Syndrome" in premature infants.

This product contains aluminum that may be toxic. Aluminum may reach toxic levels with prolonged parenteral administration if kidney function is impaired.

Premature neonates are particularly at risk because their kidneys are immature, and they require large amounts of calcium and phosphate solutions, which contain aluminum.

Research indicates that patients with impaired kidney function, including premature neonates, who receive parenteral levels of aluminum at greater than 4 to 5 mcg/kg/day accumulate aluminum at levels associated with central nervous system and bone toxicity. Tissue loading may occur at even lower rates of administration.

PRECAUTIONS

General Precautions

Vitamin B₁₂ deficiency that is allowed to progress for longer than 3 months may produce permanent degenerative lesions of the spinal cord.

Doses of folic acid greater than 0.1 mg per day may result in hematologic remission in patients with vitamin B₁₂ deficiency. Neurologic manifestations will not be prevented with folic acid, and if not treated with vitamin B₁₂, irreversible damage will result.

Doses of cyanocobalamin exceeding 10 mcg daily may produce hematologic response in patients with folate deficiency. Indiscriminate administration may mask the true diagnosis.

Information for Patients

Patients with pernicious anemia should be informed that they will require monthly

injections of vitamin B₁₂ for the remainder of their lives. Failure to do so will result in return of the anemia and in development of incapacitating and irreversible damage to the nerves of the spinal cord. Also, patients should be warned about the danger of taking folic acid in place of vitamin B₁₂, because the former may prevent anemia but allow progression of subacute combined degeneration.

A vegetarian diet which contains no animal products (including milk products or eggs) does not supply any vitamin B₁₂. Patients following such a diet, should be advised to take oral vitamin B₁₂ regularly. The need for vitamin B₁₂ is increased by pregnancy and lactation. Deficiency has been recognized in infants of vegetarian mothers who were breast fed, even though the mothers had no symptoms of deficiency at the time.

Laboratory Tests

During the initial treatment of patients with pernicious anemia, serum potassium must be observed closely the first 48 hours and potassium replaced if necessary.

Hematocrit, reticulocyte count, vitamin B₁₂, folate and iron levels should be obtained prior to treatment. Hematocrit and reticulocyte counts should be repeated daily from the fifth to seventh days of therapy and then frequently until the hematocrit is normal. If folate levels are low, folic acid should also be administered. If reticulocytes have not increased after treatment or if reticulocyte counts do not continue at least twice normal as long as the hematocrit is less than 35%, diagnosis or treatment should be reevaluated. Repeat determinations of iron and folic acid may reveal a complicating illness that might inhibit the response of the marrow.

Patients with pernicious anemia have about 3 times the incidence of carcinoma of the stomach as the general population, so appropriate tests for this condition should be carried out when indicated.

Drug/Laboratory Test Interactions

Persons taking most antibiotics, methotrexate and pyrimethamine invalidate folic acid and vitamin B₁₂ diagnostic blood assays.

Colchicine para-aminosalicylic acid and heavy alcohol intake for longer than 2 weeks may produce malabsorption of vitamin B₁₂.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Long term studies in animals to evaluate carcinogenic potential have not been done. There is no evidence from long-term use in patients with pernicious anemia that cyanocobalamin is carcinogenic. Pernicious anemia is associated with an increased incidence of carcinoma of the stomach, but this is believed to be related to the underlying pathology and not to treatment with cyanocobalamin.

Pregnancy

Teratogenic Effects

Adequate and well-controlled studies have not been done in pregnant women. However, vitamin B₁₂ is an essential vitamin and requirements are increased during pregnancy. Amounts of vitamin B₁₂ that are recommended by the Food and Nutrition Board, National Academy of Science-National Research Council for pregnant women (4 mcg

daily) should be consumed during pregnancy.

Nursing Mothers

Vitamin B₁₂ is known to be excreted in human milk. Amounts of vitamin B₁₂ that are recommended by the Food and Nutrition Board, National Academy of Science-National Research Council for lactating women (4 mcg daily) should be consumed during lactation.

Pediatric Use

Intake in children should be in the amount (0.5 to 3 mcg daily) recommended by the Food and Nutrition Board, National Academy of Science-National Research Council.

