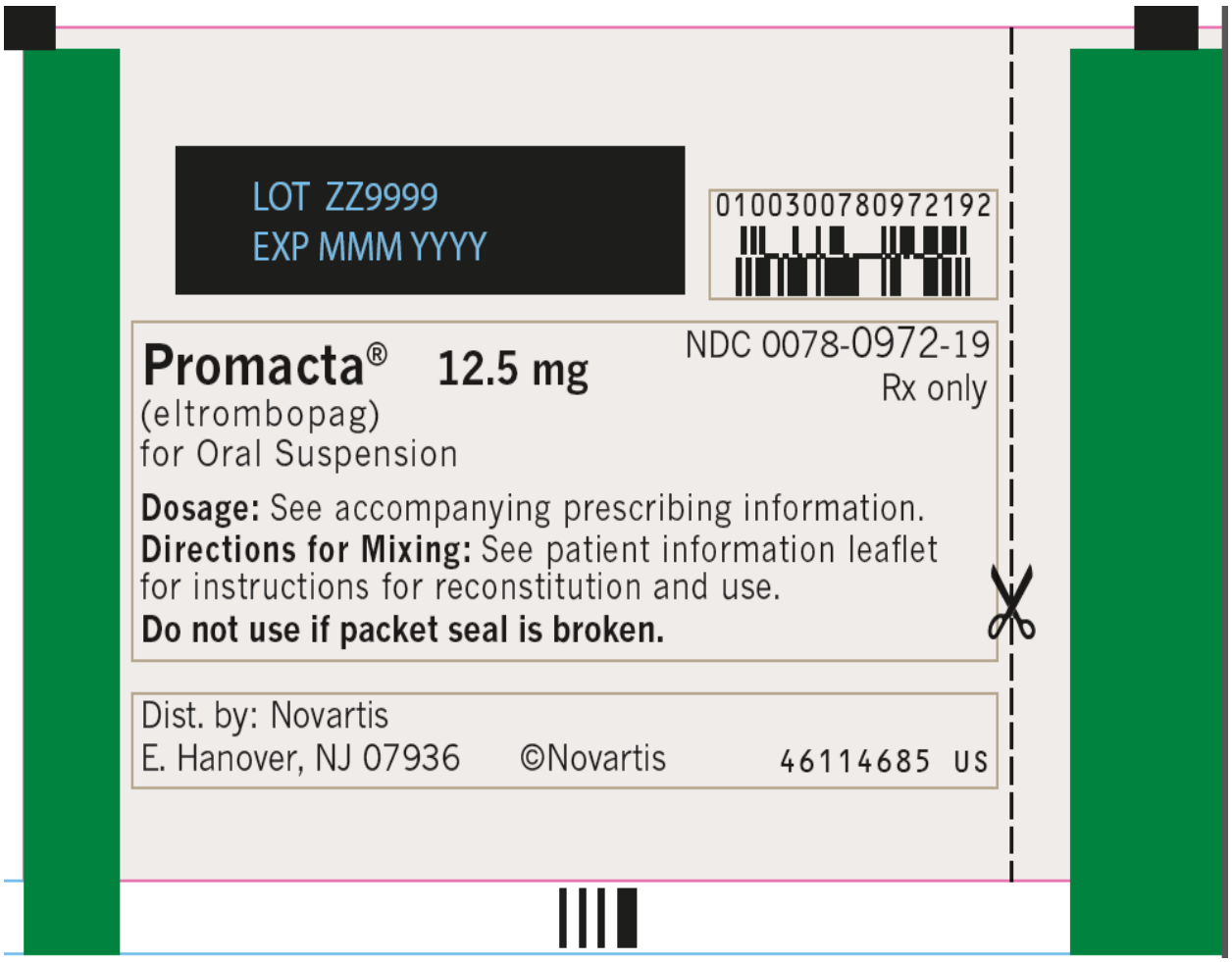
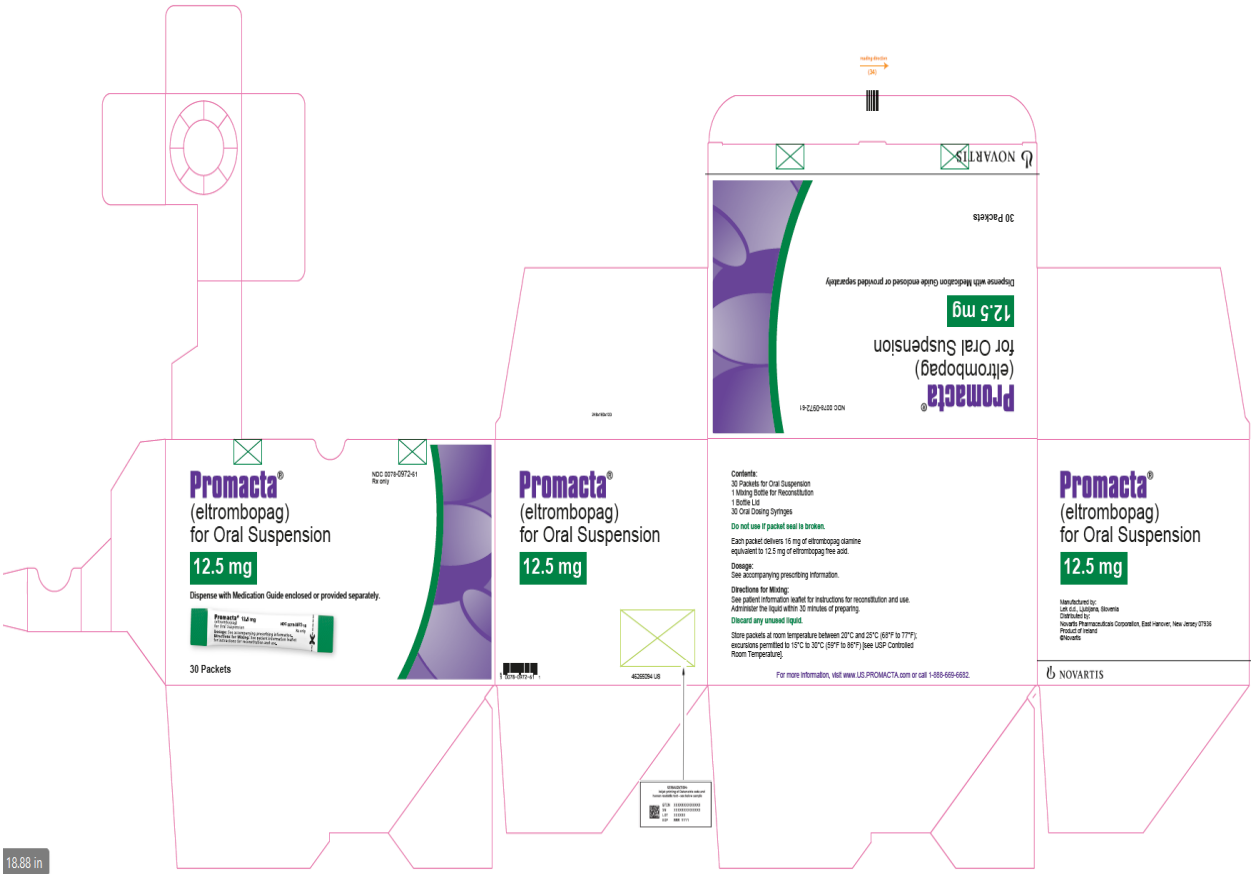
