TRESIBA®
insulin degludec injection 100 U/mL, 200 U/mL

HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use TRESIBA® safely and effectively. See full prescribing information for TRESIBA®.
TRESIBA® (insulin degludec injection), for subcutaneous use
Initial U.S. Approval: 2015

----- RECENT MAJOR CHANGES ----- 03/2018
Dosage and Administration (2)

----- INDICATIONS AND USAGE -----
TRESIBA® is a long-acting human insulin analog indicated to improve glycemic control in patients 1 year of age and older with diabetes mellitus (1).

Limitations of Use:
• Not recommended for treating diabetic ketoacidosis.
• Not recommended for pediatric patients requiring less than 5 units of TRESIBA®.

----- DOSAGE AND ADMINISTRATION ----- 03/2018

• See Full Prescribing Information for important administration instructions (2.1).
• Rotate injection sites to reduce the risk of lipodystrophy (2.1).
• In adults, inject subcutaneously once daily at any time of day (2.2).
• In pediatric patients inject subcutaneously once daily at the same time every day (2.2).
• Individualize dose based on type of diabetes, metabolic needs, blood glucose monitoring results and glycemic control goal (2.2).
• The recommended days between dose increases are 3 to 4 days (2.2).
• See Full Prescribing Information for recommended starting dose in insulin naïve patients and patients already on insulin therapy (2.3, 2.4).

----- DOSAGE FORMS AND STRENGTHS ----- 03/2018
TRESIBA® injection is available in the following package sizes:
• 100 units/mL (U-100): 3 mL FlexTouch® (3).
• 200 units/mL (U-200): 3 mL FlexTouch® (3).

----- CONTRAINDICATIONS ----- 03/2018

• During episodes of hypoglycemia (4).
• Hypersensitivity to TRESIBA® or one of its excipients (4).

----- WARNINGS AND PRECAUTIONS ----- 03/2018

• Never share a TRESIBA® FlexTouch® pen between patients, even if the needle is changed (5.1).
• Hyper- or hypoglycemia with changes in insulin regimen: Carry out under close medical supervision and increase frequency of blood glucose monitoring (5.2).
• Hypoglycemia: May be life-threatening. Increase monitoring with changes to: insulin dosage, co-administered glucose lowering medications, meal pattern, physical activity; and in patients with renal impairment or hepatic impairment or hypoglycemia unawareness (5.3, 5.4, 6.1).
• Hypoglycemia due to medication errors: Accidental mix-ups between insulin products can occur. Instruct patients to check insulin labels before injection. DO NOT transfer TRESIBA® into a syringe for administration as overdosage and severe hypoglycemia can result (5.4).
• Hypersensitivity reactions: Severe, life-threatening, generalized allergy, including anaphylaxis, can occur. Discontinue TRESIBA® and treat if indicated (5.5).
• Hypokalemia: May be life-threatening. Monitor potassium levels in patients at risk for hypokalemia and treat if indicated (5.6).
• Fluid retention and heart failure with concomitant use of Thiazolidinediones (TZDs): Observe for signs and symptoms of heart failure; consider dosage reduction or discontinuation if heart failure occurs (5.7).

----- ADVERSE REACTIONS ----- 03/2018

Adverse reactions commonly associated with TRESIBA® are:
• hypoglycemia, allergic reactions, injection site reactions, lipodystrophy, pruritus, rash, edema and weight gain (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 ----- 03/2018

• Drugs that may increase the risk of hypoglycemia: antidiabetic agents, ACE inhibitors, angiotensin II receptor blocking agents, disopyramide, fentanyl, fluoxetine, monoamine oxidase inhibitors, pentoxifylline, propoxyphene, salicylates, somatostatin analog (e.g., octreotide), and sulfonamide antibiotics (7).
• Drugs that may decrease the blood glucose lowering effect: atypical antipsychotics, corticosteroids, danazol, diuretics, estrogens, glucagon, isoniazid, niacin, oral contraceptives, phosphonates, phenothiazines, progestogens (e.g., in oral contraceptives), protease inhibitors, somatropin, symathomimetic agents (e.g., albuterol, epinephrine, terbutaline), and thyroid hormones (7).
• Drugs that may increase or decrease the blood glucose lowering effect: Alcohol, beta-blockers, clonidine, lithium salts, and pentamidine (7).
• Drugs that may blunt the signs and symptoms of hypoglycemia: beta-blockers, clonidine, guanethidine, and reserpine (7).

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 08/2018

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE
TRESIBA® is indicated to improve glycemic control in patients with type 1 and type 2 diabetes mellitus.

Limitations of Use
• Not recommended for the treatment of diabetic ketoacidosis.
• Not recommended for pediatric patients requiring less than 5 units of TRESIBA®.

2 DOSAGE AND ADMINISTRATION

2.1 Important Administration Instructions
• Always check insulin labels before administration [see Warnings and Precautions (5.4)].
• Inspect visually for particulate matter and discoloration. Only use TRESIBA® if the solution appears clear and colorless.
• Inject TRESIBA® subcutaneously into the thigh, upper arm, or abdomen.
• Rotate injection sites within the same region from one injection to the next to reduce the risk of lipodystrophy [see Adverse Reactions (6.1)].
• Use TRESIBA® with caution in patients with visual impairment that may rely on audible clicks to dial their dose.
• DO NOT administer TRESIBA® intravenously or in an insulin infusion pump.
• DO NOT dilute or mix TRESIBA® with any other insulin products or solutions.
• DO NOT transfer TRESIBA® from the TRESIBA® pen into a syringe for administration [see Warnings and Precautions (5.4)].

2.2 General Dosing Instructions
• TRESIBA® is available in 2 disposable prefilled pens:
  o TRESIBA® U-100 contains 300 units of TRESIBA® U-100. It delivers doses in 1 unit increments and can deliver up to 80 units in a single injection.
  o TRESIBA® U-200 contains 600 units of TRESIBA® U-200. It delivers doses in 1 unit increments and can deliver up to 160 units in a single injection.
• DO NOT perform dose conversion when using the TRESIBA® U-100 or U-200 pens. The dose window shows the number of insulin units to be delivered and no conversion is needed.
• In adults, inject TRESIBA® subcutaneously once-daily at any time of day.
• In pediatric patients inject TRESIBA® subcutaneously once-daily at the same time every day.
• Individualize and titrate the dose of TRESIBA® based on the patient’s metabolic needs, blood glucose monitoring results, and glycemic control goals.
• The recommended days between dose increases are 3 to 4 days.
• Dose adjustments may be needed with changes in physical activity, changes in meal patterns (i.e., macronutrient content or timing of food intake), changes in renal or hepatic function or during acute illness to minimize the risk of hypoglycemia or hyperglycemia [see Warnings and Precautions (5.3)].
• For adult patients, instruct patients who miss a dose of TRESIBA® to inject their daily dose during waking hours upon discovering the missed dose. Instruct patients to ensure that at least 8 hours have elapsed between consecutive TRESIBA® injections.
• For pediatric patients, instruct patients who miss a dose of TRESIBA® to contact their healthcare provider for guidance and to monitor blood glucose levels more frequently until the next scheduled TRESIBA® dose.

2.3 Starting Dose in Insulin Naïve Patients
Type 1 Diabetes Mellitus:
The recommended starting dose of TRESIBA® in insulin naïve patients with type 1 diabetes is approximately one-third to one-half of the total daily insulin dose. The remainder of the total daily insulin dose should be administered as a short-acting insulin and divided between each daily meal. As a general rule, 0.2 to 0.4 units of insulin per kilogram of body weight can be used to calculate the initial total daily insulin dose in insulin naïve patients with type 1 diabetes.
Type 2 Diabetes Mellitus:
The recommended starting dose of TRESIBA® in insulin naïve patients with type 2 diabetes mellitus is 10 units once daily.

2.4 Starting Dose in Patients Already on Insulin Therapy
Adults with Type 1 or Type 2 Diabetes Mellitus:
Start TRESIBA® at the same unit dose as the total daily long- or intermediate-acting insulin unit dose.
Pediatric Patients 1 Year of Age and Older with Type 1 or Type 2 Diabetes Mellitus:
Start TRESIBA® at 80% of the total daily long- or intermediate-acting insulin unit dose to minimize the risk of hypoglycemia [see Warnings and Precautions (5.2)].

3 DOSAGE FORMS AND STRENGTHS
Injection: TRESIBA® is available as a clear and colorless solution:
• 100 units/mL (U-100): 3 mL FlexTouch® disposable prefilled pen
• 200 units/mL (U-200): 3 mL FlexTouch® disposable prefilled pen

4 CONTRAINDICATIONS
TRESIBA® is contraindicated:
• During episodes of hypoglycemia [see Warnings and Precautions (5.3)].
• In patients with hypersensitivity to TRESIBA® or one of its excipients [see Warnings and Precautions (5.5)].

5 WARNINGS AND PRECAUTIONS

5.1 Never Share a TRESIBA® FlexTouch® Pen Between Patients
TRESIBA® FlexTouch® disposable prefilled pens should never be shared between patients, even if the needle is changed. Sharing poses a risk for transmission of blood-borne pathogens.

5.2 Hyperglycemia or Hypoglycemia with Changes in Insulin Regimen
Changes in insulin, manufacturer, type, or method of administration may affect glycemic control and predispose to hypoglycemia or hyperglycemia. These changes should be made cautiously and only under medical supervision and the frequency of blood glucose monitoring should be increased. For patients with type 2 diabetes, adjustments in concomitant anti-diabetic treatment may be needed. When converting from other insulin therapies to TRESIBA® follow dosing recommendations [see Dosage and Administration (2.4)].

5.3 Hypoglycemia
Hypoglycemia is the most common adverse reaction of insulin, including TRESIBA® [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). TRESIBA®, or any insulin, should not be used during episodes of hypoglycemia [see Contraindications (4)].

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.

Risk Factors for Hypoglycemia
The risk of hypoglycemia generally increases with intensity of glycemic control. The risk of hypoglycemia after an injection is related to the duration of action of the insulin [see Clinical Pharmacology (12.2)] and, in general, is highest when the glucose lowering effect of the insulin is maximal. As with all insulin preparations, the glucose lowering effect time course of TRESIBA® may vary among different individuals or at different times in the same individual depending on the specific physical conditions, including the area of injection as well as the injection site blood supply and temperature.

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

Risk Mitigation Strategies for Hypoglycemia
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.

5.4 Hypoglycemia Due to Medication Errors
Accidental mix-ups between basal insulin products and other insulins, particularly rapid-acting insulins, have been reported. To avoid medication errors between TRESIBA® and other insulins, instruct patients to always check the insulin label before each injection.

To avoid dosing errors and potential overdose, never use a syringe to remove TRESIBA® from the TRESIBA® pen into a syringe [see Dose and Administration (2.1) and Warnings and Precautions (5.3)].

5.5 Hypersensitivity and Allergic Reactions
Severe, life-threatening, generalized allergy, including anaphylaxis, can occur with insulin products, including TRESIBA®. If hypersensitivity reactions occur, discontinue TRESIBA®, treat patients following current standards of care and discontinue or dose reduction of the PPAR-gamma agonist must be considered.

