What Is BeneFix?
BeneFix, Coagulation Factor IX (Recombinant), is an injectable medicine that is used to help control and prevent bleeding in people with hemophilia B. Hemophilia B is also called congenital factor IX deficiency or Christmas disease.
BeneFix is NOT used to treat hemophilia A.

Selected Safety Information for BeneFix
- BeneFix is contraindicated in patients who have manifested life-threatening, immediate hypersensitivity reactions, including anaphylaxis, to the product or its components, including hamster protein.

Please see full Important Safety Information on back cover and accompanying full Prescribing Information in pocket.
More than 20 years* of experience in previously untreated patients (PUPs) and children

BeneFix Coagulation Factor IX (Recombinant) is FDA approved to control and prevent bleeds in both newly diagnosed and pediatric patients with hemophilia B. It's been shown to be effective across a range of ages—and a range of activity levels.

Selected Safety Information for BeneFix

- Call your health care provider right away if your bleeding is not controlled after using BeneFix.
- Allergic reactions may occur with BeneFix. Call your health care provider or get emergency treatment right away if you have any of the following symptoms: wheezing, difficulty breathing, chest tightness, your lips and gums turning blue, fast heartbeat, facial swelling, faintness, rash or hives.

Please see full Important Safety Information on back cover and accompanying full Prescribing Information in pocket.

Look at the numbers

Experience with PUPs:

- 94.1% of responses were rated excellent or good in PUPs receiving an initial infusion of BeneFix for on-demand treatment
  - 2.9% of responses were rated moderate
  - 1% of responses were rated no response
  - 2% of responses were not rated
  - 1505 infusions were given for on-demand treatment of 997 hemorrhages
  - All bleeding episodes were controlled

Control in PUPs:

- 75% of bleeds were controlled after 1 infusion with BeneFix
  - 14.9% were controlled after 2 infusions
  - 5.4% were controlled after 3 infusions
  - 2.4% were controlled after 4 infusions
  - 2.2% were controlled after >4 infusions

Viral safety is always in mind:

- Every batch has to pass more than 150 tests for quality control

Note the rate of inhibitor development

In PUPs in clinical trials: 2 out of 63 patients developed a high-titer inhibitor. Both patients were removed from the study.

The body can make antibodies called inhibitors. These may stop BeneFix from working properly. The higher the level of inhibitors, the more they may block effective treatment with BeneFix.
**Experience with previously treated patients (PTPs)**

BeneFix has been shown to work in a study with PTPs. It’s the first FDA-approved recombinant treatment that mimics the body’s naturally occurring factor IX. And, in a clinical trial, 81% of bleeds were controlled with 1 infusion of BeneFix.

**What Is BeneFix?**

BeneFix, Coagulation Factor IX (Recombinant), is an injectable medicine that is used to help control and prevent bleeding in people with hemophilia B. Hemophilia B is also called congenital factor IX deficiency or Christmas disease. BeneFix is **NOT** used to treat hemophilia A.

### Look at the numbers

#### Experience with PTPs:

- **90.9% of responses** were rated **excellent** or **good** in PTPs receiving an initial infusion of BeneFix for on-demand treatment
  - 7.1% of responses were rated moderate
  - 0.7% of responses were rated no response
  - 1.3% of responses were not rated
  - **2758 infusions** were given for on-demand treatment of 1796 hemorrhages
  - **All bleeding episodes** were controlled

#### Control in PTPs:

- **80.9% of bleeds** were controlled after 1 infusion with BeneFix
  - 11.6% were controlled after 2 infusions
  - 3.6% were controlled after 3 infusions
  - 1.8% were controlled after 4 infusions
  - 2.1% were controlled after >4 infusions

#### Note the rate of inhibitor development

In PTPs in clinical trials including children and adults: **1 out of 65 patients** developed a low-titer inhibitor.†

†This patient completed the study and had normal factor IX recovery levels (about 15 months after the inhibitor was detected).

### The body can sometimes make antibodies called inhibitors. These may stop BeneFix from working properly.

### Selected Safety Information for BeneFix

- Your body can make antibodies, called “inhibitors,” which may stop BeneFix from working properly.
- If you have risk factors for developing blood clots, such as a venous catheter through which BeneFix is given by continuous infusion, BeneFix may increase the risk of abnormal blood clots. The safety and efficacy of BeneFix administration by continuous infusion have not been established.