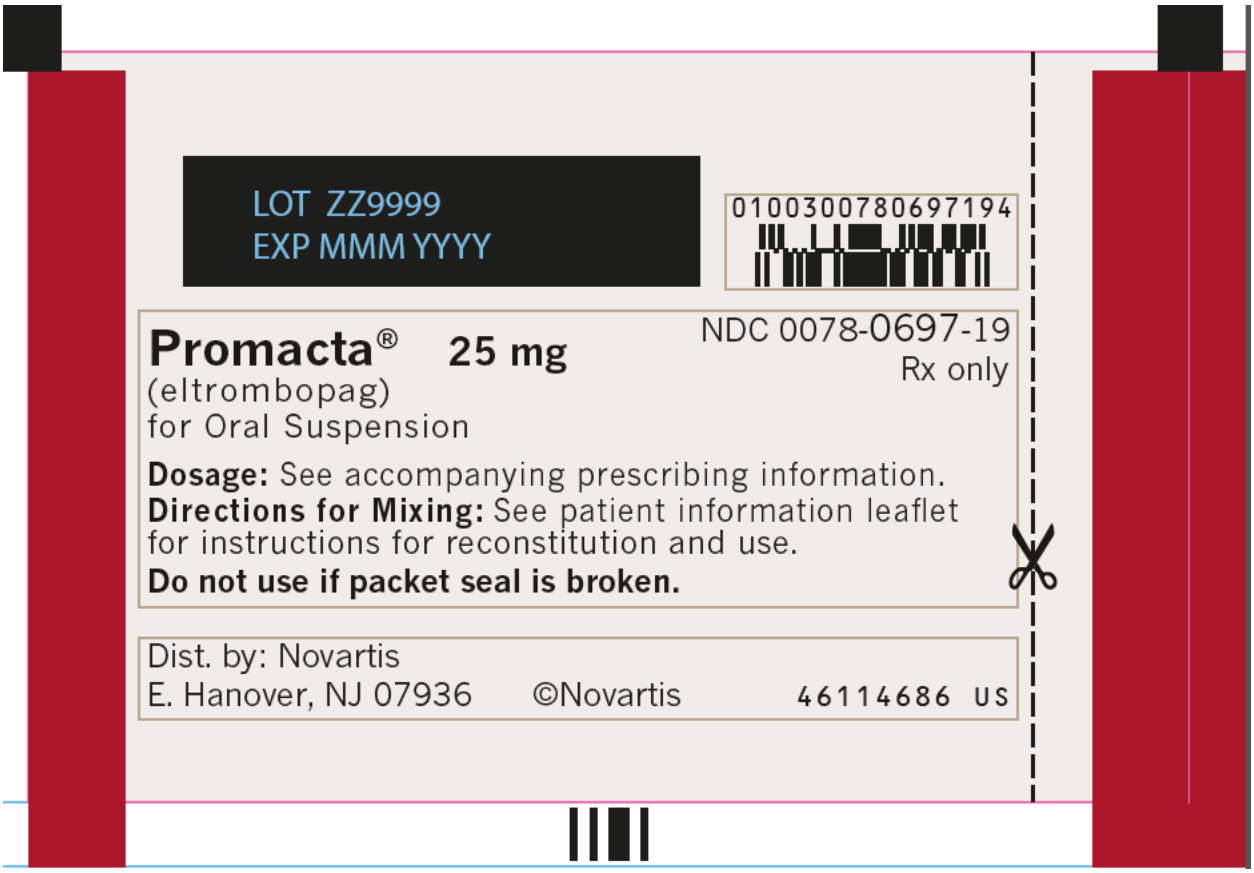

