

ELTROMBOPAG- eltrombopag tablet
Zydus Lifesciences Limited

ELTROMBOPAG tablets, for oral use

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL

NDC 70771-1925-3

Eltrombopag Tablets, 12.5 mg

30 Tablets

Rx only



NDC 70771-1926-3

Eltrombopag Tablets, 25 mg

30 Tablets

Rx only

3 N 7071013963 6

Rev.: 01/26

Eltrombopag Tablets

25 mg

Swallow tablets whole. Do not split, chew, or crush tablets

PHARMACIST: Dispense the medication guide provided separately to each patient.

zydus

30 Tablets
Rx only

Each film-coated tablet contains eltrombopag olamine equivalent to 25 mg of eltrombopag free acid.
 Dosage: See accompanying prescribing information.
 This package is child-resistant.
 Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [see USP Controlled Room Temperature].
 Dispense in original bottle.
 Important: Use safety closures when dispensing this product unless otherwise directed by physicians or requested by purchaser.
 Keep this and all drugs out of the reach of children.
 Do not use if printed safety seal under cap is broken or missing.
 Medication Guide available at www.zydususa.com/medguides or call 1-877-993-8779.
 Mfg. by: Zydus Lifesciences Ltd., Ahmedabad, India

NDC 70771-1927-3

Eltrombopag Tablets, 50 mg

30 Tablets

Rx only

3 N 7071013973 5

Rev.: 01/26

Eltrombopag Tablets

50 mg

Swallow tablets whole. Do not split, chew, or crush tablets

PHARMACIST: Dispense the medication guide provided separately to each patient.

zydus

30 Tablets
Rx only

Each film-coated tablet contains eltrombopag olamine equivalent to 50 mg of eltrombopag free acid.
 Dosage: See accompanying prescribing information.
 This package is child-resistant.
 Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [see USP Controlled Room Temperature].
 Dispense in original bottle.
 Important: Use safety closures when dispensing this product unless otherwise directed by physicians or requested by purchaser.
 Keep this and all drugs out of the reach of children.
 Do not use if printed safety seal under cap is broken or missing.
 Medication Guide available at www.zydususa.com/medguides or call 1-877-993-8779.
 Mfg. by: Zydus Lifesciences Ltd., Ahmedabad, India

NDC 70771-1928-3

Eltrombopag Tablets, 75 mg

30 Tablets

Rx only

Rev.: 01/26

Eltrombopag Tablets

75 mg

Swallow tablets whole. Do not split, chew, or crush tablets

PHARMACIST: Dispense the medication guide provided separately to each patient.

30 Tablets
Rx only

Each film-coated tablet contains eltrombopag olamine equivalent to 75 mg of eltrombopag free acid.
Dosage: See accompanying prescribing information.
This package is child-resistant.
Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [see USP Controlled Room Temperature].
Dispense in original bottle.
Important: Use safety closures when dispensing this product unless otherwise directed by physicians or requested by purchaser.
Keep this and all drugs out of the reach of children.
Do not use if printed safety seal under cap is broken or missing.
Medication Guide available at www.zydususa.com/medguides or call 1-877-993-8779.
Mfg. by: Zydus Lifesciences Ltd., Ahmedabad, India

ELTROMBOPAG

eltrombopag tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1925
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ELTROMBOPAG OLAMINE (UNII: 4U07F515LG) (ELTROMBOPAG - UNII:S56D65XJ9G)	ELTROMBOPAG	12.5 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
XYLITOL (UNII: VCQ006KQ1E)	

Product Characteristics

Color	WHITE (white to light blue)	Score	no score
Shape	ROUND	Size	5mm

Flavor		Imprint Code		I
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1925-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2026	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA		ANDA216281	01/14/2026	

ELTROMBOPAG			
eltrombopag tablet			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1926
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ELTROMBOPAG OLAMINE (UNII: 4U07F515LG) (ELTROMBOPAG - UNII:S56D65XJ9G)		ELTROMBOPAG	25 mg
Inactive Ingredients			
Ingredient Name			Strength
FERRIC OXIDE RED (UNII: 1K09F3G675)			
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)			
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)			
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			
POVIDONE K30 (UNII: U725QWY32X)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
XYLITOL (UNII: VCQ006KQ1E)			
Product Characteristics			
Color	YELLOW (light yellow)	Score	no score

Shape	ROUND	Size	7mm	
Flavor		Imprint Code	I7	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1926-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2026	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA216281		01/14/2026	

ELTROMBOPAG			
eltrombopag tablet			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1927
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ELTROMBOPAG OLAMINE (UNII: 4U07F515LG) (ELTROMBOPAG - UNII:S56D65XJ9G)		ELTROMBOPAG	50 mg
Inactive Ingredients			
Ingredient Name			Strength
FD&C BLUE NO. 2--ALUMINUM LAKE (UNII: 4AQJ3LG584)			
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)			
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			
POVIDONE K30 (UNII: U725QWY32X)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
XYLITOL (UNII: VCQ006KQ1E)			
Product Characteristics			
Color	BLUE (light blue)	Score	no score

Shape	ROUND	Size	9mm
Flavor		Imprint Code	112
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1927-7	14 in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2026	
2	NDC:70771-1927-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA216281	01/14/2026	

ELTROMBOPAG

eltrombopag tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1928
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ELTROMBOPAG OLAMINE (UNII: 4U07F515LG) (ELTROMBOPAG - UNII:S56D65XJ9G)	ELTROMBOPAG	75 mg

Inactive Ingredients

Ingredient Name	Strength
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
XYLITOL (UNII: VCQ006KQ1E)	

Product Characteristics

Color	PURPLE (light purple)	Score	no score
Shape	ROUND	Size	11mm
Flavor		Imprint Code	l17
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1928-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA216281	01/14/2026	

Labeler - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		863362789	ANALYSIS(70771-1925, 70771-1926, 70771-1927, 70771-1928) , MANUFACTURE(70771-1925, 70771-1926, 70771-1927, 70771-1928)

Revised: 1/2026

Zydus Lifesciences Limited