HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use PRANDIMET safely and effectively. See full prescribing information for PRANDIMET.

PRANDIMET® (repaglinide and metformin hydrochloride) tablets, for oral use
Initial U.S. Approval: 2008

WARNING: LACTIC ACIDOSIS
See full prescribing information for complete boxed warning

- **Post-marketing cases of metformin-associated lactic acidosis have resulted in death, hypothermia, hypotension, and resistant bradyarrhythmias. Symptoms included malaise, myalgias, respiratory distress, somnolence, and abdominal pain. Laboratory abnormalities included elevated blood lactate levels, anion gap acidosis, increased lactate/pyruvate ratio; and metformin plasma levels generally >5 mcg/mL. (5.1)**
- **Risk factors include renal impairment, concomitant use of certain drugs, age > 65 years old, radiological studies with contrast, surgery and other procedures, hypoxic states, excessive alcohol intake, and hepatic impairment. Steps to reduce the risk of and manage metformin-associated lactic acidosis in these high risk groups are provided in the Full Prescribing Information. (5.1)**
- **If lactic acidosis is suspected, discontinue PRANDIMET and institute general supportive measures in a hospital setting. Prompt hemodialysis is recommended. (5.1)**

----------RECENT MAJOR CHANGES----------

Boxed Warning	4/2017
Dosing and Administration (2.2)	4/2017
Contraindications (4)	4/2017
Warnings and Precautions (5.1)	4/2017

----------INDICATIONS AND USAGE----------
PRANDIMET is a combination of repaglinide, a glinide, and metformin, a biguanide, indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus who are already treated with a glinide and metformin or who have inadequate glycemic control on a glinide alone or metformin alone. (1)

Limitation of Use:

- Not for treatment of type 1 diabetes mellitus or diabetic ketoacidosis (1)

----------DOSAGE AND ADMINISTRATION----------

- Instruct patients to take PRANDIMET within 30 minutes before meals. Instruct patients to skip the scheduled dose of PRANDIMET if a meal is skipped. (2.1)
- For patients inadequately controlled on metformin, start with 1 mg repaglinide/500 mg metformin twice daily and gradually titrate dose based on glycemic response. (2.1)
- For patients inadequately controlled on repaglinide, start with 1 mg repaglinide/500 mg metformin or 2 mg repaglinide/500 mg metformin twice daily and gradually titrate dose to reduce gastrointestinal side effects (2.1)
- For patients taking repaglinide and metformin, start PRANDIMET at the dose of repaglinide and metformin similar to (but not exceeding) the patient's current doses. Titrate as needed to achieve glycemic control (2.1)
- Do not exceed 10 mg repaglinide/2500 mg metformin daily or 4 mg repaglinide/1000 mg metformin per meal. (2.1)
- Assess renal function with estimated glomerular filtration rate (eGFR) before initiation of PRANDIMET and periodically thereafter (2.2)
  - Contraindicated in patients with eGFR below 30 mL/minute/1.73 m^2
  - Initiation is not recommended in patients with eGFR between 30 – 45 mL/minute/1.73 m^2
  - Assess risk/benefit of continuing if eGFR falls below 45 mL/minute/1.73 m^2
  - Discontinue if eGFR falls below 30 mL/minute/1.73 m2
- PRANDIMET may need to be discontinued at time of, or prior to, iodinated contrast imaging procedures (2.3)
- Dose modifications are required when used concominantly with some medications. (2.4,7)

----------DOSAGE FORMS AND STRENGTHS----------
Tablets:

- 1 mg repaglinide/500 mg metformin (3)
- 2 mg repaglinide/500 mg metformin (3)

----------CONTRAINDICATIONS----------

- Severe renal impairment (eGFR below 30 mL/min/1.73 m2). (4, 5.1)
- Metabolic acidosis, including diabetic ketoacidosis. (4, 5.1)
- Concomitant use with gemfibrozil. (4, 7.2, 12.3)

----------WARNINGS AND PRECAUTIONS----------

- Lactic acidosis: See boxed warning. (4, 5.1)
- Hypoglycemia: PRANDIMET may cause hypoglycemia. Skip the scheduled dose if a meal is skipped to reduce the risk of hypoglycemia. Reduce the dose if hypoglycemia occurs. (5.2)
- Vitamin B_{12} deficiency: Metformin can lower vitamin B_{12} levels. Monitor hematological parameters annually. (5.3)
- Serious Cardiovascular Adverse Reactions with Concomitant NPH-insulin: PRANDIMET is not indicated for use with NPH-insulin. (5.4)
- Macrovascular outcomes: There have been no clinical studies establishing conclusive evidence of macrovascular risk reduction with PRANDIMET. (5.5)

----------ADVERSE REACTIONS----------

- Hypoglycemia and headache were the most common adverse reactions (≥10%) reported among patients treated with repaglinide in combination with metformin. (6.1)
- Gastrointestinal reactions (e.g., diarrhea, nausea and vomiting) are the most common adverse reactions with metformin (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Novo Nordisk at 1-800-727-6500 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

----------DRUG INTERACTIONS----------

- Carbonic anhydrase inhibitors may increase risk of lactic acidosis. Consider more frequent monitoring. (7)
- Drugs that reduce metformin clearance (such as ranolazine, vandetanib, dolutegravir, and cimetidine) may increase the accumulation of metformin. Consider the benefits and risks of concomitant use. (7)
- Alcohol can potentiate the effect of metformin on lactate metabolism. Warn patients against excessive alcohol intake. (7)
- Clopidogrel: Avoid concomitant use; if used concomitantly limit total daily dose of repaglinide to 4 mg (7)
- Cyclosporine: Limit daily dose of repaglinide to 6 mg and increase frequency of glucose monitoring when co-administered (7)
- CYP2C8 and CYP3A4 Inhibitors and Drugs That May Increase the Risk of Hypoglycemia: Co-administration may require dose reductions and increased frequency of glucose monitoring (7)
- CYP2C8 and CYP3A4 Inducers and Drugs That May Decrease the Blood Glucose Lowering Effect of PRANDIMET: Co-administration may require dose increases and increased frequency of glucose monitoring (7)
- Drugs That May Blunt Signs and Symptoms of Hypoglycemia: Increased frequency of glucose monitoring may be required when co-administered (7)

----------USE IN SPECIFIC POPULATIONS----------

- Nursing mothers: Discontinue PRANDIN or nursing (8.3)
- Geriatric Use: Assess renal function more frequently. (8.5)
- Hepatic Impairment: Avoid use in patients with hepatic impairment. (8.7)

See 17 for PATIENT COUNSELING INFORMATION

Revised: 12/2017

FULL PRESCRIBING INFORMATION: CONTENTS*

*Sections or subsections omitted from the full prescribing information are not listed

FULL PRESCRIBING INFORMATION

WARNING: LACTIC ACIDOSIS

Post-marketing cases of metformin-associated lactic acidosis have resulted in death, hypothermia, hypotension, and resistant bradyarrhythmias. The onset of metformin-associated lactic acidosis is often subtle, accompanied only by nonspecific symptoms such as malaise, myalgias, respiratory distress, somnolence, and abdominal pain. Metformin-associated lactic acidosis was characterized by elevated blood lactate levels (> 5 mmol/Liter), anion gap acidosis (without evidence of ketonuria or ketonemia), an increased lactate/pyruvate ratio; and metformin plasma levels generally >5 mcg/mL *[see Warnings and Precautions (5.1)]*.