5.6 Hypokalemia
All insulin products, including TRESIBA®, cause a shift in potassium from the extracellular to intracellular space, possibly leading to hypokalemia. Untreated hypokalemia may cause respiratory paralysis, ventricular arrhythmia, and death. Monitor potassium levels in patients at risk for hypokalemia if indicated (e.g., patients using potassium-lowering medications, patients taking medications sensitive to serum potassium concentrations).

5.7 Fluid Retention and Congestive Heart Failure with Concomitant Use of a PPAR Gamma Agonist
Thiazolidinediones (TZDs), which are peroxisome proliferator-activated receptor (PPAR)-gamma agonists can cause dose related fluid retention, particularly when used in combination with insulin. Fluid retention may lead to or exacerbate congestive heart failure. Patients treated with insulin, including TRESIBA® and a PPAR-gamma agonist should be observed for signs and symptoms of congestive heart failure. If congestive heart failure develops, it should be managed according to current standards of care and discontinuation or dose reduction of the PPAR-gamma agonist must be considered.

6 ADVERSE REACTIONS
The following adverse reactions are also discussed elsewhere:
• Hypoglycemia [see Warnings and Precautions (5.3)].
• Medication errors [see Warnings and Precautions (5.4)].
• Hypersensitivity and allergic reactions [see Warnings and Precautions (5.5)].
• Hypokalemia [see Warnings and Precautions (5.6)].

6.1 Clinical Trial Experience
Because clinical trials are conducted under widely 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 practice.

The safety of TRESIBA® in subjects with type 1 diabetes or type 2 diabetes was evaluated in nine trials of 6-12 month duration in adults and in one trial of 12-month duration in pediatric patients. TRESIBA® was used in pediatric patients 1 year of age and older with type 1 diabetes. The cardiovascular safety of TRESIBA® was evaluated in one double-blinded, event-driven trial of 2-year median duration in patients with type 2 diabetes at high risk of cardiovascular events [see Clinical Studies (14)].

The data in Table 1 reflect the exposure of 1102 adult with type 1 diabetes to TRESIBA® with a mean exposure duration to TRESIBA® of 34 weeks in three open-label trials. The mean age was 43
years and 1% were older than 75 years. Fifty-seven percent were male, 81% were White, 2% were Black or African American and 4% were Hispanic. The mean body mass index (BMI) was 26 kg/m². The mean duration of diabetes was 18 years and the mean HbA₁c at baseline was 7.8%. A history of neuropathy, ophthalmopathy, nephropathy and cardiovascular disease at baseline was reported in 11%, 16%, 7% and 0.5% respectively. The mean eGFR at baseline was 97 mL/min/1.73 m² and 7% of the patients had an eGFR less than 60 mL/min/1.73 m².

The data in Table 2 reflect the exposure of 2713 adults with type 2 diabetes to TRESIBA® with a mean exposure duration to TRESIBA® of 36 weeks in six open-label trials. The mean age was 58 years and 3% were older than 75 years. Fifty-eight percent were male, 79% were White, 7% were Black or African American and 15% were Hispanic. The mean BMI was 30 kg/m². The mean duration of diabetes was 11 years and the mean HbA₁c at baseline was 8.3%. A history of neuropathy, ophthalmopathy, nephropathy and cardiovascular disease at baseline was reported for 14%, 10%, 6% and 0.6% of participants respectively. At baseline, the mean eGFR was 83 mL/min/1.73 m² and 9% had an eGFR less than 60 mL/min/1.73 m².

Common adverse reactions (excluding hypoglycemia) occurring in TRESIBA® treated subjects during clinical trials in adult patients with type 1 diabetes mellitus and adults with type 2 diabetes mellitus is listed in Table 1 and Table 2, respectively. Common adverse reactions were defined as reactions occurring in ≥5% of the population studied. Hypoglycemia is not shown in these tables but discussed in a dedicated subsection below.

174 pediatric patients 1 year of age and older with type 1 diabetes were exposed to TRESIBA® with a mean exposure to TRESIBA® of 48 weeks. The mean age was 10 years: 25% were ages 1-5 years, 40% were ages 6-11 years, and 35% were ages 12-17 years. 55.2% were male, 78.2% were White, 2.9% were Black or African American and 4% were Hispanic. The mean body mass index (BMI) was 18.7 kg/m². The mean duration of diabetes was 3.9 years and the mean HbA₁c at baseline was 8.2%.

Common adverse reactions in TRESIBA® treated pediatric patients with type 1 diabetes mellitus was similar to the adverse reactions listed in Table 1.

### Table 1: Adverse Reactions Occurring in ≥5% of TRESIBA®-Treated Adult Patients with Type 1 Diabetes Mellitus

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>TRESIBA® (n=1102)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nasopharyngitis</td>
<td>23.9%</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>11.9%</td>
</tr>
<tr>
<td>Headache</td>
<td>11.8%</td>
</tr>
<tr>
<td>Sinusitis</td>
<td>5.1%</td>
</tr>
<tr>
<td>Gastroenteritis</td>
<td>5.1%</td>
</tr>
</tbody>
</table>

### Table 2: Adverse Reactions Occurring in ≥5% of TRESIBA®-Treated Adult Patients with Type 2 Diabetes Mellitus

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>TRESIBA® (n=2713)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nasopharyngitis</td>
<td>12.9%</td>
</tr>
<tr>
<td>Headache</td>
<td>8.8%</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>8.4%</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>6.3%</td>
</tr>
</tbody>
</table>

### Hypoglycemia

Hypoglycemia is the most commonly observed adverse reaction in patients using insulin, including TRESIBA® [see Warnings and Precautions (5.3)]. The rates of reported hypoglycemia depend on the definition of hypoglycemia used; diabetes type, insulin dose, intensity of glucose control, background therapies, and other intrinsic and extrinsic patient factors. For these reasons, comparing rates of hypoglycemia in clinical trials for TRESIBA® with the incidence of hypoglycemia for other products may be misleading and also, may not be representative of hypoglycemia rates that will occur in clinical practice.

In the open-label adult clinical trials of patients with type 1 and type 2 diabetes, and in the open-label pediatric clinical trial of patients with type 1 diabetes, percentages of adult and pediatric patients with type 1 diabetes randomized to TRESIBA® who experienced at least one episode of hypoglycemia in clinical trials [see Clinical Studies (14)] and adults with type 2 diabetes are shown in Tables 3 and 4, respectively.

Severe hypoglycemia in the open-label trials with adult patients was defined as an episode requiring assistance of another person to actively administer carbohydrate, glucagon, or other resuscitative actions. Severe hypoglycemia in the pediatric trial was defined as an altered mental status where the child could not assist in his own care, was semiconscious or unconscious, or in a coma ± convulsions and may require parenteral therapy (glucagon or intravenous glucose).

A Novo Nordisk hypoglycemia episode was defined as a severe hypoglycemia episode or an episode where a whole blood glucose was less than 50 mg/dL (i.e., with or without the presence of hypoglycemic symptoms).

### Table 3: Percent (%) of Type 1 Diabetes Patients Experiencing at Least One Episode of Severe Hypoglycemia or Novo Nordisk Hypoglycemia on TRESIBA® in Open-Label Adult and Pediatric Clinical Trials

<table>
<thead>
<tr>
<th>Study</th>
<th>TRESIBA® (N=472)</th>
<th>TRESIBA® at the same time each day (N=185)</th>
<th>TRESIBA® at alternating times (N=184)</th>
<th>TRESIBA® (N=174)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Severe hypoglycemia*</td>
<td>12.3%</td>
<td>12.7%</td>
<td>10.4%</td>
<td>17.8%</td>
</tr>
<tr>
<td>Percent of patients</td>
<td>95.6%</td>
<td>99.4%</td>
<td>93.9%</td>
<td>98.3%</td>
</tr>
</tbody>
</table>

Severe hypoglycemia in pediatric patients: an episode with altered mental status, where the child could not assist in his own care, was semiconscious or unconscious, or in a coma ± convulsions and may require parenteral therapy (glucagon or intravenous glucose).

A Novo Nordisk hypoglycemia: a severe hypoglycemia episode or an episode where a laboratory or a self-measured glucose calibrated to plasma was less than 56 mg/dL or where a whole blood glucose was less than 50 mg/dL (i.e., with or without the presence of hypoglycemic symptoms).

### Table 4: Percent (%) of Patients with Type 2 Diabetes Experiencing at Least One Episode of Severe Hypoglycemia or Novo Nordisk Hypoglycemia on TRESIBA® in Open-Label Adult Clinical Trials

<table>
<thead>
<tr>
<th>Study</th>
<th>TRESIBA® (N=766)</th>
<th>TRESIBA® (N=228)</th>
<th>TRESIBA® (N=284)</th>
<th>TRESIBA® (alternating time) (N=230)</th>
<th>TRESIBA® (N=753)</th>
<th>TRESIBA® (N=226)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Severe Hypoglycemia</td>
<td>0.3%</td>
<td>0</td>
<td>0.9%</td>
<td>0.4%</td>
<td>4.5%</td>
<td>0.4%</td>
</tr>
<tr>
<td>Percent of patients</td>
<td>46.5%</td>
<td>28.5%</td>
<td>50%</td>
<td>43.8%</td>
<td>50.9%</td>
<td>80.9%</td>
</tr>
</tbody>
</table>

*OAD: oral antidiabetic agent. **Novo Nordisk hypoglycemia: a severe hypoglycemia episode or an episode where a laboratory or a self-measured glucose calibrated to plasma was less than 56 mg/dL or where a whole blood glucose was less than 50 mg/dL (i.e., with or without the presence of hypoglycemic symptoms).

#### Allergic Reactions

Severe, life-threatening, generalized allergy, including anaphylaxis, generalized skin reactions, angioedema, bronchospasm, hypotension, and shock may occur with any insulin, including TRESIBA®. Severe reactions may manifest with swelling of tongue and lips, diarrhea, nausea, tiredness, and itching and urticaria were reported in 0.9% of patients treated with TRESIBA®.

#### Lipodystrophy

Long-term use of insulin, including TRESIBA®, can cause lipodystrophy at the site of repeated insulin injections. Lipodystrophy includes lipoatrophy (thinning of adipose tissue) and lipoatrophy (thinning of adipose tissue) and may affect insulin absorption [see Dosage and Administration (2.1)]. In the clinical program, lipodystrophy, lipoatrophy, or lipodystrophy was reported in 0.3% of patients treated with TRESIBA®.

#### Injection Site Reactions

Patients taking TRESIBA® may experience injection site reactions, including injection site hematoma, pain, hemorrhage, erythema, nodules, swelling, discoloration, pruritus, and injection site mass. In the clinical program, injection site reactions occurred in 3.8% of patients treated with TRESIBA®.

#### Weight Gain

Weight gain can occur with insulin therapy, including TRESIBA®, and has been attributed to the anabolic effects of insulin. In the clinical program after 52 weeks of treatment, patients with type 1 diabetes treated with TRESIBA® gained an average of 1.8 kg and patients with type 2 diabetes treated with TRESIBA® gained an average of 3.0 kg.

#### Peripheral Edema

Insulin, including TRESIBA®, may cause sodium retention and edema. In the clinical program, peripheral edema occurred in 0.9% of patients with type 1 diabetes mellitus and 3.0% of patients with type 2 diabetes mellitus treated with TRESIBA®.

