RETATRUTIDE PEN- retatrutide pen injection, solution Guangzhou Yixin Cross-border E-commerce Co., Ltd.

Active ingredient (in each 0.75 mL pen)

Retatrutide — 10 mg

Purpose

Antidiabetic / Weight management agent

Uses

To improve blood glucose control in adults with type 2 diabetes.

For long-term weight management and reduction of visceral fat in adults who are overweight or obese, with or without diabetes.

Warnings

Pancreatitis: Acute pancreatitis has been reported with GLP-1 receptor agonists. If severe or persistent abdominal pain occurs (with or without vomiting), stop using RETATRUTIDE and contact your healthcare provider immediately.

Allergic reactions: Serious hypersensitivity reactions (including anaphylaxis) have been reported.

Stop use and seek medical attention if you develop rash, swelling, or difficulty breathing.

Severe gastrointestinal disease: RETATRUTIDE has not been studied in patients with severe GI disorders; use is not recommended.

Gallbladder disease: Acute gallbladder events, including cholelithiasis, have occurred. Evaluate promptly if gallbladder disease is suspected.

Hypoglycemia warning:

Do not use RETATRUTIDE if you have a history of recurrent or severe hypoglycemia.

RETATRUTIDE may further lower blood glucose levels. Use is contraindicated in individuals with baseline hypoglycemia or conditions predisposing to hypoglycemia.

Pregnancy and breastfeeding: Do not use if pregnant or breastfeeding.

Pediatric use: Not for use in children.

Drug Interactions / Precautions

Other antidiabetic medications: Retatrutide lowers blood glucose. Using it with insulin or sulfonylureas may increase the risk of hypoglycemia. Blood glucose should be closely monitored; doses of other antidiabetic drugs may need adjustment.

Drugs affecting gastrointestinal motility: Retatrutide may delay gastric emptying. Co-

administration with prokinetic agents (e.g., domperidone) or antacids may alter absorption or worsen GI discomfort.

Cardiovascular medications: Caution with statins, digoxin, or warfarin.

Statins: absorption may be affected; monitor lipid levels.

Warfarin: changes in metabolism may alter anticoagulant effects; monitor INR levels.

General precautions: Retatrutide should be used only under medical supervision. Inform your healthcare provider about all medications, including prescription, over-the-counter, and herbal products.

Possible side effects

Common: nausea, diarrhea, decreased appetite, vomiting, constipation, indigestion, abdominal pain.

Stop use and contact a doctor if symptoms become severe or persistent.

When to stop and call your doctor

Stop using the pen and contact your physician if you experience:

- Severe or persistent abdominal pain or vomiting
- Symptoms of an allergic reaction
- Signs of pancreatitis
- Any other unexpected or severe reaction

Titration schedule:

Weeks 1-4 0.19 mL (2.5 mg) once weekly

Weeks 5-8 0.37 mL (5 mg) once weekly

Weeks 9-12 0.56 mL (7.5 mg) once weekly

Week 13 and beyond 0.75 mL (10 mg) once weekly

Directions for Use

- For subcutaneous injection only.
- Inject once weekly on the same day each week, any time of day, with or without meals.
- Recommended injection sites: abdomen, thigh, or upper arm.
- Clean the injection site with alcohol before injection.
- Rotate injection sites each week.
- Use a new needle for each injection. Do not share your pen.

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Keep out of reach of children.