Risk factors for metformin-associated lactic acidosis include renal impairment, concomitant use of certain drugs (e.g., carbonic anyhydrase inhibitors such as topiramate), age 65 years old or greater, having a radiological study with contrast, surgery and other procedures, hypoxic states (e.g., acute congestive heart failure), excessive alcohol intake, and hepatic impairment.

Steps to reduce the risk of and manage metformin-associated lactic acidosis in these high risk groups are provided in the full prescribing information *[see Dosage and Administration (2.2), Contraindications (4), Warnings and Precautions (5.1), Drug Interactions (7), and Use in Specific Populations (8.6, 8.7)]*.

If metformin-associated lactic acidosis is suspected, immediately discontinue PRANDIMET and institute general supportive measures in a hospital setting. Prompt hemodialysis is recommended *[see Warnings and Precautions (5.1)]*.

1 INDICATIONS AND USAGE

PRANDIMET is indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus who are already treated with a glinide and metformin or who have inadequate glycemic control on a glinide alone or metformin alone.

Limitation of Use:

PRANDIMET should not be used in patients with type 1 diabetes mellitus or for the treatment of diabetic ketoacidosis.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage and Administration

General Dosage and Administration information

Administer PRANDIMET orally 2 to 3 times a day with meals up to a maximum daily dose of 10 mg repaglinide/2500 mg metformin. No more than 4 mg repaglinide/1000 mg metformin should be taken per meal.

Instruct patients to take PRANDIMET within 30 minutes before meals. In patients who skip meals, instruct patients to skip the scheduled dose of PRANDIMET to reduce the risk of hypogylcemia. In patients who experience hypoglycemia, the dose of PRANDIN should be reduced *[see Warnings and Precautions (5.2)]*.

Patients Inadequately Controlled with Metformin Monotherapy

The recommended starting dose of PRANDIMET 1 mg repaglinide/500 mg metformin twice daily with meals. Gradually increase dose based on glycemic response to reduce the risk of hypoglycemia with repaglinide.

Patients Inadequately Controlled with Glinide Monotherapy

The recommended starting dose of metformin component of PRANDIMET is 500 mg metformin twice daily with meals. Gradually increase dose based on glycemic response to reduce gastrointestinal side effects associated with metformin.

Patients Currently Using Repaglinide and Metformin Concomitantly

Initiate PRANDIMET at the dose of repaglinide and metformin similar to (but not exceeding) the patient's current doses. Titrate as needed to achieve glycemic control up to the maximum daily dose.

2.2 Recommended Dosage for Patients with Renal Impairment

Assess renal function with an estimated glomerular filtration rate (eGFR) prior to initiation of PRANDIMET and periodically thereafter.

PRANDIMET is contraindicated in patients with an eGFR below 30 mL/min/1.73 m2.

Initiation of PRANDIMET in patients with an eGFR between 30 – 45 mL/min/1.73 m2 is not recommended.

In patients taking PRANDIMET whose eGFR later falls below 45 mL/min/1.73 m2, assess the benefit-risk of continuing therapy.

Discontinue PRANDIMET if the patient's eGFR later falls below 30 mL/min/1.73 m2 *[see Contraindications (4) and Warnings and Precautions (5.1)]*.

2.3 Discontinuation for Iodinated Contrast Imaging Procedures

Discontinue PRANDIMET at the time of, or prior to, an iodinated contrast imaging procedure in patients with an eGFR between 30 and 60 mL/min/1.73 m2; in patients with a history of liver disease, alcoholism or heart failure; or in patients who will be administered intra-arterial iodinated contrast. Re-evaluate eGFR 48 hours after the imaging procedure; restart PRANDIMET if renal function is stable *[See Warnings and Precautions (5.1)]*.

2.4 Dose Modifications for Drug Interactions

Concomitant use with gemfibrozil is contraindicated *[see Contraindications (4)]*.

Avoid concomitant use of PRANDIMET with clopidogrel. If concomitant use can not be avoided, initiate repaglinide at 0.5 mg before each meal. Although PRANDIMET is not available in that strength, repaglinide 0.5 mg tablets are available. Do not exceed a total daily dose of 4 mg of repaglinide *[see Drug Interactions (7), Clinical Pharmacology (12.3)]*.

Do not exceed a total daily dose of 6 mg of repaglinide in patients receiving cyclosporine *[see Drug Interactions (7), Clinical Pharmacology (12.3)]*.

Dosage adjustments are recommended in patients taking concomitant strong CYP3A4 or CYP2C8 inhibitors or strong CYP3A4 or CYP2C8 inducers *[see Drug Interactions (7), Clinical Pharmacology (12.3)]*.

3 DOSAGE FORMS AND STRENGTHS

- 1 mg repaglinide/500mg metformin tablets (yellow, biconvex, debossed with Novo Nordisk (Apis) bull symbol on one side, and 1/500 on the other side)
- 2 mg repaglinide/500mg metformin tablets (pink, biconvex, debossed with Novo Nordisk (Apis) bull symbol on one side, and 2/500 on the other side)

4 CONTRAINDICATIONS

PRANDIMET is contraindicated in patients with:

- Severe renal impairment (eGFR below 30 mL/min/1.73 m^2) *[see Warnings and Precautions (5.1)]*
- Acute or chronic metabolic acidosis, including diabetic ketoacidosis. Diabetic ketoacidosis should be treated with insulin *[see Warnings and Precautions (5.1)]*.
- Concomitant use of gemfibrozil *[see Drug Interactions (7.2)]*
- Known hypersensitivity to repaglinide, metformin or any inactive ingredients

5 WARNINGS AND PRECAUTIONS

5.1 Lactic Acidosis

There have been post-marketing cases of metformin-associated lactic acidosis, including fatal cases. These cases had a subtle onset and were accompanied by nonspecific symptoms such as malaise, myalgias, abdominal pain, respiratory distress, or increased somnolence; however, hypothermia, hypotension and resistant bradyarrhythmias have occurred with severe acidosis.

Metformin-associated lactic acidosis was characterized by elevated blood lactate concentrations (>5 mmol/Liter), anion gap acidosis (without evidence of ketonuria or ketonemia), and an increased lactate:pyruvate ratio; metformin plasma levels generally >5 mcg/mL. Metformin decreases liver uptake of lactate increasing lactate blood levels which may increase the risk of lactic acidosis, especially in patients at risk.