ADVERSE REACTIONS

Generalized: Anaphylactic shock and death have been reported with administration of parenteral vitamin B₁₂ (see **WARNINGS**).

Cardiovascular: Pulmonary edema and congestive heart failure early in treatment; peripheral vascular thrombosis

Hematological: Polycythemia vera

Gastrointestinal: Mild transient diarrhea

Dermatological: Itching; transitory exanthema

Miscellaneous: Feeling of swelling of entire body

To report SUSPECTED ADVERSE REACTIONS, contact Meitheal Pharmaceuticals Inc. at 1-844-824-8426 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

OVERDOSAGE

No overdosage has been reported with this drug.

DOSAGE AND ADMINISTRATION

Avoid using the intravenous route. Use of this product intravenously will result in almost all of the vitamin being lost in the urine.

Pernicious Anemia

Parenteral vitamin B₁₂ is the recommended treatment and will be required for the remainder of the patient's life. The oral form is not dependable. A dose of 100 mcg daily for 6 or 7 days should be administered by intramuscular or deep subcutaneous injection. If there is clinical improvement and if a reticulocyte response is observed, the same amount may be given on alternate days for seven doses, then every 3 to 4 days for another 2 to 3 weeks. By this time hematologic values should have become normal. This regimen should be followed by 100 mcg monthly for life. Folic acid should be administered concomitantly if needed.

Patients with Normal Intestinal Absorption

Where the oral route is not deemed adequate, initial treatment similar to that for patients with pernicious anemia may be indicated depending on the severity of the deficiency. Chronic treatment should be with an oral B₁₂ preparation. If other vitamin deficiencies are present, they should be treated.

Schilling Test

The flushing dose is 1,000 mcg.

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.

HOW SUPPLIED

Cyanocobalamin Injection, USP is a sterile solution of cyanocobalamin and is supplied as follows:

NDC	Cyanocobalamin Injection, USP (1,000 mcg per mL)	Package Factor
71288- 300 -02	1,000 mcg per 1 mL Multi-Dose Vial	25 vials per carton
71288- 301 -11	10,000 mcg per 10 mL Multi-Dose Vial	10 vials per carton
71288- 302 -30	30,000 mcg per 30 mL Multi-Dose Vial	1 vial per carton

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F). [See USP Controlled Room Temperature.]

Protect from light.

Sterile, Nonpyrogenic.

The container closure is not made with natural rubber latex.

meitheal®

Mfd. for Meitheal Pharmaceuticals
Chicago, IL 60631 (USA)
©2024 Meitheal Pharmaceuticals Inc.

Mfd. by Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd.
Nanjing, China 210061