### 6.2 Immunogenicity

As with all therapeutic proteins, insulin administration may cause anti-insulin antibodies to form. The detection of antibody formation is highly dependent on the sensitivity and specificity of the assay and may be influenced by several factors such as: assay methodology, sample handling, timing of sample collection, concomitant medication, and underlying disease. For these reasons, comparison of the incidence of antibodies to TRESIBA® with the incidence of antibodies in other studies or to other products may be misleading.

In a 52-week study of adult insulin-experienced type 1 diabetes patients, 68.9% of patients who received TRESIBA® were positive at baseline for anti-insulin degludec antibodies and 12.3% of the
patients developed anti-insulin degludec antibodies at least once during the study. In a 52-week study of pediatric insulin-experienced type 1 diabetes patients, 84.1% of patients who received TRESIBA® were positive at baseline for anti-insulin degludec antibodies and 5.8% of patients developed anti-insulin degludec antibodies at least once during the study. In a 52-week study of adult insulin-naïve type 2 diabetes patients, 1.7% of patients who received TRESIBA® were positive at baseline for anti-insulin degludec antibodies and 6.2% of patients developed anti-insulin degludec antibodies at least once during the study. In these trials, between 96.7% and 99.7% of patients who were positive for anti-insulin degludec antibodies were also positive for anti-human insulin antibodies.

7 DRUG INTERACTIONS

Table 5 includes clinically significant drug interactions with TRESIBA®.

Table 5: Clinically Significant Drug Interactions with TRESIBA®

Drugs That May Increase the Risk of Hypoglycemia

<table>
<thead>
<tr>
<th>Drugs:</th>
<th>Interventions:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antidiabetic agents, ACE inhibitors, angiotensin II receptor blocking agents, diisopyramide, fibrates, fluoxetine, monomeric oxidase inhibitors, pentoxifylline, pramlintide, propoxyphene, salicylates, somatostatin analogs (e.g., octreotide), and sulfonamide antibiotics, GLP-1 receptor agonists, DPP-4 inhibitors, SGLT-2 inhibitors.</td>
<td>Dose reductions and increased frequency of glucose monitoring may be required when TRESIBA® is co-administered with these drugs.</td>
</tr>
</tbody>
</table>

Drugs That May Decrease the Blood Glucose Lowering Effect of TRESIBA®

<table>
<thead>
<tr>
<th>Drugs:</th>
<th>Interventions:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atypical antipsychotics (e.g., olanzapine and clozapine), 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.</td>
<td>Dose increases and increased frequency of glucose monitoring may be required when TRESIBA® is co-administered with these drugs.</td>
</tr>
</tbody>
</table>

Drugs That May Increase or Decrease the Blood Glucose Lowering Effect of TRESIBA®

<table>
<thead>
<tr>
<th>Drugs:</th>
<th>Interventions:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alcohol, beta-blockers, clonidine, and lithium salts. Pentamidine may cause hypoglycemia, which may sometimes be followed by hyperglycemia.</td>
<td>Dose adjustment and increased frequency of glucose monitoring may be required when TRESIBA® is co-administered with these drugs.</td>
</tr>
</tbody>
</table>

Drugs That May Blunt Signs and Symptoms of Hypoglycemia

<table>
<thead>
<tr>
<th>Drugs:</th>
<th>Interventions:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Beta-blockers, clonidine, guanethidine, and reserpine</td>
<td>Increased frequency of glucose monitoring may be required when TRESIBA® is co-administered with these drugs.</td>
</tr>
</tbody>
</table>

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no available data with TRESIBA® or insulin degludec in pregnant women to inform a drug-associated risk for major birth defects and miscarriage. There are risks to the mother and fetus associated with poorly controlled diabetes in pregnancy (see Clinical Considerations). Rats and rabbits were exposed to insulin degludec in animal reproduction studies during organogenesis and postnatal development. In rats and rabbits, insulin degludec caused a decrease in fetal weight gain at doses of 1.5 and 2 times the human dose, respectively. The effects observed in the fetus were consistent with those observed in adults. In rabbits, insulin degludec decreased the fetal risk for major birth defects and miscarriage and prevented a treatment-related increase in the incidence of skeletal abnormalities. However, the relevance of these findings to pregnant women is unknown. Therefore, the use of TRESIBA® during pregnancy should be considered only if it is clearly needed. Despite the lack of available information, it is recommended that women with type 2 diabetes who are planning or become pregnant consult their health care provider about the possibility of using TRESIBA®. Ingestion of alcohol and smoking are associated with increased risk of congenital anomalies. Smoking should be discouraged in women who are planning to become pregnant. In the safety outcomes trial (DEVOTE), a total of 1429 (37.4%) of the 3818 TRESIBA®-treated patients with type 2 diabetes had an eGFR less than 60 mL/min/1.73 m² and 1 (0.1%) had an eGFR less than 30 mL/min/1.73 m². A total of 250 (9%) of the 2713 TRESIBA®-treated patients with type 2 diabetes had an eGFR less than 60 mL/min/1.73 m² and no subjects had an eGFR less than 30 mL/min/1.73 m². In the safety outcomes trial (DEVOTE), a total of 1429 (37.4%) of the 3818 TRESIBA®-treated patients with type 2 diabetes had an eGFR less than 60 mL/min/1.73 m² and 108 (2.8%) subjects had an eGFR less than 30 mL/min/1.73 m². Differences in safety or effectiveness were not observed in these subgroup analyses. No clinically relevant difference in the pharmacokinetics of TRESIBA® was identified in a study comparing healthy subjects and subjects with renal impairment including subjects with end stage renal disease (see Clinical Pharmacology [12.3]). However, as with all insulin products, glucose monitoring should be intensified and the TRESIBA® dosage adjusted on an individual basis in patients with renal impairment.

8.7 Hepatic Impairment

No difference in the pharmacokinetics of TRESIBA® was identified in a study comparing healthy subjects and subjects with hepatic impairment (mild, moderate, and severe hepatic impairment) (see Clinical Pharmacology [12.3]). However, as with all insulin products, glucose monitoring should be intensified and the TRESIBA® dosage adjusted on an individual basis in patients with hepatic impairment.

10 OVERDOSAGE

An excess of insulin relative to food intake, energy expenditure, or both may lead to severe and sometimes prolonged and life-threatening hypoglycemia and hypokalemia (see Warnings and Precautions [5.3, 5.6]). Mild episodes of hypoglycemia usually can be treated with oral glucose. Adjustments in drug dosage, meal patterns, or exercise may be needed. More severe episodes of hypoglycemia with coma, seizure, or neurologic impairment may be treated with intramuscular/ intravenous glucagon or concentrated intravenous glucose. After apparent clinical recovery from hypoglycemia, continued observation and additional carbohydrate intake may be necessary to avoid recurrence of hypoglycemia. Hypokalemia must be corrected appropriately.

11 DESCRIPTION

TRESIBA® (insulin degludec injection) is a long-acting basal human insulin analog for subcutaneous injection. Insulin degludec is produced by a process that includes expression of recombinant DNA in Saccharomyces cerevisiae followed by chemical modification. Insulin degludec differs from human insulin in that the amino acid threonine in position B30 has been modified and a side-chain consisting of glutamic acid and a C6 fatty acid has been attached (chemical name: LysB29(Nε-hexadecanoyl)−Glu) des(B30) human insulin). Insulin degludec has a molecular formula of C247H344N60O59S9 and a molecular weight of 6103.97. It has the following structure:

![Figure 1: Structural Formula of TRESIBA®](image-url)

TRESIBA® is a sterile, aqueous, clear, and colorless solution that contains insulin degludec 100 units/mL (U-100) or 200 units/mL (U-200).
Inactive ingredients for the 100 units/mL are glycerol 19.6 mg/mL, phenol 1.50 mg/mL, metacresol 1.72 mg/mL, and water for injection.

Inactive ingredients for the 200 units/mL are glycerol 19.6 mg/mL, phenol 1.50 mg/mL, metacresol 1.72 mg/mL, and water for injection.

TRESIBA® has a pH of approximately 7.6. Hydrochloric acid or sodium hydroxide may be added to adjust pH.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

The primary activity of insulin, including TRESIBA®, is regulation of glucose metabolism. Insulin and its analogs lower blood glucose by stimulating peripheral glucose uptake, especially by skeletal muscle and fat, and by inhibiting hepatic glucose production. Insulin also inhibits lipolysis and proteolysis, and enhances protein synthesis. TRESIBA® forms multi-hexamers when injected into the subcutaneous tissue resulting in a subcutaneous insulin depot. The protracted time action profile of TRESIBA® is predominantly due to delayed absorption of insulin degludec from the subcutaneous tissue to the systemic circulation and to a lesser extent due to binding of insulin-degludec to circulating albumin.

12.2 Pharmacodynamics

The glucose-lowering effect of TRESIBA® after 8 days of once-daily dosing was measured in a euglycemic glucose clamp study enrolling 21 patients with type 1 diabetes. Figure 2 shows the pharmacodynamic effect of TRESIBA® over time following 8 once-daily subcutaneous injections of 0.4 U/kg of TRESIBA® in patients with type 1 diabetes.

Figure 2: Mean GIR Profile for 0.4 units/kg Dose of TRESIBA® (Steady State) in Patients with Type 1 Diabetes Mellitus

The mean maximum glucose lowering effect (GIRmax) of a 0.4 units/kg dose of TRESIBA® was 2.0 mg/kg/min, which was observed at a median of 12 hours post-dose. The glucose lowering effect of TRESIBA® lasted at least 42 hours after the last of 8 once-daily injections.

In patients with type 1 diabetes mellitus, the steady-state, within subjects, day-to-day variability in total glucose lowering effect was 20% with TRESIBA® (within-subject coefficient of variation for AUC0-24 ss).

The total glucose-lowering effect of TRESIBA® over 24 hours measured in a euglycemic clamp study after 8 days of once-daily administration in patients with type 1 diabetes increases approximately in proportion to the dose for doses between 0.4 units/kg to 0.8 units/kg.

The total glucose-lowering effect of 0.4 units/kg of TRESIBA® U-100 and 0.4 units/kg of TRESIBA® U-200, administered at the same dose, and assessed over 24 hours in a euglycemic clamp study after 8 days of once-daily injection was comparable.

12.3 Pharmacokinetics

Absorption

In patients with type 1 diabetes, after 8 hours of once daily subcutaneous dosing with 0.4 units/kg of TRESIBA®, maximum deglucide concentrations of 4472 pmol/L were attained at a median of 9 hours (tmax). After the first dose of TRESIBA®, median onset of appearance was around one hour. Total insulin degludec concentration (i.e., exposure) increased in a dose proportional manner after subcutaneous administration of 0.4 units/kg to 0.8 units/kg TRESIBA®. Total and maximum insulin degludec exposure at steady state are comparable between TRESIBA® U-100 and TRESIBA® U-200 when each is administered at the same units/kg dose.

Insulin degludec concentration reached steady state levels after 3-4 days of TRESIBA® administration (see Dosage and Administration (2.2)).

Distribution

The affinity of insulin degludec to serum albumin corresponds to a plasma protein binding of +99% in human plasma. The results of the in vitro protein binding studies demonstrate that there is no clinically relevant interaction between insulin degludec and other protein bound drugs.