If metformin-associated lactic acidosis is suspected, general supportive measures should be instituted promptly in a hospital setting, along with immediate discontinuation of PRANDIMET.

In PRANDIMET treated patients with a diagnosis or strong suspicion of lactic acidosis, prompt hemodialysis is recommended to correct the acidosis and remove accumulated metformin (metformin is dialyzable, with a clearance of up to 170 mL/minute under good hemodynamic conditions). Hemodialysis has often resulted in reversal of symptoms and recovery.

Educate patients and their families about the symptoms of lactic acidosis and if these symptoms occur instruct them to discontinue PRANDIMET and report these symptoms to their healthcare provider.

For each of the known and possible risk factors for metformin-associated lactic acidosis, recommendations to reduce the risk of and manage metformin-associated lactic acidosis are provided below:

Renal Impairment: The post-marketing metformin-associated lactic acidosis cases primarily occurred in patients with significant renal impairment. The risk of metformin accumulation and metformin-associated lactic acidosis increases with the severity of renal impairment because metformin is substantially excreted by the kidney. Clinical recommendations based upon the patient's renal function include *[see Clinical Pharmacology (12.3)]:*

- Before initiating PRANDIMET, obtain an estimated glomerular filtration rate (eGFR).
- PRANDIMET is contraindicated in patients with an eGFR less than 30 mL/minute/1.73 m2*[see Contraindiations (4)]* .
- Initiation of PRANDIMET is not recommended in patients with eGFR between 30 – 45 mL/minute/1.73 m2.
- Obtain an eGFR at least annually in all patients taking PRANDIMET. In patients at increased risk for the development of renal impairment (e.g., the elderly), renal function should be assessed more frequently.
- In patients taking PRANDIMET whose eGFR later falls below 45 mL/min/1.73 m2, assess the benefit and risk of continuing therapy.

Drug Interactions: The concomitant use of PRANDIMETwith specific drugs may increase the risk of metformin- associated lactic acidosis: those that impair renal function, result in significant hemodynamic change, interfere with acid-base balance or increase metformin accumulation *[see Drug Interactions (7)]*. Therefore, consider more frequent monitoring of patients.

Age 65 or Greater: The risk of metformin-associated lactic acidosis increases with the patient's age because elderly patients have a greater likelihood of having hepatic, renal, or cardiac impairment than younger patients. Assess renal function more frequently in elderly patients [*see Use in Specific Populations (8.5)]*.

Radiological Studies with Contrast: Administration of intravascular iodinated contrast agents in metformin-treated patients has led to an acute decrease in renal function and the occurrence of lactic acidosis. Stop PRANDIMET at the time of, or prior to, an iodinated contrast imaging procedure in patients with an eGFR between 30 and 60 mL/min/1.73 m2; in patients with a history of hepatic impairment, alcoholism, or heart failure; or in patients who will be administered intra-arterial iodinated contrast. Re-evaluate eGFR 48 hours after the imaging procedure, and restart PRANDIMET if renal function is stable.

Surgery and Other Procedures: Withholding of food and fluids during surgical or other procedures may increase the risk for volume depletion, hypotension and renal impairment. PRANDIMET should be temporarily discontinued while patients have restricted food and fluid intake.

Hypoxic States: Several of the postmarketing cases of metformin-associated lactic acidosis occurred in the setting of acute congestive heart failure (particularly when accompanied by hypoperfusion and hypoxemia). Cardiovascular collapse (shock), acute myocardial infarction, sepsis, and other conditions associated with hypoxemia have been associated with lactic acidosis and may also cause prerenal azotemia. When such events occur, discontinue PRANDIMET.

Excessive Alcohol Intake: Alcohol potentiates the effect of metformin on lactate metabolism and this may increase the risk of metformin-associated lactic acidosis. Warn patients against excessive alcohol intake while receiving PRANDIMET.

Hepatic Impairment: Patients with hepatic impairment have developed with cases of metformin-associated lactic acidosis. This may be due to impaired lactate clearance resulting in higher lactate blood levels. Therefore, avoid use of PRANDIMET in patients with clinical or laboratory evidence of hepatic disease.

5.2 Hypoglycemia

All glinides, including PRANDIMET, can cause hypoglycemia *[see Adverse Reactions (6.1)]*. Severe hypoglycemia can cause seizures, may be life-threatening, or cause death. Hypoglycemia can impair concentration ability and reaction time; this may place an individual and others at risk in situations where these abilities are important (e.g., driving or operating other machinery).

Hypoglycemia can happen suddenly and symptoms may differ in each individual and change over time in the same individual. Symptomatic awareness of hypoglycemia may be less pronounced in patients with longstanding diabetes, in patients with diabetic nerve disease, in patients using medications that block the sympathetic nervous system (e.g., beta-blockers) *[see Drug Interactions (7)]*, or in patients who experience recurrent hypoglycemia.

Factors which may increase the risk of hypoglycemia include changes in meal pattern (e.g., macronutrient content), changes in level of physical activity, changes to co-administered medication *[see Drug Interactions (7)]*, and concomitant use with other antidiabetic agents. Patients with renal or hepatic impairment may be at higher risk of hypoglycemia *[see Use in Specific Populations (8.6, 8.7)]*.

Patients should administer PRANDIMET before meals and be instructed to skip the dose of PRANDIMET if a meal is skipped. In patients who experience hypoglycemia, the dose of PRANDIMET should be reduced *[see Dosage and Administration (2.1)]*. Patients and caregivers must be educated to recognize and manage hypoglycemia. Self-monitoring of blood glucose plays an essential role in the prevention and management of hypoglycemia. In patients at higher risk for hypoglycemia and patients who have reduced symptomatic awareness of hypoglycemia, increased frequency of blood glucose monitoring is recommended.

5.3 Vitamin B_{12} Levels

In controlled clinical trials of metformin of 29 weeks' duration, a decrease to subnormal levels of previously normal serum vitamin B_{12} levels, without clinical manifestations, was observed in approximately 7% of patients. This finding, possibly due to interference with B_{12} absorption from the B_{12}-intrinsic factor complex, is rarely associated with anemia and appears to be rapidly reversible with discontinuation of metformin or vitamin B_{12} supplementation. Measurement of hematologic parameters on an annual basis is advised in patients on PRANDIMET and any apparent abnormalities should be appropriately investigated and managed.

Certain individuals (those with inadequate vitamin B_{12} or calcium intake or absorption) appear to be predisposed to developing subnormal vitamin B_{12} levels. In these patients, routine serum vitamin B_{12} measurements at 2- to 3-year intervals may be useful.