Please see full Important Safety Information on back cover and accompanying full Prescribing Information in pocket.
Live actively? Talk to your doctor about whether BeneFix is right for you

With BeneFix, you and your doctor can decide on the right dosing with a range of vial sizes. Not only can you infuse if you have a bleed, but you also can infuse to help prevent a bleed. It’s nice to have a treatment that can fit into your routine.

Flexibility and convenience:
BeneFix® Rapid Reconstitution (R2) Kit

- Travel-ready pack
- Single-use vial
- Sterile infusion set
- Range of vial sizes

BeneFix® can be stored at room temperature or under refrigeration (2°C to 30°C/36°F to 86°F) for up to 2 years until expiration. Do not freeze the BeneFix kit.

Selected Safety Information for BeneFix

- Some common side effects of BeneFix are nausea, injection site reaction, injection site pain, headache, dizziness and rash.
- BeneFix® is contraindicated in patients who have manifested life-threatening, immediate hypersensitivity reactions, including anaphylaxis, to the product or its components, including hamster protein.

Please see full Important Safety Information on back cover and accompanying full Prescribing Information in pocket.
**Information and tools to help you with your treatment**

Pfizer is committed to helping the hemophilia community—and that goes beyond your factor therapy. It’s why we provide additional resources and support for you.

**What Is BeneFix?**

BeneFix, Coagulation Factor IX (Recombinant), is an injectable medicine that is used to help control and prevent bleeding in people with hemophilia B. Hemophilia B is also called congenital factor IX deficiency or Christmas disease. BeneFix is **NOT** used to treat hemophilia A.

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**Selected Safety Information for BeneFix**

- Call your health care provider right away if your bleeding is not controlled after using BeneFix.
- Allergic reactions may occur with BeneFix. Call your health care provider or get emergency treatment right away if you have any of the following symptoms: wheezing, difficulty breathing, chest tightness, your lips and gums turning blue, fast heartbeat, facial swelling, faintness, rash or hives.

Please see full Important Safety Information on back cover and accompanying full Prescribing Information in pocket.
EXPERIENCE MATTERS

With more than 20 years* of experience and dosing flexibility to fit your lifestyle, BeneFix was the most prescribed recombinant factor IX treatment for hemophilia B in 2019.

A history of sustained supply—More than 10 billion IU of product manufactured

Designed with viral safety in mind—More than 150 quality control tests are done on each batch of BeneFix

Convenient storage—BeneFix can be stored at room temperature or under refrigeration [2°C to 30°C/36°F to 86°F] for up to 2 years until expiration. Do not freeze the BeneFix kit

Availability—BeneFix is available in hundreds of institutions nationwide

Learn more today at BeneFix.com

Important Safety Information for BeneFix

• BeneFix is contraindicated in patients who have manifested life-threatening, immediate hypersensitivity reactions, including anaphylaxis, to the product or its components, including hamster protein.

• Call your health care provider right away if your bleeding is not controlled after using BeneFix.

• Allergic reactions may occur with BeneFix. Call your health care provider or get emergency treatment right away if you have any of the following symptoms: wheezing, difficulty breathing, chest tightness, your lips and gums turning blue, fast heartbeat, facial swelling, faintness, rash or hives.

• Your body can make antibodies, called “inhibitors,” which may stop BeneFix from working properly.

• If you have risk factors for developing blood clots, such as a venous catheter through which BeneFix is given by continuous infusion, BeneFix may increase the risk of abnormal blood clots. The safety and efficacy of BeneFix administration by continuous infusion have not been established.

• Some common side effects of BeneFix are nausea, injection site reaction, injection site pain, headache, dizziness and rash.