Elimination

The half-life after subcutaneous administration is determined primarily by the rate of absorption from the subcutaneous tissue. On average, the half-life at steady state is approximately 25 hours independent of dose. Degradation of TRESIBA® is similar to that of insulin human, all metabolites formed are inactive. The mean apparent clearance of insulin degludec is 0.03 L/kg (2.1 L/h in 70 kg individual) after single subcutaneous dose of 0.4 units/kg.

Specific Populations

Pediatrics-

Population pharmacokinetic analysis was conducted for TRESIBA® using data from 199 pediatric subjects (1 to <18 years of age) with type 1 diabetes. Body weight was a significant covariate affecting the clearance of TRESIBA®. After adjusting for body weight, the total exposure of TRESIBA® at steady state was independent of age.

Geriatrics-

Pharmacokinetic and pharmacodynamic response of TRESIBA® was compared in 13 younger adult (18–35 years) and 14 geriatric (≥65 years) subjects with type 1 diabetes following two 6-day periods of once-daily subcutaneous dosing with 0.4 units/kg dose of TRESIBA® or insulin glargine. On average, the pharmacokinetic and pharmacodynamic properties of TRESIBA® at steady-state were similar in younger adult and geriatric subjects, albeit with greater between subject variability among the geriatric subjects.

Gender-

The effect of gender on the pharmacokinetics of TRESIBA® was examined in an across-trial analysis of the pharmacokinetic and pharmacodynamic studies conducted using unit/kg doses of TRESIBA®. Overall, there were no clinically relevant differences in the pharmacokinetic properties of insulin degludec between female and male subjects.

Obesity-

The effect of BMI on the pharmacokinetics of TRESIBA® was explored in a cross-trial analysis of pharmacokinetic and pharmacodynamic studies conducted using unit/kg doses of TRESIBA®. For subjects with type 1 diabetes, no relationship between exposure of TRESIBA® and BMI was observed. For subjects with type 1 and type 2 diabetes a trend decrease in glucone-producing effect of TRESIBA® with increasing BMI was observed.

Race and Ethnicity-

TRESIBA® has been studied in a pharmacokinetic and pharmacodynamic study in Black or African American subjects not of Hispanic or Latino origin (n=18). While subjects of Hispanic or Latino origin (n=22) and White subjects not of Hispanic or Latino origin (n=23) with type 2 diabetes mellitus conducted using unit/kg doses of TRESIBA®. There were no statistically significant differences in the pharmacokinetic and pharmacodynamic properties of TRESIBA® between the racial and ethnic groups investigated.

Pregnancy-

The effect of pregnancy on the pharmacokinetics and pharmacodynamics of TRESIBA® has not been studied (see Use in Specific Populations (8.1)).

Renal Impairment

TRESIBA® pharmacokinetics was studied in 32 subjects (n=4-8/group) with normal or impaired renal function/end-stage renal disease following administration of a single subcutaneous dose (0.4 units/kg) of TRESIBA®. Renal function was defined using creatinine clearance (ClCcr) as follows: >90 mL/min (normal), 60-89 mL/min (moderate), 30-59 mL/min (severe) and <30 mL/min (severe). Subjects requiring dialysis were classified as having end-stage renal disease (ESRD). Total (AUCinf hourly) and peak exposure of TRESIBA® were comparable to or higher, respectively in subjects with mild to severe renal impairment except subjects with ESRD who showed similar exposure as compared to subjects with normal renal function. No systematic trend was noted for this increase in exposure across different renal impairment subgroups. Hemodialysis did not affect clearance of TRESIBA® (CL/FUD SS) in subjects with ESRD (see Use in Specific Populations (8.6)).

Hepatic Impairment-

TRESIBA® has been studied in a pharmacokinetic study in 24 subjects (n=6/group) with normal or impaired hepatic function (mild, moderate, and severe hepatic impairment) following administration of a single subcutaneous dose (0.4 units/kg) of TRESIBA®. Hepatic function was defined using Child-Pugh Scores ranging from 5 (mild hepatic impairment) to 15 (severe hepatic impairment). No differences in the pharmacokinetics of TRESIBA® were identified between healthy subjects and subjects with hepatic impairment (see Use in Specific Populations (8.7)).

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Standard 2-year carcinogenicity studies in animals have not been performed to evaluate the carcinogenic potential of insulin degludec. In a 52-week study including human (NPH insulin) as comparator (6.7 units/kg/day), Sprague-Dawley rats were dosed subcutaneously with insulin degludec at 3.3, 6.7, and 10 units/kg/day, resulting in 5 times the human exposure (AUC) when compared to a human subcutaneous dose of 0.75 units/kg/day. Human insulin was dosed at 6.7 units/kg/day. No treatment-related increases in incidences of hyperplasia and neoplastic tumors were recorded in female mammary glands from rats dosed with insulin degludec and no treatment related changes in the female mammary gland cell proliferation were found using BrdU incorporation. Further, no treatment related changes in the occurrence of hyperplastic or neoplastic lesions were seen in other tissues in animals dosed with insulin degludec when compared to vehicle or human insulin.

Genotoxicity testing of insulin degludec was not performed.

In a combined fertility and embryo-fetal study in male and female rats, treatment with insulin degludec up to 21 units/kg/day (approximately 5 times the human subcutaneous dose of 0.75 units/kg/day, based on units/body surface area) prior to mating and in female rats during gestation had no effect on mating performance and fertility.

14 CLINICAL STUDIES

The efficacy of TRESIBA® administered once–daily either at the same time each day or at any time each day in patients with type 1 diabetes and used in combination with a metformal insulin was evaluated in three randomized, open-label, treat-to-target, active-controlled trials in adults and one randomized, open-label, treat-to-target, active-controlled trial in pediatric patients 1 year of age and older. The efficacy of TRESIBA® administered once–daily either at the same time each day or at any time each day in adults with type 2 diabetes and used in combination with a metformal insulin or in combination with common oral anti-diabetic agents was evaluated in six randomized, open-label, treat-to-target active-controlled trials. Adult patients treated with TRESIBA® achieved levels of glycemic control similar to those achieved with LANTUS® (insulin glargine 100 units/mL) and LEVEMIR® (insulin detemir) and achieved statistically significant improvement in HbA1c compared to sitagliptin.

14.1 Type 1 Diabetes – Adult

TRESIBA® Administered at the Same Time Each Day in Combination with a Rapid–Acting Insulin Analogue at Mealtimes in Adult Patients

Study A

The efficacy of TRESIBA® was evaluated in a 52-week randomized, open-label, multicenter trial in 629 patients with type 1 diabetes mellitus (Study A). Patients were randomized to TRESIBA® once-daily with the evening meal or insulin glargine U-100 once-daily according to the approved labeling. Insulin aspart was administered before each meal in both treatment arms.
The mean age of the trial population was 43 years and mean duration of diabetes was 18.9 years. 58.5% were male. 93% were White, 1.9% Black or African American. 5.1% were Hispanic. 8.6% of patients had eGFR ≤ 60 mL/min/1.73m². The mean BMI was approximately 23.9 kg/m².

At week 26, the difference in HbA₁c reduction from baseline between TRESIBA® and insulin glargine U-100 was -0.09% with a 95% confidence interval of [-0.23%; 0.05%] and met the pre-specified non-inferiority margin (0.4%). See Table 6, Study A.

Study B

The efficacy of TRESIBA® was evaluated in a 26-week randomized, open-label, multicenter trial in 455 patients with type 1 diabetes mellitus (Study B). Patients were randomized to TRESIBA® or insulin detemir once-daily in the evening. After 8 weeks, insulin detemir could be dosed twice-daily. 67.1% used insulin detemir once daily at end of trial. 32.9% used insulin detemir twice daily at end of trial. Insulin aspart was administered before each meal in both treatment arms.

The mean age of the trial population was 41.3 years and mean duration of diabetes was 13.9 years. 51.9% were male. 44.6% were White. 0.4% Black or African American. 4.4% were Hispanic. 4.4% of patients had eGFR ≤ 60 mL/min/1.73m². The mean BMI was approximately 23.9 kg/m².

At week 26, the difference in HbA₁c reduction from baseline between TRESIBA® and insulin detemir was -0.09% with a 95% confidence interval of [-0.23%; 0.05%] and met the pre-specified non-inferiority margin (0.4%). See Table 6, Study B.

Table 6: Results at Week 52 in a Trial Comparing TRESIBA® to Insulin Glargine U-100 (Study A) and Week 26 in a Trial Comparing TRESIBA® to Insulin Detemir (Study B) in Adult Patients with Type 1 Diabetes Mellitus Receiving Insulin Aspart at Mealtimes

<table>
<thead>
<tr>
<th>Study A</th>
<th>Study B</th>
</tr>
</thead>
<tbody>
<tr>
<td>TRESIBA® + Insulin aspart</td>
<td>Insulin glargine U-100 + Insulin aspart</td>
</tr>
<tr>
<td>TRESIBA® + Insulin aspart</td>
<td>Insulin aspart</td>
</tr>
<tr>
<td>N</td>
<td>472</td>
</tr>
<tr>
<td>HbA₁c (%)</td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>7.7</td>
</tr>
<tr>
<td>End of trial</td>
<td>7.3</td>
</tr>
<tr>
<td>Adjusted mean change from baseline*</td>
<td>-0.36</td>
</tr>
<tr>
<td>Estimated treatment difference (95%CI)</td>
<td>-0.01 [-0.14; 0.01]</td>
</tr>
<tr>
<td>Proportion Achieving HbA₁c &lt; 7% at Trial End</td>
<td>39.8%</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>165</td>
</tr>
<tr>
<td>End of trial</td>
<td>141</td>
</tr>
<tr>
<td>Adjusted mean change from baseline</td>
<td>-27.6</td>
</tr>
<tr>
<td>Daily basal insulin dose</td>
<td></td>
</tr>
<tr>
<td>Baseline mean</td>
<td>28 U</td>
</tr>
<tr>
<td>Mean dose at end of study</td>
<td>29 U</td>
</tr>
<tr>
<td>Daily bolus insulin dose</td>
<td></td>
</tr>
<tr>
<td>Baseline mean</td>
<td>29 U</td>
</tr>
<tr>
<td>Mean dose at end of study</td>
<td>32 U</td>
</tr>
</tbody>
</table>

*At Week 52

†At Week 26

The change from baseline to end of treatment visit in HbA₁c was analyzed using ANOVA with treatment, region, sex, and anti-diabetic treatment at screening as fixed effects, and age and baseline HbA₁c as covariates. In Study C, there were 15.8% and 15.9% of subjects in the TRESIBA® (same time and alternating times respectively) and 73% insulin-glargin arms for whom data was missing at the time of the HbA₁c measurement.

14.2 Type 1 Diabetes – Pediatric Patients 1 Year of Age and Older

Study J: TRESIBA® Administered at the Same Time Each Day in Combination with a Rapid-Acting Insulin Analog at Mealtimes in Pediatric Patients 1 Year of Age and Older

The efficacy of TRESIBA® was evaluated in a 26-week, randomized, open-label, multicenter trial in 350 patients with type 1 diabetes mellitus (Study J). Patients were randomized to TRESIBA® once-daily or insulin detemir once or twice-daily. Subjects on a twice-daily insulin detemir regimen were dosed at breakfast and in the evening either with the main evening meal or at bedtime. Insulin aspart was administered before each main meal in both treatment arms. At end of trial, 36% used insulin detemir once daily and 64% used insulin detemir twice daily.