5.4 Serious Cardiovascular Adverse Reactions with Concomitant Use with NPH-insulin

Across seven controlled trials, there were six serious adverse events of myocardial ischemia in patients treated with repaglanide plus NPH-insulin from two studies, and one event in patients using insulin formulations alone from another study *[See Adverse Reactions (6.1)]*. PRANDIMET is not indicated for use in combination with NPH-insulin.

5.5 Macrovascular Outcomes

There have been no clinical studies establishing conclusive evidence of macrovascular risk reduction with PRANDIMET.

6 ADVERSE REACTIONS

The following adverse reactions are discussed in more detail in other sections of the labeling.

- Lactic acidosis *[see Warnings and Precautions (5.1)]*
- Hypoglycemia *[see Warnings and Precautions (5.2)]*
- Vitamin B_{12} levels *[see Warnings and Precautions (5.3)]*

6.1 Clinical Trial Experience

Because clinical trials are conducted under varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in clinical practice.

PRANDIMET was administered to 374 patients with type 2 diabetes during clinical trials. Table 1 summarizes the most common adverse reactions occurring in a 6-month randomized trial of repaglinide as add-on therapy to metformin in patients inadequately controlled on metformin alone.

Table 1: Adverse Reactions (%) occurring ≥10% in Patients Treated with Repaglinide and Metformin Together or Repaglinide and Metformin Monotherapy during a 6 Month Trial			
	Repaglinide and Metformin	Metformin Monotherapy	Repaglinide Monotherapy
	N = 27	N = 27	N = 28
Diarrhea Nausea	19 15	30 7	7 4
Symptomatic Hypoglycemia*	33	0	11
Headache	22	15	11
Upper Respiratory Tract Infection	11	11	11
* Hypoglycemia with symptoms which included, but were not limited to, anxious feeling, dizziness, sweating, tremor, hunger and difficulty in concentration. None of the symptomatic hypoglycemia events listed in the table required the assistance of another person.			

Hypoglycemia

In clinical trials with repaglinide, hypoglycemia is the most commonly observed adverse reaction. Mild or moderate hypoglycemia occurred in 31% of repaglinide treated patients and 7% of placebo treated patients.

Gastrointestinal Adverse Reactions

Gastrointestinal reactions (e.g., diarrhea, nausea, vomiting) are the most common adverse reactions (> 5%) with metformin treatment and are more frequent at higher metformin doses.

Weight Gain

In a clinical trial, 83 patients were randomized to add-on repaglinide, repaglinide monotherapy, or continued treatment with metformin monotherapy. A statistically significant weight gain was observed for the combination of repaglinide and metformin in a pairwise comparison with metformin monotherapy (see Table 2).

Table 2: Repaglinide as Add-on to Metformin: Mean Changes from Baseline in Body Weight After 4 to 5 Months of Treatment[1]			
	Repaglinide add-on to Metformin	Repaglinide monotherapy	Metformin HCl monotherapy
N	27	28	27
Weight (kg)			
Baseline	93	87	91
Change from Baseline	2.4#	3.0	-0.9
[1]: based on intent-to-treat analysis #: $p < 0.05$, for pairwise comparison with metformin monotherapy.			

Cardiovascular Events in repaglinide monotherapy trials

The incidence of total serious cardiovascular adverse events, including ischemia, was higher for repaglinide (51/1228 or 4%) than for sulfonylurea drugs (13/498 or 3%) in controlled clinical trials.

Seven controlled clinical trials included repaglinide combination therapy with NPH-insulin (n=431), insulin formulations alone (n=388) or other combinations (sulfonylurea plus NPH-insulin or repaglinide plus metformin) (n=120). There were six serious adverse events of myocardial ischemia in patients treated with repaglinide plus NPH-insulin (1.4%) from two studies, and one event in patients using insulin formulations alone from another study (0.3%) *[see Warnings and Precautions (5.4)]*.

6.2 Postmarketing Experience

The following additional adverse reactions have been identified during postapproval use. Because these reactions are reported voluntarily from a population of uncertain size, it is generally not possible to reliably estimate their frequency or a causal relationship to drug exposure.

Repaglinide

Alopecia, hemolytic anemia, pancreatitis, Stevens-Johnson Syndrome, and severe hepatic dysfunction including jaundice and hepatitis.

Metformin

Cholestatic, hepatocellular, and mixed hepatocellular liver injury

7 DRUG INTERACTIONS

Table 3 includes a list of drugs with clinically important drug interactions when administered concomitantly with PRANDIMET and instructions for preventing or managing them.

Table 3: Clinically Important Drug Interactions with PRANDIMET

Carbonic Anhydrase Inhibitors	
Clinical Impact:	Carbonic anhydrase inhibitors frequently causes a decrease in serum bicarbonate and induce non-anion gap, hyperchloremic metabolic acidosis. Concomitant use of these drugs with PRANDIMET may increase the risk for lactic acidosis.
Intervention:	Consider more frequent monitoring of these patients.
Examples:	Topiramate, zonisamide, acetazolamide or dichlorphenamide.
Drugs that reduce metformin clearance	
Clinical Impact:	Concomitant use of drugs that interfere with common renal tubular transport systems involved in the renal elimination of metformin could increase systemic exposure to metformin and may increase the risk for lactic acidosis *[see Clinical Pharmacology (12.3)]*.
Intervention:	Consider the benefits and risks of concomitant use.
Examples:	E.g., organic cationic transporter-2 [OCT2] / multidrug and toxin extrusion [MATE] inhibitors such as ranolazine, vandetanib, dolutegravir, and cimetidine.
Alcohol	
Clinical Impact:	Alcohol is known to potentiate the effect of metformin on lactate metabolism.
Intervention:	Warn patients against excessive alcohol intake while receiving PRANDIMET.
Gemfibrozil	
Clinical Impact:	Gemfibrozil significantly increased repaglinide exposures by 8.1 fold *[see Clinical Pharmacology (12.3)]*
Intervention:	Do not administer PRANDIMET to patients receiving gemfibrozil *[see Contraindications (4)]*.
Clopidogrel	
Clinical Impact:	Clopidogrel increased repaglinide exposures by 3.9-5.1 fold *[see Clinical Pharmacology (12.3)]*
Intervention:	Avoid concomitant use of PRANDIMET with clopidogrel. If concomitant use can not be avoided, initiate repaglinide at 0.5 mg before each meal. Although PRANDIMET is not available in that strength, repaglinide 0.5 mg tablets are available. Do not exceed a total daily repaglinide dose of 4 mg. Increased frequency of glucose monitoring may be required during concomitant use *[see Dosage and Administration (2.4)]*.
Cyclosporine	
Clinical Impact:	Cyclosporine increased low dose repaglinide exposures by 2.5 fold *[see Clinical Pharmacology (12.3)]*
Intervention:	Daily maximum repaglinide dose should be limited to 6 mg, and increased frequency of glucose monitoring may be required when PRANDIMET is co-administered with cyclosporine.
CYP2C8 and CYP3A4 Inhibitors	
Intervention:	PRANDIMET dose reductions and increased frequency of glucose monitoring may be required when co-administered.
Examples:	Drugs that are known to inhibit CYP3A4 include antifungal agents (ketoconazole, itraconazole) and antibacterial agents (clarithromycin, erythromycin). Drugs that are known to inhibit CYP2C8 include trimethoprim, gemfibrozil, montelukast, deferasirox, and clopidiogrel.
CYP2C8 and CYP3A4 Inducers	
Intervention:	PRANDIMET dose increases and increased frequency of glucose monitoring may be required when co-administered.
Examples:	Drugs that induce the CYP3A4 and/or 2C8 enzyme systems include rifampin, barbiturates, and carbamezapine
Drugs That May Increase the Risk of Hypoglycemia	
Intervention:	PRANDIMET dose reductions and increased frequency of glucose monitoring may be required when co-administered.
Examples:	Antidiabetic agents, ACE inhibitors, angiotensin II receptor blocking agents, disopyramide, fibrates, fluoxetine, monoamine oxidase inhibitors, nonsteroidal anti-inflammatory agents (NSAIDs), pentoxifylline, pramlintide, propoxyphene, salicylates, somatostatin analogs (e.g., octreotide), and sulfonamide antibiotics
Drugs That May Decrease the Blood Glucose Lowering Effect of PRANDIMET	
Intervention:	PRANDIMET dose increases and increased frequency of glucose monitoring may be required when co-administered.
Examples:	Atypical antipsychotics (e.g., olanzapine and clozapine), calcium channel antagonists, corticosteroids, danazol, diuretics, estrogens, glucagon, isoniazid, niacin, oral contraceptives, phenothiazines, progestogens (e.g., in oral contraceptives), protease inhibitors, somatropin, sympathomimetic agents (e.g., albuterol, epinephrine, terbutaline), and thyroid hormones.
Drugs That May Blunt Signs and Symptoms of Hypoglycemia	
Intervention:	Increased frequency of glucose monitoring may be required when PRANDIMET is co-administered with these drugs.
Examples:	beta-blockers, clonidine, guanethidine, and reserpine