September 2024

8H1AAM9-00

PRINCIPAL DISPLAY PANEL - Cyanocobalamin Injection, USP, 1 mL Vial Label

NDC 71288-**300**-01

Rx Only

Cyanocobalamin Injection, USP

1,000 mcg per mL

For Intramuscular or Subcutaneous Use Only

1 mL Multi-Dose Vial



PRINCIPAL DISPLAY PANEL - Cyanocobalamin Injection, USP, 1 mL Carton

NDC 71288-300-02

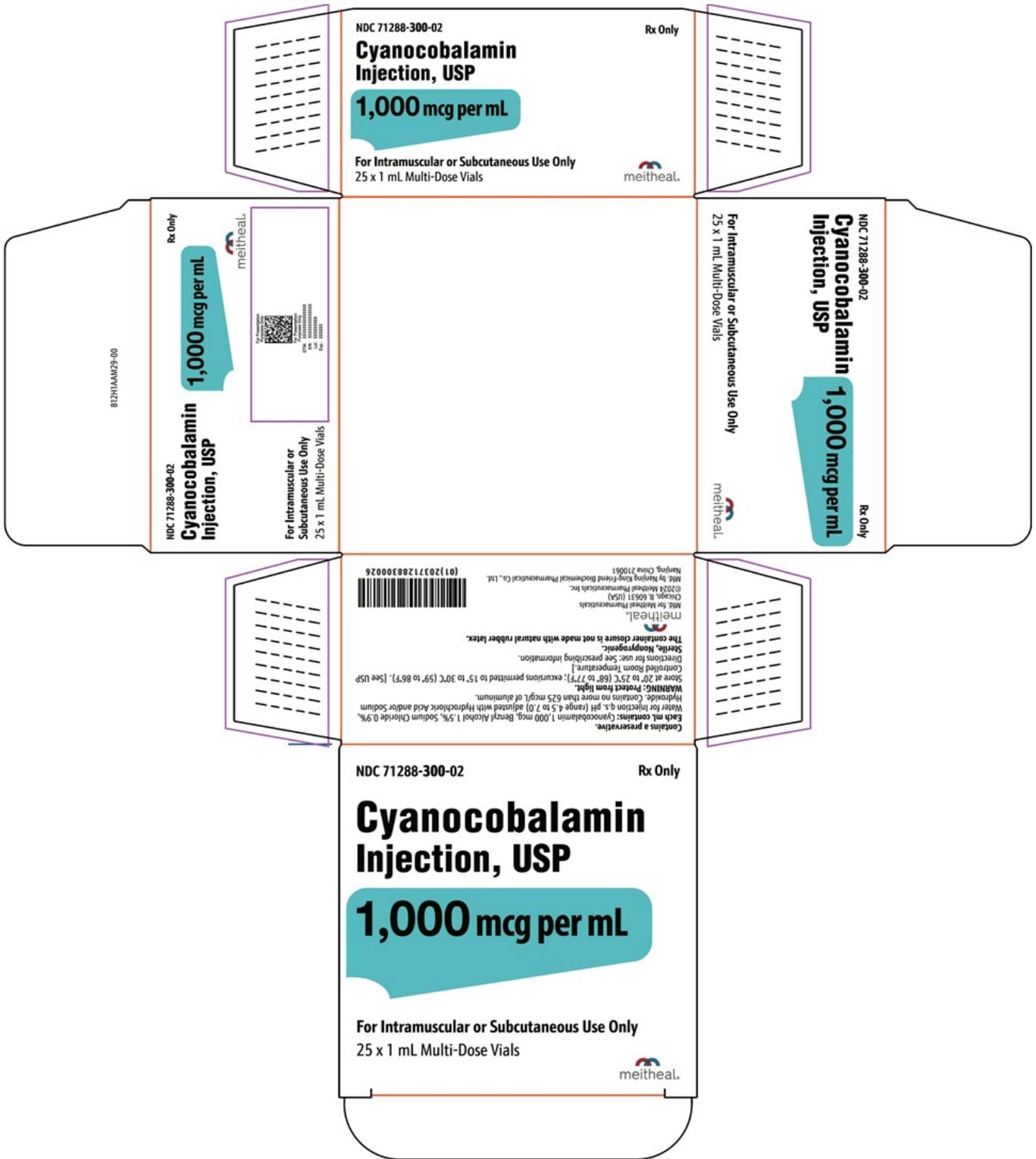
Rx Only

Cyanocobalamin Injection, USP

1,000 mcg per mL

For Intramuscular or Subcutaneous Use Only

25 x 1 mL Multi-Dose Vials



PRINCIPAL DISPLAY PANEL - Cyanocobalamin Injection, USP, 10 mL Vial Label

NDC 71288-301-10

Rx Only

Cyanocobalamin Injection, USP

10,000 mcg per 10 mL

(1,000 mcg per mL)

For Intramuscular or Subcutaneous Use Only

10 mL Multi-Dose Vial

NDC 71288-301-10 Rx Only

Cyanocobalamin Injection, USP

10,000 mcg per 10 mL
(1,000 mcg per mL)

For Intramuscular or Subcutaneous Use Only
10 mL Multi-Dose Vial

Contains a preservative.
Each mL contains:
Cyanocobalamin 1,000 mcg,
Benzyl Alcohol 1.5%, Sodium Chloride 0.9%, Water for Injection q.s. pH adjusted with Hydrochloric Acid and/or Sodium Hydroxide.
Contains no more than 625 mcg/L of aluminum. **WARNING: Protect from light.** Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature.] Directions for use: See package insert.



(01)00371288301104

Lot:
Exp.:

801H1CAM90-00

 **meitheal.**
Mfd. for Meitheal Pharmaceuticals
Chicago, IL 60631 (USA)
Made in China
©2024 Meitheal Pharmaceuticals Inc.

