

## Promacta - (50 mg and 75 mg ; Tablets)

<b>Generic Name</b>	Eltrombopag Olamine	<b>Innovator</b>	Novartis
<b>Dosage</b>	50 mg and 75 mg ; Tablets	<b>Branded US Sales</b>	More Than \$1000 mn
<b>Probable FTF</b>	None	<b>Known Para IV Filers</b>	Less Than 5
<b>Other ANDA developers</b>	More Than 5	<b>Tentative Approvals</b>	None
<b>Final Approvals</b>	None	<b>Generic Launches</b>	None
<b>Indication</b>	PROMACTA is a thrombopoietin receptor agonist indicated: • for the treatment of thrombocytopenia in adult and pediatric patients 1 year and older with persistent or chronic immune thrombocytopenia (ITP) who have had an insufficient response to corticosteroids, immunoglobulins, or splenectomy. PROMACTA should be used only in patients with ITP whose degree of thrombocytopenia and clinical condition increase the risk for bleeding. • for the treatment of thrombocytopenia in patients with chronic hepatitis C to allow the initiation and maintenance of interferon-based therapy. PROMACTA should be used only in patients with chronic hepatitis C whose degree of thrombocytopenia prevents the initiation of interferon-based therapy or limits the ability to maintain interferon-based therapy. • in combination with standard immunosuppressive therapy for the firstline treatment of adult and pediatric patients 2 years and older with severe aplastic anemia. • for the treatment of patients with severe aplastic anemia who have had an insufficient response to immunosuppressive therapy.		
<b>Complexities</b>	Yes		

### Chronology Of Events

Please Contact [contact@researchdelta.com](mailto:contact@researchdelta.com) to get Detailed Information.

### Executive Summary

Please Contact [contact@researchdelta.com](mailto:contact@researchdelta.com) to get Detailed Information.

### Patent Status

Please Contact [contact@researchdelta.com](mailto:contact@researchdelta.com) to get Detailed Information.

### Launch Timelines and Competition

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## Chronology Of Events

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## Promacta - (12.5 mg and 25 mg ; Tablets)

<b>Generic Name</b>	Eltrombopag Olamine	<b>Innovator</b>	Novartis
<b>Dosage</b>	12.5 mg and 25 mg ; Tablets	<b>Branded US Sales</b>	More Than \$1000 mn
<b>Probable FTF</b>	None	<b>Known Para IV Filers</b>	Less Than 5
<b>Other ANDA developers</b>	More Than 5	<b>Tentative Approvals</b>	None
<b>Final Approvals</b>	None	<b>Generic Launches</b>	None
<b>Indication</b>	PROMACTA is a thrombopoietin receptor agonist indicated: • for the treatment of thrombocytopenia in adult and pediatric patients 1 year and older with persistent or chronic immune thrombocytopenia (ITP) who have had an insufficient response to corticosteroids, immunoglobulins, or splenectomy. PROMACTA should be used only in patients with ITP whose degree of thrombocytopenia and clinical condition increase the risk for bleeding. • for the treatment of thrombocytopenia in patients with chronic hepatitis C to allow the initiation and maintenance of interferon-based therapy. PROMACTA should be used only in patients with chronic hepatitis C whose degree of thrombocytopenia prevents the initiation of interferon-based therapy or limits the ability to maintain interferon-based therapy. • in combination with standard immunosuppressive therapy for the firstline treatment of adult and pediatric patients 2 years and older with severe aplastic anemia. • for the treatment of patients with severe aplastic anemia who have had an insufficient response to immunosuppressive therapy.		
<b>Complexities</b>	Yes		

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