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Pregnancy Category C.

There are no adequate and well-controlled studies in pregnant women with PRANDIMET or its individual components. Because animal reproduction studies are not always predictive of human response, PRANDIMET like other antidiabetic medications, should be used during pregnancy only if clearly needed.

No animal studies have been conducted with the combined products in PRANDIMET. The following data are based on findings in studies performed with repaglinide or metformin individually.

Repaglinide

Repaglinide was not teratogenic in rats at doses 40 times, and rabbits approximately 0.8 times the clinical exposure (on a mg/m^2 basis) throughout pregnancy. Offspring of rat dams exposed to repaglinide at 15 times clinical exposure on a mg/m^2 basis during days 17 to 22 of gestation and during lactation developed nonteratogenic skeletal deformities consisting of shortening, thickening, and bending of the humerus during the postnatal period. This effect was not seen at doses up to 2.5 times clinical exposure (on a mg/m^2 basis) on days 1 to 22 of pregnancy or at higher doses given during days 1 to 16 of pregnancy. Relevant human exposure has not occurred to date and therefore the safety of repaglinide administration throughout pregnancy or lactation cannot be established.

Metformin

Metformin alone was not teratogenic in rats or rabbits at doses up to 600 mg/kg/day. This represents an exposure of approximately two and six times the near-maximal efficacious human daily dose of 2000 mg of the metformin component of PRANDIMET based on body surface area comparisons for rats and rabbits, respectively. Determination of fetal concentrations demonstrated a partial placental barrier to metformin.

8.3 Nursing Mothers

No studies in lactating animals have been conducted with the PRANDIMET fixed dose combination. In studies performed with individual components, both repaglinide and metformin are excreted into milk of lactating rats. It is not known whether repaglinide or metformin are excreted in human milk. PRANDIMET is not recommended in nursing mothers because it may potentially cause hypoglycemia in nursing infants.

Repaglinide

In rat reproduction studies, measurable levels of repaglinide were detected in the breast milk of the dams and lowered blood glucose levels were observed in the pups. Cross fostering studies indicated that skeletal changes could be induced in control pups nursed by treated dams, although this occurred to a lesser degree than those pups treated in utero.

Metformin

Studies in lactating rats with metformin show that it is excreted into milk and reaches levels comparable to those in plasma.

8.4 Pediatric Use

Safety and effectiveness have not been established in pediatric patients.

8.5 Geriatric Use

In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy and the higher risk of lactic acidosis. Assess renal function more frequently in elderly patients. *[see Warnings and Precautions (5.1), Contraindications (4), Clinical Pharmacology (12.3)]*

8.6 Renal Impairment

Metformin is substantially excreted by the kidney, and the risk of metformin accumulation and lactic acidosis increases with the degree of renal impairment. PRANDIMET is contraindicated in severe renal impairment, patients with an estimated glomerular filtration rate (eGFR) below 30 mL/min/1.73 m2. *[See Dosage and Administration (2.2), Contraindications (4), Warnings and Precautions (5.1) and Clinical Pharmacology (12.3)]*

8.7 Hepatic Impairment

Use of metformin in patients with hepatic impairment has been associated with some cases of lactic acidosis. PRANDIMET is not recommended in patients with hepatic impairment. *[See Warnings and Precautions (5.1)]*

10 OVERDOSAGE

Repaglinide

Severe hypoglycemic reactions with coma, seizure, or other neurological impairment may occur and constitute medical emergencies requiring immediate hospitalization. Hypoglycemic symptoms without loss of consciousness or neurologic findings should be treated aggressively with oral glucose and adjustments in drug dosage and/or meal patterns. Close monitoring may continue until the physician is assured that the patient is out of danger. Patients should be closely monitored for a minimum of 24 to 48 hours, since hypoglycemia may recur after apparent clinical recovery. There is no evidence that repaglinide is dialyzable using hemodialysis.

Metformin

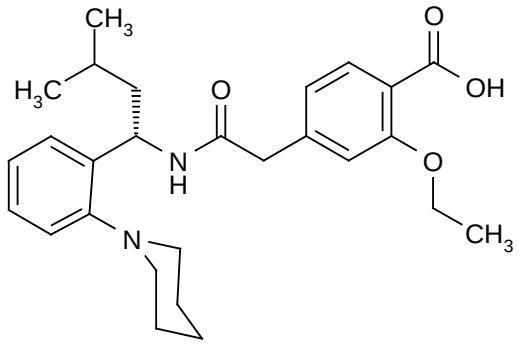
Overdose of metformin has occurred, including ingestion of amounts greater than 50 grams. Hypoglycemia was reported in approximately 10% of cases, but no causal association with metformin has been established. Lactic acidosis has been reported in approximately 32% of metformin overdose cases *[see Warnings and Precautions (5.1)]*. Metformin is dialyzable with a clearance of up to 170 mL/min under good hemodynamic conditions. Therefore, hemodialysis may be useful for removal of accumulated drug from patients in whom metformin overdosage is suspected.