The mean age of the trial population was 10 years; 24% were ages 1-5 years; 39% were ages 6-11 years and 36% were ages 12-17 years. The mean duration of diabetes was 4 years. 55.4% were male. 74.6% were White. 2.9% Black or African American. 2.9% were Hispanic. The mean z-score for body weight was 0.31.

At week 26, the difference in HbA₁c reduction from baseline between TRESIBA® and insulin detemir was 0.15% with a 95% confidence interval of [-0.03%; 0.33%] and met the pre-specified non-inferiority margin (0.4%). See Table 8.

Table 7: Results at Week 26 in a Trial Comparing TRESIBA® to Insulin Detemir in Pediatric Patients 1 Year of Age and Older with Type 1 Diabetes Mellitus Receiving Insulin Aspart at Mealtimes

<table>
<thead>
<tr>
<th>TRESIBA® at the same time each day + Insulin aspart</th>
<th>TRESIBA® at alternating times + Insulin aspart</th>
<th>Insulin glargine U-100 + Insulin aspart</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>165</td>
<td>164</td>
</tr>
<tr>
<td>HbA₁c (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>7.7</td>
<td>7.7</td>
</tr>
<tr>
<td>End of trial</td>
<td>7.3</td>
<td>7.3</td>
</tr>
<tr>
<td>Adjusted mean change from baseline*</td>
<td>-0.41</td>
<td>-0.40</td>
</tr>
<tr>
<td>Estimated treatment difference [95%CI]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>TRESIBA® - insulin glargine U-100</td>
<td>0.17 [0.04; 0.30]</td>
<td></td>
</tr>
<tr>
<td>Proportion Achieving HbA₁c &lt; 7% at Trial End</td>
<td>37.0%</td>
<td>37.2%</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>179</td>
<td>173</td>
</tr>
<tr>
<td>End of trial</td>
<td>133</td>
<td>149</td>
</tr>
<tr>
<td>Adjusted mean change from baseline</td>
<td>-41.8</td>
<td>-24.7</td>
</tr>
</tbody>
</table>

Table 8: Results at Week 26 in a Trial Comparing TRESIBA® to Insulin Detemir in Pediatric Patients 1 Year of Age and Older with Type 1 Diabetes Mellitus Receiving Insulin Aspart at Mealtimes

<table>
<thead>
<tr>
<th>TRESIBA® + Insulin aspart</th>
<th>Insulin detemir + Insulin aspart</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>174</td>
</tr>
<tr>
<td>HbA₁c (%)</td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>8.2</td>
</tr>
<tr>
<td>End of 26 weeks</td>
<td>8.0</td>
</tr>
<tr>
<td>Adjusted mean change from baseline after 26 weeks*</td>
<td></td>
</tr>
<tr>
<td>Estimated treatment difference [95%CI]</td>
<td>0.15 [-0.03; 0.33]</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>162</td>
</tr>
<tr>
<td>Mean dose after 26 weeks</td>
<td>150</td>
</tr>
<tr>
<td>Adjusted mean change from baseline after 26 weeks</td>
<td>52.0</td>
</tr>
<tr>
<td>Daily basal insulin dose</td>
<td></td>
</tr>
<tr>
<td>Baseline mean</td>
<td>15 U (0.37 U/kg)</td>
</tr>
<tr>
<td>Mean dose after 26 weeks</td>
<td>16 U (0.37 U/kg)</td>
</tr>
<tr>
<td>Daily bolus insulin dose</td>
<td></td>
</tr>
<tr>
<td>Baseline mean</td>
<td>20 U (0.50 U/kg)</td>
</tr>
<tr>
<td>Mean dose after 26 weeks</td>
<td>23 U (0.56 U/kg)</td>
</tr>
</tbody>
</table>

*The change from baseline to end of treatment visit in HbA₁c was analyzed using ANOVA with treatment, region, sex, and age group as fixed factors, and baseline HbA₁c as covariate.

In Study J, there were 25% of subjects in TRESIBA® and 6.3% insulin detemir arms for whom data was missing at the 26-week HbA₁c measurement.
14.3 Type 2 Diabetes – Adult

Study D: TRESIBA® Administered at the Same Time Each Day as an Add-on to Metformin with or without a DPP-4 Inhibitor in Insulin Naïve Adult Patients

The efficacy of TRESIBA® was evaluated in a 52-week randomized, open-label, multicenter trial that enrolled 1000 insulin naïve patients with type 2 diabetes mellitus inadequately controlled on one or more oral antidiabetic agents (OADs). Patients were randomized to TRESIBA® once-daily with the evening meal or insulin glargine U-100 once-daily according to the approved labeling. Metformin alone (82.5%) or in combination with a DPP-4 inhibitor (17.5%) was used as background therapy in both treatment arms.

The mean age of the trial population was 59.1 years and mean duration of diabetes was 9.2 years. 61.9% were male, 86.4% were White, 7.1% Black or African American, 17.2% were Hispanic. 91.6% of patients had eGFR < 60 mL/min/1.73m². The mean BMI was approximately 31.1 kg/m².

At week 52, the difference in HbA₁c reduction from baseline between TRESIBA® and insulin glargine U-100 was 0.09% with a 95% confidence interval of [-0.04%; 0.22%] and met the pre-specified non-inferiority margin (0.4%). See Table 9.

Table 9: Results at Week 52 in a Trial Comparing TRESIBA® to Insulin Glargine U-100 in Adult Patients with Type 2 Diabetes Mellitus on OAD(s)*

<table>
<thead>
<tr>
<th></th>
<th>TRESIBA® + OAD(s)*</th>
<th>Insulin glargine U-100 + OAD(s)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>773</td>
<td>257</td>
</tr>
<tr>
<td>HbA₁c (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>8.2</td>
<td>8.2</td>
</tr>
<tr>
<td>End of trial</td>
<td>7.1</td>
<td>7.0</td>
</tr>
<tr>
<td>Adjusted mean change from baseline**</td>
<td>-1.06</td>
<td>-1.15</td>
</tr>
<tr>
<td>Estimated treatment difference (95%CI)</td>
<td>TRESIBA® - insulin glargine U-100</td>
<td>-0.09 [-0.04; 0.22]</td>
</tr>
<tr>
<td>Proportion Achieving HbA₁c &lt; 7% at Trial End</td>
<td>51.7%</td>
<td>54.1%</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>174</td>
<td>174</td>
</tr>
<tr>
<td>End of trial</td>
<td>106</td>
<td>115</td>
</tr>
<tr>
<td>Adjusted mean change from baseline</td>
<td>-68.0</td>
<td>-60.2</td>
</tr>
<tr>
<td>Daily insulin dose</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline mean (starting dose)</td>
<td>10 U</td>
<td>10 U</td>
</tr>
<tr>
<td>Mean dose after 52 weeks</td>
<td>56 U</td>
<td>58 U</td>
</tr>
</tbody>
</table>

*OAD: oral antidiabetic agent
**The change from baseline to end of treatment visit in HbA₁c was analyzed using ANOVA with treatment, region, sex, and anti-diabetic treatment at screening as fixed effects, and age and baseline HbA₁c as covariates.

Study E: TRESIBA® U-200 Administered at the Same Time Each Day as an Add-on to Metformin with or without a DPP-4 Inhibitor in Insulin Naïve Adult Patients

The efficacy of TRESIBA® U-200 was evaluated in a 26-week randomized, open-label, multicenter trial in 457 insulin naïve patients with type 2 diabetes mellitus inadequately controlled on one or more oral antidiabetic agents (OADs) at baseline. Patients were randomized to TRESIBA® U-200 once-daily in the evening or insulin glargine U-100 once-daily according to the approved labeling. Both treatment arms were receiving metformin alone (84%) or in combination with a DPP-4 inhibitor and insulin glargine U-100 was 0.04% with a 95% confidence interval of [-0.12%; 0.20%]. This difference from baseline to end of treatment visit in HbA₁c was analyzed using ANOVA with treatment, region, sex, and anti-diabetic treatment at screening as fixed effects, and age and baseline HbA₁c as covariates. In Study E, there were 10% of subjects in the TRESIBA® and 6.8% Insulin glargine arms for whom data was missing at the time of the HbA₁c measurement.

Study F: TRESIBA® Administered at the Same Time Each Day in Insulin Naïve Adult Patients as an Add-on to One or More of the Following Oral Agents: Metformin, Sulfonylurea, Glinides or Alpha-Glucosidase Inhibitors

The efficacy of TRESIBA® was evaluated in a 26-week randomized, open-label, multicenter trial in Asia in 435 insulin naïve patients with type 2 diabetes mellitus inadequately controlled on one or more oral antidiabetic agents (OADs) at baseline. Patients were randomized to TRESIBA® once-daily in the evening or insulin glargine U-100 once-daily according to the approved labeling. Pre-trial oral antidiabetic agents were continued as background therapy except for DPP-4 inhibitors or thiazolidinediones in both treatment arms.

The mean age of the trial population was 58.6 years and mean duration of diabetes was 11.6 years. 53.6% were male. All patients were Asian. 10.9% of patients had eGFR < 60 mL/min/1.73m². The mean BMI was approximately 25.0 kg/m².

At week 26, the difference in HbA₁c reduction from baseline between TRESIBA® and insulin glargine U-100 was 0.11% with a 95% confidence interval of [-0.03%; 0.24%] and met the pre-specified non-inferiority margin (0.4%). See Table 11.

Table 10: Results at Week 26 in a Trial Comparing TRESIBA® U-200 to Insulin Glargine U-100 in Adult Patients with Type 2 Diabetes Mellitus on OAD(s)*

<table>
<thead>
<tr>
<th></th>
<th>TRESIBA® ± DPP-4</th>
<th>Insulin glargine U-100 ± DPP-4</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>228</td>
<td>229</td>
</tr>
<tr>
<td>HbA₁c (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>8.3</td>
<td>8.2</td>
</tr>
<tr>
<td>End of trial</td>
<td>7.0</td>
<td>6.9</td>
</tr>
<tr>
<td>Adjusted mean change from baseline**</td>
<td>-1.18</td>
<td>-1.22</td>
</tr>
<tr>
<td>Estimated treatment difference (95%CI)</td>
<td>TRESIBA® - insulin glargine U-100</td>
<td>0.04 [-0.01; 0.19]</td>
</tr>
<tr>
<td>Proportion Achieving HbA₁c &lt; 7% at Trial End</td>
<td>52.2%</td>
<td>55.9%</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>172</td>
<td>174</td>
</tr>
<tr>
<td>End of trial</td>
<td>106</td>
<td>113</td>
</tr>
<tr>
<td>Adjusted mean change from baseline</td>
<td>-7.1</td>
<td>-6.3</td>
</tr>
<tr>
<td>Daily insulin dose</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline mean</td>
<td>10 U</td>
<td>10 U</td>
</tr>
<tr>
<td>Mean dose after 26 weeks</td>
<td>59 U</td>
<td>62 U</td>
</tr>
</tbody>
</table>

*OAD: oral antidiabetic agent
**The change from baseline to end of treatment visit in HbA₁c was analyzed using ANOVA with treatment, region, sex, and anti-diabetic treatment at screening as fixed effects, and age and baseline HbA₁c as covariates. In Study E, there were 12.3% of subjects in the TRESIBA® and 12.7% Insulin glargine arms for whom data was missing at the time of the HbA₁c measurement.

