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

NDC 70771-1927-3

Eltrombopag Tablets, 50 mg

30 Tablets

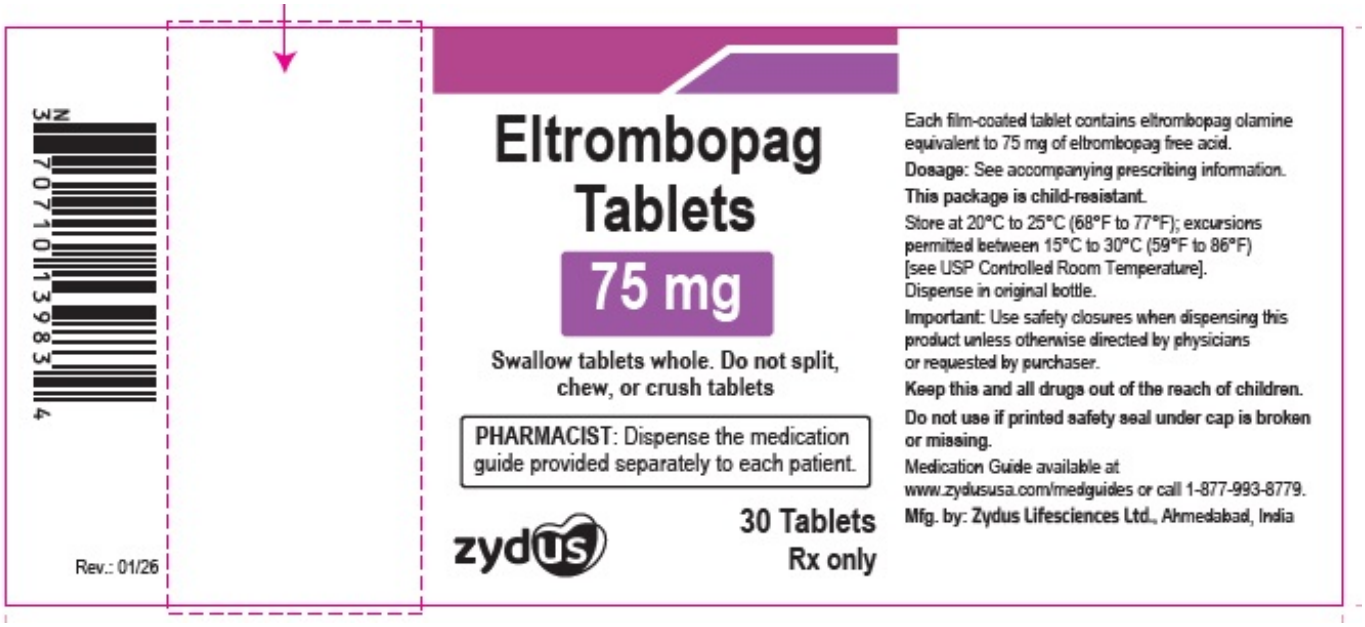
Rx only

NDC 70771-1928-3

Eltrombopag Tablets, 75 mg

30 Tablets

Rx only



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

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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**	I17
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