11 DESCRIPTION

PRANDIMET (repaglinide and metformin hydrochloride) tablets for oral use contain repaglinide, a glinide, and metformin, a biguanide.

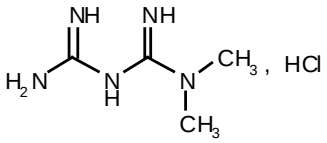
Repaglinide, S(+)2-ethoxy-4(2((3-methyl-1-(2-(1-piperidinyl) phenyl)-butyl) amino)-2-oxoethyl) benzoic acid, is chemically unrelated to the oral sulfonylurea insulin secretagogues. Repaglinide is a white to off-white powder with molecular formula C_{27} H_{36} N_2 O_4 and a molecular weight of 452.6 with the structural formula as shown below. Repaglinide is freely soluble in methanol and ethanol. The pKa of repaglinide in acid is 3.9, and the pKa in amine is 6.0.

Structural formula of Repaglinide



Metformin hydrochloride (N,N-dimethylimidodicarbonimidic diamide hydrochloride) is a white to off-white crystalline compound with a molecular formula of $C_4H_{11}N_5$•HCl and a molecular weight of 165.63. Metformin HCl is freely soluble in water and is practically insoluble in acetone, ether, and chloroform. The pKa of metformin hydrochloride is 12.4. The pH of a 1% aqueous solution of metformin hydrochloride is 6.68. The structural formula of metformin hydrochloride is:

Structural formula of Metformin Hydrochloride



PRANDIMET tablets contain 1 mg repaglinide with 500 mg metformin HCl (1 mg/500 mg) or 2 mg repaglinide with 500 mg metformin HCl (2 mg/500 mg) formulated with the following inactive ingredients: poloxamer 188**,** microcrystalline cellulose, polacrillin potassium, magnesium stearate, hypromellose 3cp or 6cp, povidone, meglumine, sorbitol, talc, titanium dioxide, red or yellow iron oxide, and polyethylene glycol. Propylene glycol is present in the 2 mg/500 mg PRANDIMET tablets.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

PRANDIMET

Repaglinide lowers blood glucose levels by stimulating the release of insulin from the pancreas. This action is dependent upon functioning beta (ß) cells in the pancreatic islets.

Repaglinide closes ATP-dependent potassium channels in the ß-cell membrane by binding at characterizable sites. This potassium channel blockade depolarizes the ß-cell, which leads to an opening of calcium channels. The resulting increased calcium influx induces insulin secretion. The ion channel mechanism is highly tissue selective with low affinity for heart and skeletal muscle.

Metformin improves glucose tolerance in patients with type 2 diabetes by lowering both the basal and postprandial plasma glucose. Metformin decreases hepatic glucose production, decreases intestinal absorption of glucose, and improves insulin sensitivity by increasing peripheral glucose uptake and utilization. With metformin therapy, insulin secretion remains unchanged while fasting insulin levels and day-long plasma insulin response may actually decrease.

12.3 Pharmacokinetics

PRANDIMET

The pharmacokinetic profiles of repaglinide and metformin from PRANDIMET doses are listed in Table 4 below. The results of a pharmacokinetic single-dose crossover study in healthy subjects demonstrated that repaglinide has dose proportional pharmacokinetics (AUC and C_{max}) for the combination of repaglinide/metformin in PRANDIMET (2 mg/500 mg and 1 mg/500 mg).

Table 4: Pharmacokinetic Parameters for Repaglinide and Metformin

		Repaglinide		Metformin	
PRANDIMET	N	AUC (ng·h/mL)	C_{max} (ng/mL)	AUC (ng·h/mL)	C_{max} (ng/mL)
2 mg/500 mg tablet	55	34.5 (13.3)	26.0 (13.7)	6041.9 (1494.6)	838.8 (210.2)
1 mg/500 mg tablet	55	17.6 (6.6)	12.9 (6.9)	5948.9 (1442.0)	799.4 (174.6)

Absorption and Bioavailability

Repaglinide: After single and multiple oral doses in healthy subjects or in patients with type 2 diabetes, peak plasma drug levels (C_{max}) occur within 1 hour (T_{max}). Repaglinide is eliminated from the blood stream with a half-life of approximately 1 hour. The mean absolute bioavailability is 56%. When repaglinide was given with food, the mean T_{max} was not changed, but the mean C_{max} and AUC (area under the time/plasma concentration curve) were decreased 20% and 12.4%, respectively.

Metformin: The absolute bioavailability of a 500 mg metformin tablet given under fasting conditions is approximately 50% to 60%. Studies using single oral doses of metformin tablets of 500 mg to 1,500 mg, and 850 mg to 2,550 mg, indicate that there is a lack of dose proportionality with increasing doses, which is due to decreased absorption rather than an alteration in elimination. Food decreases the extent of and slightly delays the absorption of metformin, as shown by approximately a 40% lower peak concentration (C_{max}), a 25% lower area under plasma concentration (AUC) and a 35-minute prolongation of time to peak plasma concentration (T_{max}) following administration of a single 850 mg tablet of metformin with food, compared to the same tablet strength administered fasting. The clinical relevance of these decreases is unknown.

Distribution

Repaglinide: After intravenous (IV) dosing in healthy subjects, the volume of distribution at steady state (Vss) was 31 L, and the total body clearance (CL) was 38 L/h. Protein binding and binding to human serum albumin was greater than 98%.

Metformin: The apparent volume of distribution (V/F) of metformin following single oral dose of 850 mg averaged 654 ± 358 L. Metformin is negligibly bound to plasma proteins. Metformin partitions into erythrocytes, most likely as a function of time. At usual clinical doses and dosing schedules of metformin, steady state plasma concentrations of metformin are reached within 24-48 hours and are generally < 1 µg/mL. During controlled clinical trials, maximum metformin plasma levels did not exceed 5 µg/mL, even at maximum doses.

Metabolism and Elimination

Repaglinide: Repaglinide is completely metabolized by oxidative biotransformation and direct conjugation with glucuronic acid after either an intravenous or oral dose. The major metabolites are an oxidized dicarboxylic acid (M2), the aromatic amine (M1), and the acyl glucuronide (M7). The cytochrome P-450 enzyme system, specifically 2C8 and 3A4, have been shown to be involved in the N-dealkylation of repaglinide to M2 and the further oxidation to M1. Metabolites do not contribute to the glucose-lowering effect of repaglinide. Within 96 hours after dosing with ^{14}C-repaglinide as a single, oral dose, approximately 90% of the radiolabel was recovered in the feces and approximately 8% in the urine. Only 0.1% of the dose is cleared in the urine as parent compound. The major metabolite (M2) accounted for 60% of the administered dose. Less than 2% of parent drug was recovered in feces. Repaglinide appears to be a substrate for active hepatic uptake transporter (organic anion transporting protein OATP1B1).