Table 11: Results at Week 26 in a Trial Comparing TRESIBA® to Insulin Glargine U-100 in Adult Patients with Type 2 Diabetes Mellitus on OAD(s)*

<table>
<thead>
<tr>
<th></th>
<th>TRESIBA® + OAD(s)*</th>
<th>Insulin glargine U-100 + OAD(s)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>268</td>
<td>146</td>
</tr>
<tr>
<td>HbA₁c (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>8.4</td>
<td>8.5</td>
</tr>
<tr>
<td>End of trial</td>
<td>7.2</td>
<td>7.1</td>
</tr>
<tr>
<td>Adjusted mean change from baseline**</td>
<td>-1.42</td>
<td>-1.52</td>
</tr>
<tr>
<td>Estimated treatment difference (95%CI)</td>
<td>TRESIBA® - insulin glargine U-100</td>
<td>0.11 [-0.03; 0.24]</td>
</tr>
<tr>
<td>Proportion Achieving HbA₁c &lt; 7% at Trial End</td>
<td>40.8%</td>
<td>46.8%</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>152</td>
<td>156</td>
</tr>
<tr>
<td>End of trial</td>
<td>100</td>
<td>102</td>
</tr>
<tr>
<td>Adjusted mean change from baseline</td>
<td>-54.6</td>
<td>-53.0</td>
</tr>
<tr>
<td>Daily insulin dose</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline mean</td>
<td>9 U</td>
<td>9 U</td>
</tr>
<tr>
<td>Mean dose after 26 weeks</td>
<td>19 U</td>
<td>24 U</td>
</tr>
</tbody>
</table>

*OAD: oral antidiabetic agent
**The change from baseline to end of treatment visit in HbA₁c was analyzed using ANOVA with treatment, region, sex, and anti-diabetic treatment at screening as fixed effects, and age and baseline HbA₁c as covariates.
The efficacy of TRESIBA® was evaluated in a 26-week randomized, open-label, multicenter trial in 447 patients with type 2 diabetes mellitus inadequately controlled on one or more oral antidiabetic agents (OADs) at baseline. Patients were randomized to TRESIBA® once-daily at any time of day or sitagliptin once-daily according to the approved labeling. One or two of the following oral antidiabetic agents (metformin, sulfonylurea or pioglitazone) were also administered in both treatment arms.

The mean age of the trial population was 55.7 years and mean duration of diabetes was 7.7 years. 58.6% were male. 61.3% were White, 7.6% Black or African American. 21.0% were Hispanic. 6% of patients had eGFR < 60 mL/min/1.73 m². The mean BMI was approximately 30.4 kg/m².

At the end of 26 weeks, TRESIBA® provided greater reduction in mean HbA1c compared to sitagliptin (p < 0.001). See Table 14.

### Table 14: Results at Week 26 in a Trial Comparing TRESIBA® to Sitagliptin in Adult Patients with Type 2 Diabetes Mellitus on OADs*

<table>
<thead>
<tr>
<th>N</th>
<th>TRESIBA® + OAD(s)*</th>
<th>Sitagliptin + OAD(s)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>225</td>
<td></td>
<td></td>
</tr>
<tr>
<td>222</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HbA1c (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>8.8</td>
<td>9.0</td>
</tr>
<tr>
<td>End of trial</td>
<td>7.9</td>
<td>7.9</td>
</tr>
<tr>
<td>Adjusted mean change from baseline**</td>
<td>-1.52</td>
<td>-1.09</td>
</tr>
</tbody>
</table>

### Daily insulin dose

<table>
<thead>
<tr>
<th>Mean dose after 26 weeks</th>
<th>N/A</th>
</tr>
</thead>
</table>

*p < 0.001; 1-sided p-value evaluated at 2.5% level for superiority.

### 14.4 Safety Outcomes Trial

DEVOTE (NCT01959529) Cardiovascular Outcomes Trial of TRESIBA® Administered Once-Daily Between Dinner and Bedtime in Combination with Standard of Care in Subjects with Type 2 Diabetes and Atherosclerotic Cardiovascular Disease

DEVOTE was a multi-center, multi-national, randomized, double-blind, active-controlled, treat-to-target, event-driven trial. 7,637 patients with inadequately controlled type 2 diabetes and atherosclerotic cardiovascular disease were randomized to either TRESIBA® or insulin glargine U-100. Each was administered once-daily between dinner and bedtime in addition to standard care for diabetes and cardiovascular disease for a median duration of 2 years.

Patients eligible to enter the trial were: 50 years of age or older and had established stable, cardiovascular, cerebrovascular, peripheral artery disease, chronic kidney disease or NYHA class II and III heart failure (6% of the enrolled population) or were 60 years of age or older and had other specified risk factors for cardiovascular disease (15% of the enrolled population).

At baseline, demographic and disease characteristics were balanced between treatment groups. The mean age of the trial population was 65 years and mean duration of diabetes was 16.4 years. The population was 62.6% male, 75.6% White 10.9% Black or African American, 10.2% Asian. 14.9% had Hispanic ethnicity. The mean HbA1c was 9.4% and the mean BMI was 33.6 kg/m². The baseline mean estimated glomerular filtration rate (eGFR) was 68 mL/min/1.73 m². 41% of patients had eGFR 60-90 mL/min/1.73 m² and 3% of patients had eGFR < 60 mL/min/1.73 m². Previous history of severe hypoglycemia was not captured in the trial.

At baseline, patients treated their diabetes with oral antidiabetic drugs (72%) and with an insulin regimen (84%). Types of insulins included long acting insulin (60%), intermediate acting insulin (14%) short acting insulin (37%) and premixed insulin (10%). 16% of patients were insulin naive. The most common background oral antidiabetic drugs used at baseline were metformin (60%), sulfonylureas (29%) and DPP-4 inhibitors (12%).

During the trial, investigators could modify anti-diabetic and cardiovascular medications to achieve local standard of care treatment targets for lipids and blood pressure.

### Cardiovascular Outcomes - Patients with T2DM and Atherosclerotic CVD

The incidence of major cardiovascular events with TRESIBA® was evaluated in DEVOTE. Subjects treated with TRESIBA® had a similar incidence of major adverse cardiovascular events (MACE) when compared to those treated with insulin glargine U-100. The primary endpoint in DEVOTE was time from randomization to the first occurrence of a 3-component major adverse cardiovascular event (MACE): cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke. The study was designed to exclude a pre-specified risk margin of 1.3 for the hazard ratio of MACE comparing TRESIBA® to insulin glargine U-100. The primary outcome at end of trial was available for 98.2% of participants in each treatment group.

The time to first occurrence of MACE with TRESIBA® as compared to insulin glargine U-100 was non-inferior (HR: 0.91, 95% CI [0.78;1.06]; see Figure 3). The results of the primary composite MACE endpoint and a summary of its individual components are shown in Table 15.
Table 15: Analysis of the Composite 3-point MACE and Individual Cardiovascular Endpoints in DEVOTE

<table>
<thead>
<tr>
<th>N</th>
<th>TRESIBA®</th>
<th>Insulin glargine U-100</th>
</tr>
</thead>
<tbody>
<tr>
<td>3818</td>
<td>3819</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Number of Patients (%)</th>
<th>Rate per 100 PYO*</th>
<th>Number of Patients (%)</th>
<th>Rate per 100 PYO*</th>
<th>Hazard Ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Composite of first event of CV death, non-fatal MI, or non-fatal stroke (3-Point MACE)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>325 (8.5)</td>
<td>4.41</td>
<td>356 (9.3)</td>
<td>4.86</td>
<td>0.91 [0.78; 1.06]</td>
</tr>
<tr>
<td>CV death</td>
<td>136 (3.6)</td>
<td>1.65</td>
<td>142 (3.7)</td>
<td>1.94</td>
</tr>
<tr>
<td>Non-fatal MI</td>
<td>144 (3.8)</td>
<td>1.95</td>
<td>169 (4.4)</td>
<td>2.31</td>
</tr>
<tr>
<td>Non-fatal stroke</td>
<td>71 (1.9)</td>
<td>0.96</td>
<td>79 (2.1)</td>
<td>1.08</td>
</tr>
</tbody>
</table>

*PYO = patient-years of observation until first MACE, death, or trial discontinuation

Figure 3: Cumulative Event Probability for Time to First MACE in DEVOTE

**Hypoglycemia Outcomes - Patients with T2DM and Atherosclerotic CVD**

The pre-specified secondary endpoints of event and incidence rates of severe hypoglycemia were sequentially tested.

Severe hypoglycemia was defined as an episode requiring assistance of another person to actively administer carbohydrate, glucagon, or other resuscitative actions and during which plasma glucose concentration may not have been available, but where neurological recovery following the return of plasma glucose to normal was considered sufficient evidence that the event was induced by a low plasma glucose concentration.

The incidence of severe hypoglycemia was lower in the TRESIBA® group as compared to the insulin glargine U-100 group (Table 16). Glycemic control between the two groups was similar at baseline and throughout the trial.

Table 16: Severe Hypoglycemic Episodes in Patients Treated with TRESIBA® or Insulin Glargine U-100 in DEVOTE

<table>
<thead>
<tr>
<th>N</th>
<th>TRESIBA®</th>
<th>Insulin glargine U-100</th>
</tr>
</thead>
<tbody>
<tr>
<td>3818</td>
<td>3819</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Severe Hypoglycemia</th>
<th>Percent of patients with events</th>
<th>Estimated odds ratio (95% CI) TRESIBA®/Insulin glargine U-100</th>
<th>Events per 100 Patient Years of Observation</th>
<th>Estimated rate ratio (95% CI) TRESIBA®/Insulin glargine U-100</th>
</tr>
</thead>
<tbody>
<tr>
<td>Percent of patients with events</td>
<td>4.9%</td>
<td>6.6%</td>
<td>0.73 [0.60; 0.89]*</td>
<td>3.70</td>
</tr>
</tbody>
</table>

*Test for superiority evaluated at 5% level for significance, (2-sided p<0.001)

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

TRESIBA® is available as a clear and colorless solution in the following package sizes (see Table 17).

Table 17: Presentations of TRESIBA®

<table>
<thead>
<tr>
<th>TRESIBA®</th>
<th>Total volume</th>
<th>Concentration</th>
<th>Total units available in presentation</th>
<th>NDC number</th>
<th>Max dose per injection</th>
<th>Dose increment</th>
<th>Package Size</th>
</tr>
</thead>
<tbody>
<tr>
<td>U-100 FlexTouch®</td>
<td>3 mL</td>
<td>100 units/mL</td>
<td>300 Units</td>
<td>0169-2660-15</td>
<td>80 Units</td>
<td>1 Unit</td>
<td>5 pens/pack</td>
</tr>
<tr>
<td>U-200 FlexTouch®</td>
<td>3 mL</td>
<td>200 units/mL</td>
<td>600 Units</td>
<td>0169-2550-13</td>
<td>160 Units</td>
<td>2 Unit</td>
<td>3 pens/pack</td>
</tr>
</tbody>
</table>

16.2 Recommended Storage

Unused TRESIBA® should be stored in a refrigerator (36°F to 46°F [2°C to 8°C]). Do not store in the freezer or directly adjacent to the refrigerator cooling element. Do not freeze. Do not use TRESIBA® if it has been frozen.