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Inactive ingredients

Sterile Water for Injection

PRINCIPAL DISPLAY PANEL

10mg减肥注射器-水剂款-双插盒外盒

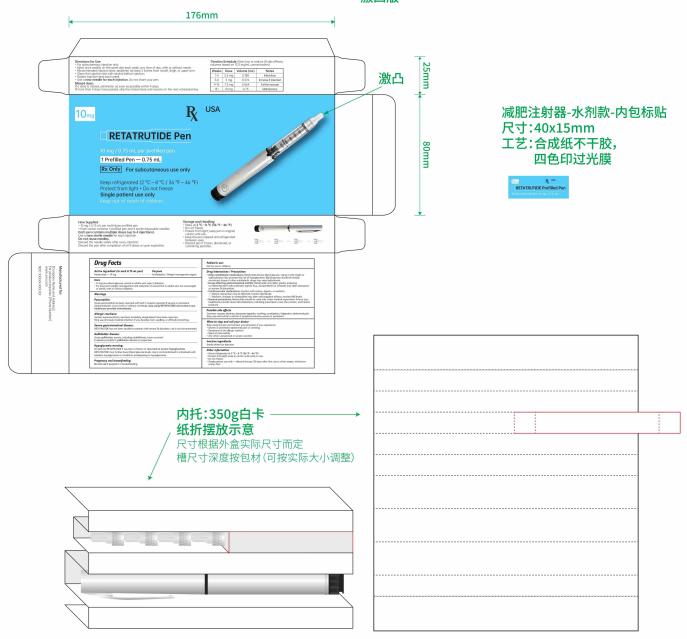
尺寸:176x25x80mm

工艺:350g白卡过光膜,四色印,指定位置激凸

内托:350g白卡纸折卡托



激凸版



30mg减肥注射器-水剂款-双插盒外盒

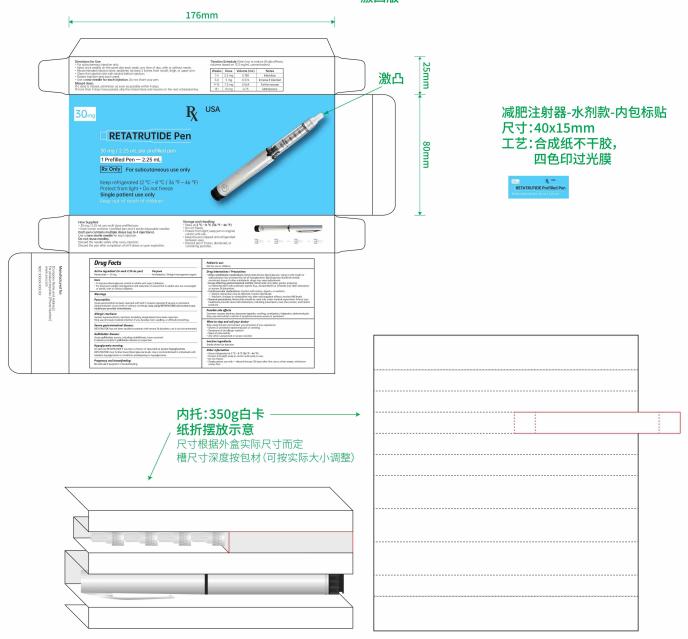
尺寸:176x25x80mm

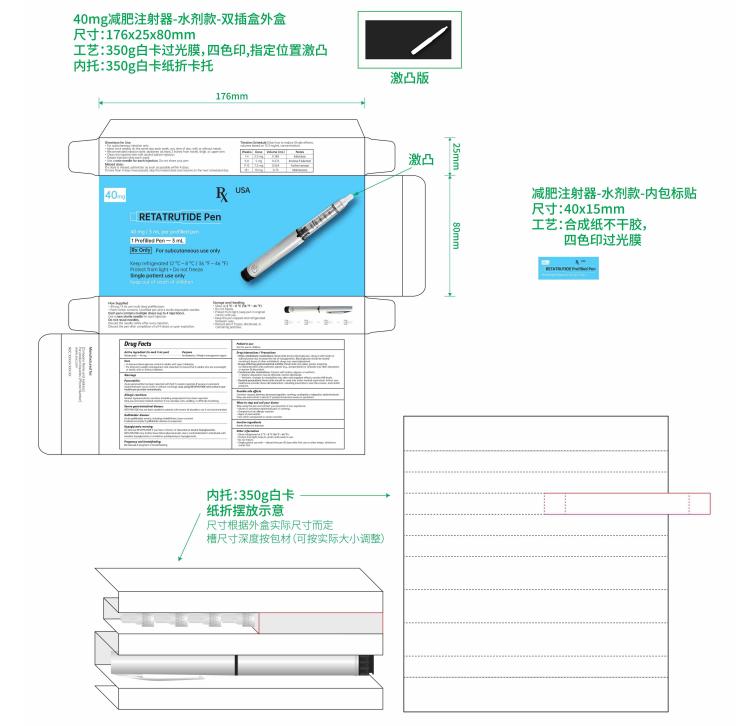
工艺:350g白卡过光膜,四色印,指定位置激凸

内托:350g白卡纸折卡托



激凸版





retatrutide pen injection, solution Product Information Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:84778-106 Route of Administration SUBCUTANEOUS Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
RETATRUTIDE (UNII: NOP2Y096GV) (RETATRUTIDE - UNII:NOP2Y096GV)	RETATRUTIDE	10 mg in 100 mL

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

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:84778- 106-01	4 in 1 BOX	10/29/2025		
1		0.75 mL in 1 PAIL; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)			
2	NDC:84778- 106-02	4 in 1 BOX	10/29/2025		
2		2.25 mL in 1 SYRINGE; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)			
3	NDC:84778- 106-03	4 in 1 BOX	10/29/2025		
3		3 mL in 1 SYRINGE; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Export only		10/29/2025	

Labeler - Guangzhou Yixin Cross-border E-commerce Co., Ltd. (455800881)

Registrant - Guangzhou Yixin Cross-border E-commerce Co., Ltd. (455800881)

Establishment				
Name	Address	ID/FEI	Business Operations	
Guangzhou Yixin Cross-border E-commerce Co., Ltd.		455800881	label(84778-106) , manufacture(84778- 106)	

Revised: 10/2025 Guangzhou Yixin Cross-border E-commerce Co., Ltd.