Metformin: Intravenous single-dose studies in normal subjects demonstrate that metformin is excreted unchanged in the urine and does not undergo hepatic metabolism (no metabolites have been identified in humans) or biliary excretion. Renal clearance is approximately 3.5 times greater than creatinine clearance, which indicates that tubular secretion is the major route of metformin elimination. Following oral administration, approximately 90% of the absorbed drug is eliminated via the renal route within the first 24 hours, with a plasma elimination half-life of approximately 6.2 hours. In blood, the elimination half-life is approximately 17.6 hours, suggesting that the erythrocyte mass may be a compartment of distribution.

Specific Populations

Renal Impairment

Repaglinide

Single-dose and steady-state pharmacokinetics of repaglinide were compared between patients with type 2 diabetes and normal renal function (CrCl > 80 mL/min), mild to moderate renal function impairment (CrCl = 40 – 80 mL/min), and severe renal function impairment (CrCl = 20 – 40 mL/min). Both AUC and C_{max} of repaglinide were similar in patients with normal and mild to moderately impaired renal function (mean values 56.7 ng/mL*hr vs 57.2 ng/mL*hr and 37.5 ng/mL vs 37.7 ng/mL, respectively). Patients with severely reduced renal function had elevated mean AUC and C_{max} values (98.0 ng/mL*hr and 50.7 ng/mL, respectively), but this study showed only a weak correlation between repaglinide levels and creatinine clearance.

Metformin

In patients with decreased renal function , the plasma and blood half-life of metformin is prolonged and the renal clearance is decreased *[see Contraindications (4) and Warnings and Precautions (5.1)]*.

Hepatic Impairment

Repaglinide

A single-dose, open-label study was conducted in 12 healthy subjects and 12 patients with chronic liver disease (CLD) classified by Child-Pugh scale and caffeine clearance. Patients with moderate to severe impairment of liver function had higher and more prolonged serum concentrations of both total and unbound repaglinide than healthy subjects ($AUC_{healthy}$: 91.6 ng/mL*hr; $AUC_{CLD\ patients}$: 368.9 ng/mL*hr; $C_{max,\ healthy}$: 46.7 ng/mL; $C_{max,\ CLD\ patients}$: 105.4 ng/mL). AUC was statistically correlated with caffeine clearance. No difference in glucose profiles was observed across patient groups.

Metformin

No pharmacokinetics studies with metformin have been conducted in patients with hepatic impairment.

Geriatric Patients

Healthy volunteers treated with repaglinide 2 mg before each of 3 meals, showed no significant differences in repaglinide pharmacokinetics between the group of patients <65 years of age and those ≥65 years of age.

Limited data from controlled pharmacokinetic studies of metformin in healthy elderly subjects suggest that total plasma clearance is decreased, the half-life is prolonged, and C_{max} is increased, compared to healthy young subjects. From these data, it appears that the change in metformin pharmacokinetics with aging is primarily accounted for by a change in renal function *[see Warnings and Precautions (5.1)]*.

Drug Interactions

Table 5: Effect of Other Drugs on AUC and C_{max} of Metformin

Study Drug*	Metformin AUC	Metformin C_{max}
Cimetidine	40% ↑	60% ↑
Furosemide	15% ↑	22% ↑
Nifedipine	9% ↑	20% ↑
Propranolol-metformin	10% ↓	6% ↓
Ibuprofen-metformin	5% ↑	7% ↑

Unless indicated all drug interactions were observed with single dose co-administration

*single and multiple dose co-administration

↑ indicates increase

↓ indicates decrease

Table 6: Effect of Other Drugs on AUC and C_{max} of Repaglinide

Study Drug	Dosing	Repaglinide Dosing[1]	Repaglinide	
			AUC	C_{max}
Clarithromycin*	250 mg BID for 4 days		40% ↑	67% ↑
Clopidogrel*	300 mg (Day 1) 75 mg QD (Day 2-3)	0.25 mg (Day 1 and 3)	(day 1) 5.1 fold ↑ (3.9-6.6) (day 3) 3.9 fold ↑ (2.9-5.3)	2.5 fold ↑ (1.8-3.5) 2.0 fold ↑ (1.3-3.1)
Cyclosporine	100 mg (2 doses 12 hours apart)		2.5 fold ↑	1.8 fold ↑
Deferasirox*	30 mg/kg QD for 4 days	0.5 mg	2.3 fold ↑	62% ↑
Fenofibrate	200 mg QD for 5 days		0%	0%
Gemfibrozil*	600 mg BID for 3 days		8.1 fold ↑	2.4 fold ↑
Itraconazole*	100 mg BID for 3 days		1.4 fold ↑	1.5 fold ↑
Gemfibrozil + **Itraconazole*** Co-administration	Gem: 600 mg BID for 3 days Itra: 100 mg BID for 3 days		19 fold ↑	2.8 fold ↑
Ketoconazole	200 mg QD for 4 days	2 mg	15% ↑	16% ↑
Levonorgestrel/ethinyl Estradiol	(0.15 mg/0.03 mg) Combination tablet QD for 21 days	2 mg	0%	20% ↑
Nifedipine*	10 mg TID for 4 days	2 mg	0%	0%
Rifampin*	600 mg QD for 6-7 days	4 mg	32 – 80% ↓	17 - 79% ↓
Simvastatin	20 mg QD for 4 days	2 mg	0%	26% ↑
Trimethoprim*	160 mg BID for 2 days 160 mg QD for 1 day		61% ↑	41% ↑

[1] Unless indicated all drug interactions were observed with single dose of 0.25 mg repaglinide
↑ indicates increase
↓ indicates decrease
* Indicates data are from published literature

Table 7: Effect of Metformin or Repaglinide on AUC and C_{max} of Other Drugs

Other Drugs	AUC	C_{max}
Furosemide [1]	12% ↓	31% ↓
Ethinyl Estradiol [2]	20% ↑	20% ↑
Fenofibrate	0%	18% ↑

[1] When administered with metformin
[2] Co-administration of a combination tablet (0.15 mg levonorgestrel/0.03 mg ethinyl estradiol) once daily for 21 days with 2 mg repaglinide administered TID (days 1-4) and a single dose on day 5.
↓ indicates decrease
↑ indicates increase

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

PRANDIMET

No animal studies have been conducted with the combined products in PRANDIMET to evaluate carcinogenesis, mutagenesis and impairment of fertility. The following data are based on findings in studies performed with the individual components.