The storage conditions are summarized in Table 18:

Table 18: Storage Conditions for TRESIBA® FlexTouch®

<table>
<thead>
<tr>
<th>Not in-use (unopened)</th>
<th>In-use (opened)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Refrigerated (36°F to 46°F [2°C to 8°C])</td>
<td>Room Temperature (below 86°F [30°C])</td>
</tr>
<tr>
<td>Room Temperature (below 86°F [30°C])</td>
<td>Refrigerated (36°F to 46°F [2°C to 8°C])</td>
</tr>
</tbody>
</table>

3 mL TRESIBA® U-100 FlexTouch® | Until expiration date | 56 days (8 weeks) | 56 days (8 weeks) | 56 days (8 weeks) |
| 3 mL TRESIBA® U-200 FlexTouch® | Until expiration date | 56 days (8 weeks) | 56 days (8 weeks) | 56 days (8 weeks) |

17 PATIENT COUNSELING INFORMATION

Advise the patient and/or caregiver to read the FDA-approved patient labeling (Patient Information and Instructions for Use).

Never Share a TRESIBA® FlexTouch® Pen Between Patients

Advise patients that they should never share a TRESIBA® FlexTouch® pen device with another person, even if the needle is changed, because doing so carries a risk for transmission of blood-borne pathogens [see Warnings and Precautions (5.1)].

Hypoglycemia or Hypoglycemia

Inform patients that hypoglycemia is the most common adverse reaction with insulin. Inform patients of the symptoms of hypoglycemia. Inform patients that the ability to concentrate and react may be impaired as a result of hypoglycemia. This may present a risk in situations where these abilities are especially important, such as driving or operating other machinery. Advise patients who have frequent hypoglycemia or reduced or absent warning signs of hypoglycemia to use caution when driving or operating machinery.

Advise patients that changes in insulin regimen can predispose to hyper- or hypoglycemia. Advise patients that changes in insulin regimen should be made under close medical supervision [see Warnings and Precautions (5.2)].

Medication Errors

Inform patients to always check the insulin label before each injection [see Warnings and Precautions (5.4)]. Inform patients that the dose counter of TRESIBA® FlexTouch® pen shows the number of units of TRESIBA® to be injected. NO dose re-calculation is required [see Dosage and Administration (2.2)]. Instruct patients to never use a syringe to remove TRESIBA® from the FlexTouch® disposable insulin prefilled pen.

Women of Reproductive Potential

Advise patients to inform their health care professional if they are pregnant or are contemplating pregnancy.
Do not share your TRESIBA® FlexTouch® insulin delivery device with other people, even if the needle has changed. You may give other people a serious infection, or get a serious infection from them.

What is TRESIBA®?
• TRESIBA® is a man-made insulin that is used to control high blood sugar in adults and children who are 1 year of age and older with diabetes mellitus.
• TRESIBA® is not for people with diabetic ketoacidosis (increased ketones in the blood or urine).
• TRESIBA® is not for children who need less than 5 units of TRESIBA® each day.
• It is not known if TRESIBA® is safe and effective in children under 1 year of age.
• TRESIBA® is available in 2 concentrations: The 100 units/mL pen can be injected from 1 to 80 units in a single injection, in increments of 1 unit. The 200 units/mL pen can be injected from 2 to 160 units in a single injection, in increments of 2 units.

Who should not take TRESIBA®?
Do not take TRESIBA® if you:
• are having an episode of low blood sugar (hypoglycemia).
• have an allergy to TRESIBA® or any of the ingredients in TRESIBA®.

Before taking TRESIBA®, tell your healthcare provider about all your medical conditions including, if you are:
• pregnant, planning to become pregnant, or are breastfeeding.
• taking new prescription or over-the-counter medicines, vitamins, or herbal supplements.

Before you start taking TRESIBA®, talk to your healthcare provider about low blood sugar and how to manage it.

How should I take TRESIBA®?
• Read the Instructions for Use that come with your TRESIBA®.
• Take TRESIBA® exactly as your healthcare provider tells you to.
• Do not do any conversion of your dose. The dose counter always shows the selected dose in units. Both the 100 units/mL and 200 units/mL TRESIBA® FlexTouch® pens are made to deliver your insulin dose units in doses.
• Know the type and strength of insulin you take. Do not change the type of insulin you take unless your healthcare provider tells you to. The amount of insulin and the best time for you to take your insulin may need to change if you take different types of insulin.
• Adults: If you miss or are delayed in taking your dose of TRESIBA®:
  o Take your dose as soon as you remember then continue with your regular dosing schedule.
  o Make sure there are at least 8 hours between your doses.
• If children miss a dose of TRESIBA®:
  o Call the healthcare provider for information and instructions about checking blood sugar levels more often until the next scheduled dose of TRESIBA®.
• Check your blood sugar levels. Ask your healthcare provider what your blood sugars should be and when you should check your blood sugar levels.
• Do not reuse or share your needles with other people. You may give other people a serious infection or get a serious infection from them.
• Never inject TRESIBA® into a vein or muscle.
• Never use a syringe to remove TRESIBA® from the FlexTouch® pen.

What should I avoid while taking TRESIBA®?
While taking TRESIBA® do not:
• Drive or operate heavy machinery, until you know how TRESIBA® affects you.
• Drink alcohol or use prescription or over-the-counter medicines that contain alcohol.

What are the possible side effects of TRESIBA®?
TRESIBA® may cause serious side effects that can lead to death, including:
• Low blood sugar (hypoglycemia). Signs and symptoms that may indicate low blood sugar include:
  o dizziness or light-headedness
  o sweating
  o confusion
  o fast heartbeat
• Low potassium in your blood (hypokalemia).
• Heart failure. Taking certain diabetes pills called thiazolidinediones or “TZDs” with TRESIBA® may cause heart failure in some people. This can happen even if you have never had heart failure or heart problems before. If you already have heart failure, it may get worse while you take TZDs with TRESIBA®. Your healthcare provider should monitor you closely while you are taking TZDs with TRESIBA®. Tell your healthcare provider if you have any new or worse symptoms of heart failure including shortness of breath, tiredness, swelling of your ankles or feet and sudden weight gain. Treatment with TZDs and TRESIBA® may need to be adjusted or stopped by your healthcare provider if you have new or worse heart failure.

Your insulin dose may need to change because of:
• change in level of physical activity or exercise
• increased stress
• change in diet
• weight gain or loss
• illness

Common side effects of TRESIBA® may include:
• serious allergic reactions (whole body reactions), reactions at the injection site, skin thickening or pits at the injection site (lipodystrophy), itching, rash, swelling of your hands and feet, and weight gain.

Get emergency medical help if you have:
• trouble breathing, shortness of breath, fast heartbeat, swelling of your face, tongue, or throat, sweating, extreme drowsiness, dizziness, confusion.

These are not all the possible side effects of TRESIBA®. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

General information about the safe and effective use of TRESIBA®.
Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. You can ask your pharmacist or healthcare provider for information about TRESIBA® that is written for health professionals. Do not use TRESIBA® for a condition for which it was not prescribed. Do not give TRESIBA® to other people, even if they have the same symptoms that you have. It may harm them.

What are the ingredients in TRESIBA®?
Active Ingredient: insulin degludec
Inactive Ingredients: zinc, metacresol, glycerol, phenol, and water for injection. Hydrochloric acid or sodium hydroxide may be added.
Manufactured by: Novo Nordisk A/S DK-2880 Bagsvaerd, Denmark
For more information, go to www.novonordisk-us.com or call 1-800-727-6500.

This Patient Information has been approved by the U.S. Food and Drug Administration
Revised: 12/2016

Novo Nordisk®, TRESIBA®, and FlexTouch® are registered trademarks of Novo Nordisk A/S.
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US18TS00318
September 2018
Instructions for Use
TRESIBA® (tre-SI-bah) FlexTouch® Pen 200 units/mL (insulin degludec injection)
• Do not share your TRESIBA® FlexTouch® Pen with other people, even if the needle is changed. You may give other people a serious infection, or get a serious infection from them.
• TRESIBA® FlexTouch® Pen 200 units/mL ("Pen") is a prefilled disposable pen containing 600 units of TRESIBA® (insulin degludec injection) 200 units/mL insulin. You can inject from 2 to 160 units in a single injection. The units can be increased by 2 units at a time.
• This Pen is not recommended for use by the blind or visually impaired without the assistance of a person trained in the proper use of the product.

Supplies you will need to give your TRESIBA® injection:
• TRESIBA® FlexTouch® Pen
• a new NovoFine® or NovoTwist® needle
• alcohol swab
• a sharps container for throwing away used Pens and needles. See “After your injection” at the end of these instructions.

Preparing your TRESIBA® FlexTouch® Pen:
• Wash your hands with soap and water.
• Before you start to prepare your injection, check the TRESIBA® FlexTouch® Pen label to make sure you are taking the right type of insulin. This is especially important if you take more than 1 type of insulin.
• TRESIBA® should look clear and colorless. Do not use TRESIBA® if it is cloudy or colored.
• Do not use TRESIBA® past the expiration date printed on the label or 56 days after you start using the Pen.
• Always use a new needle for each injection to help ensure sterility and prevent blocked needles. Do not reuse or share needles with another person. You may give other people a serious infection, or get a serious infection from them.

NovoFine® Outer needle cap Inner needle cap Needle Paper tab
NovoTwist® Outer needle cap Inner needle cap Needle Paper tab

Step 1:
• Pull Pen cap straight off (See Figure B).

Step 2:
• Check the liquid in the Pen (See Figure C). TRESIBA® should look clear and colorless. Do not use it if it looks cloudy or colored.

Step 3:
• Select a new needle.
• Pull off the paper tab from the outer needle cap (See Figure D).

Step 4:
• Push the capped needle straight onto the Pen and twist the needle on until it is tight (See Figure E).

Step 5:
• Pull off the outer needle cap. Do not throw it away (See Figure F).

Step 6:
• Pull off the inner needle cap and throw it away (See Figure G).

Step 7:
• Turn the dose selector to select 2 units (See Figure H).

Step 8:
• Hold the Pen with the needle pointing up. Tap the top of the Pen gently a few times to let any air bubbles rise to the top (See Figure I).

Step 9:
• Hold the Pen with the needle pointing up. Press and hold in the dose button until the dose counter shows “0”. The “0” must line up with the dose pointer.
• A drop of insulin should be seen at the needle tip (See Figure J).
  o If you do not see a drop of insulin, repeat steps 7 to 9, no more than 6 times.
  o If you still do not see a drop of insulin, change the needle and repeat steps 7 to 9.

Step 10:
• Turn the dose selector to select the number of units you need to inject. The dose pointer should line up with your dose (See Figure K).
  o If you select the wrong dose, you can turn the dose selector forwards or backwards to the correct dose.
  o Each line on the dial is an even number.

Examples
6 units selected
24 units selected

To see how much insulin is left in your TRESIBA® FlexTouch® Pen:
• Turn the dose selector until it stops. The dose counter will line up with the number of units of insulin that is left in your Pen. If the dose counter shows 160, there are at least 160 units left in your Pen.
• If the dose counter shows less than 160, the number shown in the dose counter is the number of units left in your Pen.
Giving your injection:
- Inject your TRESIBA® exactly as your healthcare provider has shown you. Your healthcare provider should tell you if you need to pinch the skin before injecting.
- TRESIBA® can be injected under the skin (subcutaneously) of your upper legs (thighs), upper arms, or stomach area (abdomen).
- Change (rotate) your injection sites within the area you choose for each dose. Do not use the same injection site for each injection.