Please see accompanying full Prescribing Information in pocket.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1.800.FDA.1088.
For intravenous use after reconstitution only

**INDICATIONS AND USAGE**

BeneFIX is an antihemophilic factor (recombinant) indicated for:
- Control and prevention of bleeding episodes in adult and pediatric patients with hemophilia B. (1)
- Peri-operative management in adult and pediatric patients with hemophilia B. (1)

**Limitations of Use:**

BeneFIX is not indicated for the treatment of other factor deficiencies (e.g., factors II, VII, VIII, and X), hemophilia A patients with inhibitors to factor VIII, reversal of coumarin-induced anticoagulation, and bleeding due to low levels of liver-dependent coagulation factors. (1)

**DOSE AND ADMINISTRATION**

For intravenous use after reconstitution only

- One international unit (IU) of BeneFIX per kilogram of body weight increased the circulating activity of factor IX as follows:
  - Adults: 0.8 ± 0.2 IU/dL [range 0.4 to 1.2 IU/dL]. (2.1)
  - Pediatric (<15 years): 0.7 ± 0.3 IU/dL [range 0.2 to 2.1 IU/dL]. (2.1)

- Determine the initial estimated dose using the following formula:

  
  \[
  \text{Dosage} = \text{body weight (kg)} \times \text{desired factor IX increase (IU/dL or \% of normal)} \times \text{reciprocal of observed recovery (IU/kg per IU/dL). (2.1)}
  \]

- Dosage and duration of treatment with BeneFIX depends on the severity of the factor IX deficiency, the location and extent of bleeding, and the patient’s clinical condition, age, and recovery of factor IX. (2)

**DOSAGE FORMS AND STRENGTHS**

BeneFIX is available as lyophilized powder in single use vials containing nominally 250, 500, 1000, 2000, or 3000 IU. (3)

**ADVERSE REACTIONS**

**Clinical Trials Experience**

- Injection site reactions, including anaphylaxis, to the product or its components, including hamster protein. (4)

**Postmarketing Experience**

- Hypersensitivity reactions, including anaphylaxis, are possible. Should symptoms occur, discontinue treatment with the product and seek emergency treatment. Patients may develop hypersensitivity to hamster (CHO) protein as BeneFIX contains trace amounts. (1.1)

- BeneFIX has been associated with the development of thromboembolic complications, including in patients receiving continuous infusion through a central venous catheter. (5.2)

- Nephrotic syndrome has been reported following immune tolerance induction with factor IX products in hemophilia B patients with factor IX inhibitors and a history of allergic reactions to factor IX. (5.3)

- Development of neutralizing antibodies (inhibitors) to BeneFIX may occur. If expected plasma factor IX activity levels are not attained, or if patient presents with allergic reaction, or if bleeding is not controlled with an expected dose, perform an assay that measures factor IX inhibitor concentration. (5.4)

**USE IN SPECIFIC POPULATIONS**

- Pregnancy: No human or animal data. Use only if clearly needed. (8.1)

**ADVERSE REACTIONS**

The most common adverse reactions (incidence >5%) from clinical trials were nausea, injection site reaction, injection site pain, headache, dizziness and rash. (6.1)

**CONTRAINDICATIONS**

Do not use in patients who have manifested life-threatening, immediate hypersensitivity reactions, including anaphylaxis, to the product or its components, including hamster protein. (4)

**WARNINGS AND PRECAUTIONS**

- Dosage and duration of treatment with BeneFIX depend on the severity of the factor IX deficiency, the location and extent of bleeding, and the patient’s clinical condition, age, and recovery of factor IX. (5.1)

**CLINICAL STUDIES**

**REFERENCES**

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

**PREFERENCE**

Revised: 7/2019
Preparation and Reconstitution

The procedures below are provided as general guidelines for the preparation and reconstitution of BeneFIX.

Preparation
1. Always wash hands before performing the following procedures.
2. Use aseptic technique (meaning clean and germ-free) during the reconstitution procedure.
3. Use all components in the reconstitution and administration of this product as soon as possible after opening their sterile containers to minimize unnecessary exposure to the atmosphere.
4. Pooling: If needing more than one vial of BeneFIX per infusion, reconstitute each vial according to the following instructions. Remove the diluent syringe leaving the vial adapter in place, and use a separate large luer lock syringe to draw back the reconstituted contents of each vial. Do not detach the diluent syringes or the large luer lock syringe until you are ready to attach the large luer lock syringe to the next vial adapter.