PRINCIPAL DISPLAY PANEL - Cyanocobalamin Injection, USP, 10 mL Carton

NDC 71288-301-11

Rx Only

Cyanocobalamin Injection, USP

10,000 mcg per 10 mL

(1,000 mcg per mL)

For Intramuscular or Subcutaneous Use Only

10 x 10 mL Multi-Dose Vials



PRINCIPAL DISPLAY PANEL - Cyanocobalamin Injection, USP, 30 mL Vial Label

NDC 71288-302-30

Rx Only

Cyanocobalamin Injection, USP

30,000 mcg per 30 mL

(1,000 mcg per mL)

For Intramuscular or Subcutaneous Use Only

30 mL Multi-Dose Vial

NDC 71288-302-30 Rx Only

Cyanocobalamin Injection, USP

30,000 mcg per 30 mL
(1,000 mcg per mL)

For Intramuscular or Subcutaneous Use Only

30 mL Multi-Dose Vial

Contains a preservative.

Each mL contains: Cyanocobalamin 1,000 mcg, Benzyl Alcohol 1.5%, Sodium Chloride 0.9%, Water for Injection q.s. pH adjusted with Hydrochloric Acid and/or Sodium Hydroxide. Contains no more than 625 mcg/L of aluminum.

WARNING: Protect from light.

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F). [See USP Controlled Room Temperature.]

Directions for use: See package insert.

Sterile, Nonpyrogenic.

This container closure is not made with natural rubber latex.

Lot:
Exp.:



Mfd. for Meitheal Pharmaceuticals
Chicago, IL 60631 (USA)
Made in China
©2024 Meitheal Pharmaceuticals Inc.

801H1EAM90-00



(01)00371288302309

PRINCIPAL DISPLAY PANEL - Cyanocobalamin Injection, USP, 30 mL Carton

NDC 71288-302-30

Rx Only

Cyanocobalamin Injection, USP

30,000 mcg per 30 mL

(1,000 mcg per mL)

For Intramuscular or Subcutaneous Use Only

1 x 30 mL Multi-Dose Vial



CYANOCOBALAMIN

cyanocobalamin injection

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:71288-300
Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of	Strength
-----------------	----------	----------

Ingredient Name		Strength	Strength	
CYANOCOBALAMIN (UNII: P6YC3EG204) (CYANOCOBALAMIN - UNII:P6YC3EG204)		CYANOCOBALAMIN	1000 ug in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		9 mg in 1 mL		
BENZYL ALCOHOL (UNII: LKG8494WBH)		15 mg in 1 mL		
WATER (UNII: 059QF0KO0R)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71288-300-02	25 in 1 CARTON	11/15/2024	
1	NDC:71288-300-01	1 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA214316	11/15/2024		

CYANOCOBALAMIN			
cyanocobalamin injection			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:71288-301
Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
CYANOCOBALAMIN (UNII: P6YC3EG204) (CYANOCOBALAMIN - UNII:P6YC3EG204)		CYANOCOBALAMIN	1000 ug in 1 mL
Inactive Ingredients			
Ingredient Name		Strength	
SODIUM CHLORIDE (UNII: 451W47IQ8X)		9 mg in 1 mL	
BENZYL ALCOHOL (UNII: LKG8494WBH)		15 mg in 1 mL	

WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71288-301-11	10 in 1 CARTON	11/15/2024	
1	NDC:71288-301-10	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214316	11/15/2024	

CYANOCOBALAMIN

cyanocobalamin injection

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:71288-302
Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CYANOCOBALAMIN (UNII: P6YC3EG204) (CYANOCOBALAMIN - UNII:P6YC3EG204)	CYANOCOBALAMIN	1000 ug in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	9 mg in 1 mL
BENZYL ALCOHOL (UNII: LKG8494WBH)	15 mg in 1 mL
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:71288			

1	NDC: 71288-302-30	1 in 1 CARTON	11/15/2024	
1		30 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA214316		11/15/2024	

Labeler - Meitheal Pharmaceuticals Inc. (080548348)

Establishment			
Name	Address	ID/FEI	Business Operations
Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd.		421297554	MANUFACTURE(71288-300, 71288-301, 71288-302)

Revised: 11/2024

Meitheal Pharmaceuticals Inc.