Repaglinide

In a 104-week carcinogenicity study in rats at doses up to 120 mg/kg/day, the incidences of benign adenomas of the thyroid and liver were increased in male rats. The higher incidences of thyroid and liver tumors in male rats were not seen at lower dose of 30 mg/kg/day and 60 mg/kg/day respectively (which are over 15 and 30 times, respectively, clinical exposures on a mg/m^2 basis).
In a 104-week carcinogenicity study in mice at doses up to 500 mg/kg/day, no evidence of carcinogenicity was found in mice (which is approximately 125 times clinical exposure on a mg/m^2 basis).
Repaglinide was non-genotoxic in a battery of *in vivo* and *in vitro* studies: Bacterial mutagenesis (Ames test), *in vitro* forward cell mutation assay in V79 cells (HGPRT), *in vitro* chromosomal aberration assay in human lymphocytes, unscheduled and replicating DNA synthesis in rat liver, and *in vivo* mouse and rat micronucleus tests.
In a rat fertility study, repaglinide was administered to male and female rats at doses up to 300 and 80 mg/kg/day, respectively. No adverse effects on fertility were observed (which are over 40 times clinical exposure on a mg/m^2 basis).

Metformin

In a 104-week carcinogenicity study in rats at doses up to 900 mg/kg/day, the incidences of benign stromal uterine polyps were increased in female rats at 900 mg/kg/day (which is approximately four times the maximal recommended human daily dose of 2000 mg of metformin component of PRANDIMET on a mg/m^2 basis).
In a 91-week carcinogenicity study in mice at doses up to 1500 mg/kg/day, no evidence of carcinogenicity was found in mice (which is approximately four times the maximal recommended human daily dose of 2000 mg of metformin component of PRANDIMET on a mg/m^2 basis).
There was no evidence of a mutagenic potential of metformin alone in the following *in vitro* tests: Ames test (*S. typhimurium*), gene mutation test (mouse lymphoma cells), or chromosomal aberrations test (human lymphocytes). Results in the *in vivo* mouse micronucleus test were also negative.
In a rat fertility study, metformin was administered to male and female rats at doses up to 600 mg/kg/day. No adverse effects on fertility were observed (which is approximately three times the maximal recommended human daily dose of 2000 mg of metformin component of PRANDIMET on a mg/m^2 basis).

14 CLINICAL STUDIES

There have been no clinical efficacy studies conducted with PRANDIMET; however, bioequivalence of PRANDIMET to repaglinide and metformin co-administered as individual tablets was demonstrated in healthy subjects.

14.1 Repaglinide as Add-on Combination Therapy With Metformin

A total of 83 patients with type 2 diabetes and inadequate glycemic control on metformin monotherapy were randomized to repaglinide as add-on therapy to metformin, repaglinide monotherapy, or continued treatment with metformin monotherapy. The repaglinide dosage was titrated for 4 to 8 weeks, followed by a 3-month maintenance period. Combination therapy with repaglinide and metformin resulted in a statistically significant improvement in HbA1c and fasting plasma glucose (FPG) compared to repaglinide or metformin monotherapy (Table 8). In this study where metformin dosage was kept constant, the combination therapy of repaglinide and metformin showed dose-sparing effects with respect to repaglinide. The improvement in HbA1c and FPG of the combination group was achieved at a lower daily repaglinide dosage than in the repaglinide monotherapy group.

Table 8: Repaglinide in Combination With Metformin: Mean Changes from Baseline After 4 to 5 Months of Treatment[1]

	Repaglinide in Combination with Metformin	Repaglinide monotherapy	Metformin monotherapy
N	27	28	27
Median Final Dose (mg/day)	6 (repaglinide) 1500 (metformin)	12	1500
HbA_{1c} (%)			
Baseline	8.3	8.6	8.6
Change from baseline	-1.4*	-0.4	-0.3
Fasting plasma glucose (mg/dL)			
Baseline	184	174	194
Change from baseline	-39*	+9	-5

[1] based on intent-to-treat analysis

* $p < 0.05$, for pairwise comparisons with repaglinide and metformin monotherapy.

16 HOW SUPPLIED/STORAGE AND HANDLING

PRANDIMET tablets are supplied as biconvex tablets available in 1 mg/500 mg (yellow) and 2 mg/500 mg (pink) strengths. Tablets are debossed with the Novo Nordisk (Apis) bull symbol on one side and tablet strength on the other side.

1 mg repaglinide/500 mg metformin HCl tablets (yellow)	Bottles of 20	NDC	0169-0093-21
	Bottles of 100	NDC	0169-0093-01
2 mg repaglinide/500 mg metformin HCl tablets (pink)	Bottles of 20	NDC	0169-0092-21
	Bottles of 100	NDC	0169-0092-01

Do not store above 25° C (77° F).

Protect from moisture. Keep bottles tightly closed.

Dispense in tight containers with safety closures.

17 PATIENT COUNSELING INFORMATION

Lactic Acidosis

Explain the risks of lactic acidosis due to the metformin component, its symptoms and conditions that predispose to its development, as noted in *Warnings and Precautions (5.1).* Advise patients to discontinue PRANDIMET immediately and to promptly notify their healthcare provider if unexplained hyperventilation, myalgia, malaise, unusual somnolence, dizziness, slow or irregular heartbeat, sensation of feeling cold (especially in the extremities), or other nonspecific symptoms occur. Gastrointestinal symptoms are common during initiation of metformin treatment and may occur during initiation of PRANDIMET therapy; however, patients should consult their physician if they develop unexplained symptoms. Although gastrointestinal symptoms that occur after stabilization are unlikely to be drug related, such an occurrence of symptoms should be evaluated to determine if it may be due to lactic acidosis or other serious disease.

Counsel patients against excessive alcohol intake while receiving PRANDIMET.

Inform patients about the importance of regular testing of renal function and hematological parameters when receiving treatment with PRANDIMET.

Instruct patients to inform their doctor that they are taking PRANDIMET prior to any surgical or radiological procedure, as temporary discontinuation of PRANDIMET may be required until renal function has been confirmed to be normal *[see Warnings and Precautions (5.1)]*.

Hypoglycemia

Inform patients that PRANDIMET can cause hypoglycemia and instruct patients and their caregivers on self-management procedures including glucose monitoring and management of hypoglycemia. Inform patients that their ability to concentrate and react may be impaired as a result of hypoglycemia. In patients at higher risk for hypoglycemia and patients who have reduced symptomatic awareness of hypoglycemia, increased frequency of blood glucose monitoring is recommended *[see Warnings and Precautions (5.2)]*.

Administration

Instruct patients to take PRANDIMET within 30 minutes before meals. Instruct patients to skip their dose of PRANDIMET when a meal is skipped.

Drug Interactions

Discuss potential drug interactions with patients and inform them of potential drug-drug interactions with PRANDIMET. *[see Drug Interactions (7)]*.

Version: 7

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