LOT ZZ9999
EXP MMM YYYY
0100300780972192
Promacta® 12.5 mg
(eltrombopag)
for Oral Suspension
NDC 0078-0972-19
Rx only
Dosage: See accompanying prescribing information.
Directions for Mixing: See patient information leaflet for instructions for reconstitution and use.
Do not use if packet seal is broken.
Dist. by: Novartis
E. Hanover, NJ 07936
©Novartis
46114685 US

NDC 0078-0972-23
Rx only
Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Dispense with Medication Guide enclosed with this package or provided separately.
Store packets at room temperature between 20°C and 25°C (68°F to 77°F); excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled Room Temperature].
N 3 0078-0972-23 9
Manufactured by:
Lek d.d.
Ljubljana, Slovenia
Distributed by:
Novartis Pharmaceuticals Corporation
East Hanover, New Jersey 07936
©Novartis
46265089 US
54x54x111
NOVARTIS
Contents:
30 Packets for Oral Suspension
Do not use if packet seal is broken.
Each packet delivers 16 mg of eltrombopag olamine equivalent to 12.5 mg of eltrombopag free acid.
Dosage:
See accompanying prescribing information.
Directions for Mixing:
See patient information leaflet for instructions for reconstitution and use.
Administer the liquid within 30 minutes of preparing.
Discard any unused liquid.
For more information, visit www.US.PROMACTA.com or call 1-888-669-6682.
Variable data printing
MMM YYYY
XXXXXXX


Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Dispense with Medication Guide enclosed or provided separately.
30 Packets
NDC 0078-0972-61
Rx only
Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Contents:
30 Packets for Oral Suspension
1 Mixing Bottle for Reconstitution
1 Bottle Lid
30 Oral Dosing Syringes
Do not use if packet seal is broken.
Each packet delivers 16 mg of eltrombopag olamine
equivalent to 12.5 mg of eltrombopag free acid.
Dosage:
See accompanying prescribing information.
Directions for Mixing:
See patient information leaflet for instructions for reconstitution and use.
Administer the liquid within 30 minutes of preparing.
Discard any unused liquid.
Store packets at room temperature between 20°C and 25°C (68°F to 77°F);
excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled
Room Temperature].
For more information, visit www.US.PROMACTA.com or call 1-888-669-6682.
Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Manufactured by:
Lek d.d., Ljubljana, Slovenia
Distributed by:
Novartis Pharmaceuticals Corporation, East Hanover, New Jersey 07936
Product of Ireland
©Novartis
NOVARTIS
18.88 in

Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Dispense with Medication Guide enclosed or provided separately.
30 Packets
NDC 0078-0972-61
Rx only
Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
46268293 US
Contents:
30 Packets for Oral Suspension
1 Mixing Bottle for Reconstitution
1 Bottle List
30 Oral Dosing Syringes
Do not use if packet seal is broken.
Each packet delivers 16 mg of eltrombopag olamine
equivalent to 12.5 mg of eltrombopag free acid.
Dosage:
See accompanying prescribing information.
Directions for Mixing:
See patient information leaflet for instructions for reconstitution and use.
Administer the liquid within 30 minutes of preparing.
Discard any unused liquid.
Store packets at room temperature between 20°C and 25°C (68°F to 77°F);
excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled
Room Temperature].
For more information, visit www.US.PROMACTA.com or call 1-888-669-6682.
Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Manufactured by:
Lab S.C., Ljubljana, Slovenia
Distributed by:
Novartis Pharmaceuticals Corporation, East Hanover, New Jersey 07936
Product of Switzerland
@Novartis
NOVARTIS
NOVARTIS
Promacta®
(eltrombopag)
for Oral Suspension
12.5 mg
Dispense with Medication Guide enclosed or provided separately
30 Packets
NDC 0078-0972-61
reading direction
(33)


LOT ZZ9999
EXP MMM YYYY
01003007806971 94
Promacta® 25 mg
(eltrombopag)
for Oral Suspension
NDC 0078-0697-19
Rx only
Dosage: See accompanying prescribing information.
Directions for Mixing: See patient information leaflet for instructions for reconstitution and use.
Do not use if packet seal is broken.
Dist. by: Novartis
E. Hanover, NJ 07936
©Novartis
46114686 US

NDC 0078-0697-23
Rx only
Promacta®
(eltrombopag)
for Oral Suspension
25 mg
Dispense with Medication Guide enclosed with this package or provided separately.
Store packets at room temperature between 20°C and 25°C (68°F to 77°F); excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled Room Temperature].
N 3 0078-0697-23 1
Manufactured by:
Lek d.d.
Ljubljana, Slovenia
Distributed by:
Novartis Pharmaceuticals Corporation
East Hanover, New Jersey 07936
©Novartis
NOVARTIS
Contents:
30 Packets for Oral Suspension
Do not use if packet seal is broken.
Each packet delivers 32 mg of eltrombopag olamine equivalent to 25 mg of eltrombopag free acid.
Dosage:
See accompanying prescribing information.
Directions for Mixing:
See patient information leaflet for instructions for reconstitution and use.
Administer the liquid within 30 minutes of preparing.
Discard any unused liquid.
For more information, visit www.US.PROMACTA.com or call 1-888-669-6682.
46265088 US
54x54x111
Variable data printing
MMM YYYY
XXXXXXX

Promacta®
(eltrombopag)
for Oral Suspension
25 mg
Dispense with Medication Guide enclosed or provided separately.
30 Packets
NDC 0078-0697-61
Rx only
Contents:
30 Packets for Oral Suspension
1 Mixing Bottle for Reconstitution
1 Bottle Lid
30 Oral Dosing Syringes
Do not use if packet seal is broken.
Each packet delivers 32 mg of eltrombopag olamine
equivalent to 25 mg of eltrombopag free acid.
Dosage:
See accompanying prescribing information.
Directions for Mixing:
See patient information leaflet for instructions for reconstitution and use.
Administer the liquid within 30 minutes of preparing.
Discard any unused liquid.
Store packets at room temperature between 20°C and 25°C (68°F to 77°F);
excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled
Room Temperature].
For more information, visit www.US.PROMACTA.com or call 1-888-669-6682.
Manufactured by:
Lek d.d., Ljubljana, Slovenia
Distributed by:
Novartis Pharmaceuticals Corporation, East Hanover, New Jersey 07936
Product of Ireland
©Novartis
NOVARTIS

Promacta®
(eltrombopag)
for Oral Suspension
25 mg
NDC 0078-0697-61
Rx only
Dispense with Medication Guide enclosed or provided separately.
30 Packets
Promacta®
(eltrombopag)
for Oral Suspension
25 mg
0078-0697-61
4635291 US
Contents:
30 Packets for Oral Suspension
1 Mixing Bottle for Reconstitution
1 Bottle Lid
30 Oral Dosing Syringes
Do not use if packet seal is broken.
Each packet delivers 32 mg of eltrombopag olamine
equivalent to 25 mg of eltrombopag free acid.
Dosage:
See accompanying prescribing information.
Directions for Mixing:
See patient information leaflet for instructions for reconstitution and use.
Administer the liquid within 30 minutes of preparing.
Discard any unused liquid.
Store packets at room temperature between 20°C and 25°C (68°F to 77°F);
excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled
Room Temperature].
For more information, visit www.US.PROMACTA.com or call 1-888-669-6682.
Promacta®
(eltrombopag)
for Oral Suspension
25 mg
Manufactured by:
Lek d.d., Ljubljana, Slovenia
Distributed by:
Novartis Pharmaceuticals Corporation, East Hanover, New Jersey 07936
Product of Switzerland
©Novartis
NOVARTIS
NOVARTIS
Promacta®
(eltrombopag)
for Oral Suspension
25 mg
Dispense with Medication Guide enclosed or provided separately.
30 Packets
NDC 0078-0697-61
mailing direction