Step 11:
- Choose your injection site and wipe the skin with an alcohol swab (See Figure M). Let the injection site dry before you inject your dose.

Step 12:
- Insert the needle into your skin (See Figure N).
  - Make sure you can see the dose counter. Do not cover it with your fingers, this can stop your injection.
- Press and hold down the dose button until the dose counter shows “0” (See Figure O).
  - The “0” must line up with the dose pointer. You may then hear or feel a click.
- Keep the needle in your skin after the dose counter has returned to “0” and slowly count to 6 (See Figure P).
  - When the dose counter returns to “0”, you will not get your full dose until 6 seconds later.
  - If the needle is removed before you count to 6, you may see a stream of insulin coming from the needle tip.
  - If you see a stream of insulin coming from the needle tip you will not get your full dose. If this happens you should check your blood sugar levels more often because you may need more insulin.

Step 13:
- Press and hold down the dose button until the dose counter shows “0” (See Figure O).
  - If you see blood after you take the needle out of your skin, press the injection site lightly with a piece of gauze or an alcohol swab. Do not rub the area.
- Count slowly: 1-2-3-4-5-6 (See Figure P).

Step 14:
- Pull the needle out of your skin (See Figure Q).
  - If you see blood after you take the needle out of your skin, press the injection site lightly with a piece of gauze or an alcohol swab. Do not rub the area.

Step 15:
- Carefully remove the needle from the Pen and throw it away (See Figure R).
  - Do not recap the needle. Recapping the needle can lead to needle stick injury.
  - If you do not have a sharps container, carefully slip the needle into the outer needle cap (See Figure S). Safely remove the needle and throw it away as soon as you can.
  - Do not store the Pen with the needle attached. Storing without the needle attached helps prevent leaking, blocking of the needle, and air from entering the Pen.

Step 16:
- Replace the Pen cap by pushing it straight on (See Figure T).

After your injection:
- Put your used TRESIBA® FlexTouch® Pen and needles in a FDA-cleared sharps disposal container right away after use. Do not throw away (dispose of) loose needles and Pens in your household trash.
- If you do not have a FDA-cleared sharps disposal container, you may use a household container that is:
  - made of a heavy-duty plastic
  - can be closed with a tight-fitting, puncture-resistant lid, without sharps being able to come out
  - upright and stable during use
  - leak-resistant
  - properly labeled to warn of hazardous waste inside the container
- When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to dispose of your sharps disposal container. There may be state or local laws about how you should throw away used needles and syringes. Do not reuse or share needles or syringes with another person. For more information about the safe sharps disposal, and for specific information about sharps disposal in the state that you live in, go to the FDA’s website at: http://www.fda.gov/safesharpsdisposal.
- Do not dispose of your used sharps disposal container in your household trash unless your community guidelines permit this. Do not recycle your used sharps disposal container.
Instructions for Use
TRESIBA® (tre-SI-bah) FlexTouch® Pen 100 units/mL (insulin degludec injection)

- Do not share your TRESIBA® FlexTouch® Pen with other people, even if the needle is changed. You may give other people a serious infection, or get a serious infection from them.
- TRESIBA® FlexTouch® Pen 100 units/mL ("Pen") is a prefilled disposable pen containing 300 units of TRESIBA® (insulin degludec injection) 100 units/mL insulin. You can inject from 1 to 80 units in a single injection. The units can be increased by 1 unit at a time.
- This Pen is not recommended for use by the blind or visually impaired without the assistance of a person trained in the proper use of the product.

Supplies you will need to give your TRESIBA® injection:
- TRESIBA® FlexTouch® Pen
- a new NovoFine® or NovoTwist® needle
- alcohol swab
- a sharps container for throwing away used Pens and needles. See “After your injection” at the end of these instructions.

Preparing your TRESIBA® FlexTouch® Pen:
- Wash your hands with soap and water.
- Before you start to prepare your injection, check the TRESIBA® FlexTouch® Pen label to make sure you are taking the right type of insulin. This is especially important if you take more than 1 type of insulin.
- TRESIBA® should look clear and colorless. Do not use TRESIBA® if it is cloudy or colored.
- Do not use TRESIBA® past the expiration date printed on the label or 56 days after you start using the Pen.
- Always use a new needle for each injection to help ensure sterility and prevent blocked needles. Do not reuse or share needles with another person. You may give other people a serious infection, or get a serious infection from them.

NovoFine®

- Outer needle cap
- Inner needle cap
- Needle
- Paper tab

NovoTwist®

- Outer needle cap
- Inner needle cap
- Needle
- Paper tab
- Pen cap

Insulin scale
Insulin window
Dose counter
Dose selector
Dose pointer
Dose button

Step 1:
- Pull Pen cap straight off (See Figure B).

Step 2:
- Check the liquid in the Pen (See Figure C). TRESIBA® should look clear and colorless. Do not use it if it looks cloudy or colored.

Step 3:
- Select a new needle.
  - Pull off the paper tab from the outer needle cap (See Figure D).

Step 4:
- Push the capped needle straight onto the Pen and twist the needle on until it is light (See Figure E).

Step 5:
- Pull off the outer needle cap. Do not throw it away (See Figure F).

Step 6:
- Pull off the inner needle cap and throw it away (See Figure G).

Step 7:
- Turn the dose selector to select 2 units (See Figure H).

Step 8:
- Hold the Pen with the needle pointing up. Tap the top of the Pen gently a few times to let any air bubbles rise to the top (See Figure I).

Step 9:
- Hold the Pen with the needle pointing up. Press and hold in the dose button until the dose counter shows “0”. The “0” must line up with the dose pointer.
- A drop of insulin should be seen at the needle tip (See Figure J).
  - If you do not see a drop of insulin, repeat steps 7 to 9, no more than 6 times.
  - If you still do not see a drop of insulin, change the needle and repeat steps 7 to 9.

Selecting your dose:

Step 10:
- TRESIBA® FlexTouch® Pen 100 units/mL is made to deliver the number of insulin units that your healthcare provider prescribed. Do not perform any dose conversion.
- Check to make sure the dose selector is set at 0.
  - Turn the dose selector to select the number of units you need to inject. The dose pointer should line up with your dose (See Figure K).
  - If you select the wrong dose, you can turn the dose selector forwards or backwards to the correct dose.
  - The even numbers are printed on the dial.
  - The odd numbers are shown as lines.

- The TRESIBA® FlexTouch® Pen insulin scale will show you how much insulin is left in your Pen (See Figure L).

Example: Approx. 200 units left

- To see how much insulin is left in your TRESIBA® FlexTouch® Pen:
  - Turn the dose selector until it stops. The dose counter will line up with the number of units of insulin that is left in your Pen. If the dose counter shows 80, there are at least 80 units left in your Pen.
  - If the dose counter shows less than 80, the number shown in the dose counter is the number of units left in your Pen.
Giving your injection:

- Inject your TRESIBA® exactly as your healthcare provider has shown you. Your healthcare provider should tell you if you need to pinch the skin before injecting.
- TRESIBA® can be injected under the skin (subcutaneously) of your upper legs (thighs), upper arms, or stomach area (abdomen).
- Change (rotate) your injection sites within the area you choose for each dose. Do not use the same injection site for each injection.

**Step 11:**

- Choose your injection site and wipe the skin with an alcohol swab (See Figure M). Let the injection site dry before you inject your dose.

**Step 12:**

- Insert the needle into your skin (See Figure N).
  - Make sure you can see the dose counter. Do not cover it with your fingers; this can stop your injection.
- Press and hold down the dose button until the dose counter shows “0” (See Figure O).
  - The “0” must line up with the dose pointer. You may then hear or feel a click.
- Keep the needle in your skin after the dose counter has returned to “0” and slowly count to 6 (See Figure P).
  - If the needle is removed before you count to 6, you may see a stream of insulin coming from the needle tip.
  - If you see a stream of insulin coming from the needle tip you will not get your full dose. If this happens you should check your blood sugar levels more often because you may need more insulin.

**Step 13:**

- Count slowly:

(Legend)

**Step 14:**

- Pull the needle out of your skin (See Figure Q).
  - If you see blood after you take the needle out of your skin, press the injection site lightly with a piece of gauze or an alcohol swab. Do not rub the area.
  - Do not recap the needle. Recapping the needle can lead to needle stick injury.
- If you do not have a sharps container, carefully slip the needle into the outer needle cap (See Figure S).
  - Safely remove the needle and throw it away as soon as you can.
  - Do not store the Pen with the needle attached. Storing without the needle attached helps prevent leaking, blocking of the needle, and air from entering the Pen.
- If you see a stream of insulin coming from the needle tip, and air from blocking of the needle, and air from entering the Pen.

**Step 15:**

- Carefully remove the needle from the Pen and throw it away (See Figure R).
  - Do not recap the needle. Recapping the needle can lead to needle stick injury.

**Step 16:**

- Replace the Pen cap by pushing it straight on (See Figure T).

**How should I store my TRESIBA® FlexTouch® Pen?**

**Before use:**
- Store unused TRESIBA® FlexTouch® Pens in the refrigerator at 36°F to 46°F (2°C to 8°C).
- Do not freeze TRESIBA®. Do not use TRESIBA® if it has been frozen.
- Unused Pens may be used until the expiration date printed on the label, if kept in the refrigerator.

**Pen in use:**
- Store the Pen you are currently using in the refrigerator between 36°F to 46°F (2°C to 8°C) or keep at room temperature below 86°F (30°C).
- Keep TRESIBA® away from heat or light.
- The TRESIBA® FlexTouch® Pen you are using should be thrown away after 56 days if it is refrigerated or kept at room temperature, even if it still has insulin left in it and the expiration date has not passed.

**General Information about the safe and effective use of TRESIBA®:**
- Keep TRESIBA® FlexTouch® Pens and needles out of the reach of children.
- Always use a new needle for each injection.
- Do not share TRESIBA® FlexTouch® Pens or needles with other people. You may give other people a serious infection, or get a serious infection from them.

- Put your used TRESIBA® FlexTouch® Pen and needles in a FDA-cleared sharps disposal container right away after use. Do not throw away (dispose of) loose needles and Pens in your household trash.
- If you do not have a FDA-cleared sharps disposal container, you may use a household container that is:
  - made of a heavy-duty plastic
  - can be closed with a tight-fitting, puncture-resistant lid, without sharps being able to come out
  - upright and stable during use
  - leak-resistant
  - properly labeled to warn of hazardous waste inside the container
- When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to dispose of your sharps disposal container. There may be state or local laws about how you should throw away used needles and syringes. Do not reuse or share needles or syringes with another person. For more information about the safe sharps disposal, and for specific information about sharps disposal in the state that you live in, go to the FDA’s website at http://www.fda.gov/safesharpsdisposal.
- Do not dispose of your used sharps disposal container in your household trash unless your community guidelines permit this. Do not recycle your used sharps disposal container.

This Instructions for Use has been approved by the U.S. Food and Drug Administration.

**Manufactured by:**
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DK-2880 Bagsvaerd, Denmark
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For more information go to
www.TRESIBA.com

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