Reconstitution
1. If refrigerated, allow the vial of lyophilized BeneFIX and the pre-filled diluent syringe to reach room temperature.
2. Remove the plastic flip-top cap from the BeneFIX vial to expose the central portions of the rubber stopper.
3. Slowly depress the plunger rod to inject all the diluent into the BeneFIX vial.

Doses administered should be titrated to the patient's clinical response. Patients may vary in their pharmacokinetic (e.g., half-life, in vivo recovery) and clinical responses to BeneFIX. Although the dose can be estimated by the calculations above, it is highly recommended that, whenever possible, appropriate laboratory tests, including serial factor IX activity assays, be performed.

Dosing Guide for Control and Prevention of Bleeding Episodes and Peri-operative Management

Table 1

<table>
<thead>
<tr>
<th>Type of Hemorrhage</th>
<th>Circulating Factor IX Activity Required (% or IU/dL)</th>
<th>Dosing Interval (hours)</th>
<th>Duration of Therapy (days)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Minor</td>
<td>Uncomplicated hematomas, superficial muscle, or soft tissue</td>
<td>20-30</td>
<td>12-24</td>
</tr>
<tr>
<td>Moderate</td>
<td>Intramuscle or soft tissue with dissection, mucous membranes, dental extractions, or hematuria</td>
<td>25-50</td>
<td>12-24</td>
</tr>
<tr>
<td>Major</td>
<td>Pharynx, retropharynx, retroperitoneum, CNS, surgery</td>
<td>50-100</td>
<td>12-24</td>
</tr>
</tbody>
</table>

Adapted from: Roberts and Eberst1
12. Invert the vial and slowly draw the solution into the syringe.

13. Detach the syringe from the vial adapter by gently pulling and turning the syringe counter-clockwise. Discard the vial with the adapter attached.

14. The reconstituted solution should be clear and colorless. If it is not, discard and use a new kit. If the solution is not to be used immediately, recap the syringe. Do not touch the syringe tip or the inside of the cap.

15. Store the reconstituted solution at room temperature and use it within 3 hours.

Note: BeneFIX, when reconstituted, contains polysorbate-80, which is known to increase the rate of di-(2-ethylhexyl) phthalate (DEHP) extraction from polyvinyl chloride (PVC). This should be considered during the preparation and administration of BeneFIX, including storage time elapsed in a PVC container following reconstitution. It is important that the recommendations for dosage and administration be followed closely [see Dosage and Administration (2, 2.3)].

Note: The tubing of the infusion set included with this kit does not contain DEHP.

2.3 Administration

For intravenous use after reconstitution only.

The safety and efficacy of administration by continuous infusion have not been established [see Warnings and Precautions (5.2)].

- Inspect BeneFIX solution for particulate matter and discoloration prior to administration, whenever solution and container permit.
- Administer BeneFIX using the tubing provided in this kit, and the pre-filled diluent syringe provided, or a single sterile disposable plastic syringe.
- Do not mix or administer BeneFIX in the same tubing or container with other medicinal products.

Administration

1. Attach the syringe to the luer end of the infusion set tubing provided.

2. Apply a tourniquet and prepare the injections site by wiping the skin well with an alcohol swab provided in the kit.

3. Perform venipuncture. Insert the needle on the infusion set tubing into the vein, and remove the tourniquet. Inject the reconstituted BeneFIX intravenously over several minutes. Adjust the rate of administration based on the patient’s comfort level.

4. Following completion of BeneFIX treatment, remove and discard the infusion set. Dispose of all unused solution, empty vial(s), and used needles and syringes in an appropriate container.

Note: Agglutination of red blood cells in the tubing/syringe has been reported with the administration of BeneFIX. No adverse events have been reported in association with this observation. To minimize the possibility of agglutination, it is important to limit the amount of blood entering the tubing. Blood should not enter the syringe. If red blood cell agglutination is observed in the tubing or syringe, discard all material (tubing, syringe and BeneFIX solution) and resume administration with a new package.

5. DOSAGE FORMS AND STRENGTHS

BeneFIX is supplied as a white lyophilized powder in the following dosages:

- 250 IU
- 500 IU
- 1000 IU
- 2000 IU
- 3000 IU

4. CONTRAINDICATIONS

BeneFIX is contraindicated in patients who have manifested life-threatening, immediate hypersensitivity reactions, including anaphylaxis, to the product or its components, including hamster protein.

5. WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity Reactions

Hypersensitivity reactions, including anaphylaxis, have been reported with BeneFIX and have manifested as pruritus, rash, urticaria, hives, facial swelling, dizziness, hypotension, nausea, chest discomfort, cough, dyspnea, wheezing, flushing, discomfort (generalized) and fatigue. Frequently, these events have occurred in close temporal association with the development of factor IX inhibitors. Closely monitor patients for signs and symptoms of acute hypersensitivity reactions, particularly during the early phases of initial exposure to product. Because of the potential for allergic reactions with factor IX concentrates, perform the initial (approximately 10 - 20) administrations of factor IX under medical supervision where proper medical care for allergic reactions could be provided. Advise patients to discontinue use of the product and contact their physician and/or seek immediate emergency care. Immediately discontinue the administration and initiate appropriate treatment if symptoms occur.

BeneFIX contains trace amounts of hamster (CHO) proteins. Patients treated with this product may develop hypersensitivity to these non-human mammalian proteins.

5.2 Thromboembolic Complications

There have been post-marketing reports of thrombotic events in patients receiving continuous-infusion BeneFIX through a central venous catheter, including life-threatening superior vena cava (SVC) syndrome in critically ill neonates [see Adverse Reactions (6.2)]. The safety and efficacy of BeneFIX administration by continuous infusion have not been established [see Dosage and Administration (2, 2.3)].

5.3 Nephrotic Syndrome

Nephrotic syndrome has been reported following immune tolerance induction with factor IX products in hemophilia B patients with factor IX inhibitors and a history of allergic reactions to factor IX. The safety and efficacy of using BeneFIX for immune tolerance induction have not been established.

5.4 Neutralizing Antibodies (Inhibitors)

Neutralizing antibodies (inhibitors) have been reported following administration of BeneFIX [see Adverse Reactions (6.1)]. Evaluate patients using BeneFIX for the development of factor IX inhibitors by appropriate clinical observations and laboratory tests. If expected plasma factor IX activity levels are not attained, or if bleeding is not controlled with an expected dose, perform an assay that measures factor IX inhibitor concentration. Patients with factor IX inhibitors are at an increased risk of severe hypersensitivity reactions or anaphylaxis upon subsequent challenge with factor IX. Evaluate patients experiencing allergic reactions for the presence of an inhibitor and closely monitor patients with inhibitors for signs and symptoms of acute hypersensitivity reactions, particularly during the early phases of initial exposure to product [see Warnings and Precautions (5.1)].

5.5 Monitoring Laboratory Tests

- Monitor patients for factor IX activity levels by the one-stage clotting assay to confirm that adequate factor IX levels have been achieved and maintained, when clinically indicated [see Dosage and Administration (2.1)].
- Monitor patients for the development of inhibitors if expected factor IX activity plasma levels are not attained, or if bleeding is not controlled with the recommended dose of BeneFIX. Determine factor IX inhibitor levels in Bethesda Units (BUs).

6. ADVERSE REACTIONS

The most serious adverse reactions are systemic hypersensitivity reactions, including bronchospastic reactions and/or hypotension and anaphylaxis and the development of high-titer inhibitors necessitating alternative treatments to factor IX replacement therapy.

The most common adverse reactions observed in clinical trials (frequency > 5% of PTPs or PUPs) were headaches, dizziness, nausea, injections site reaction, injection site pain and skin-related hypersensitivity reactions (e.g., rash, hives).

6.1 Clinical Trials 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 clinical practice.

During uncontrolled, open-label clinical studies with BeneFIX conducted in previously treated patients (PTPs), 113 adverse reactions with known or unknown relation to BeneFIX therapy were reported among 38.5% (25 of 65) of subjects (with some subjects reporting more than one event) who received a total of 7,573 infusions. These adverse reactions are summarized in Table 2.
Blood and lymphatic system disorders Factor IX inhibition\(^1\) 2 (3.2%)

Eye disorders Blurred vision 1 (1.6%)

Gastrointestinal disorders Nausea 5 (7.7%)

General disorders and administration site conditions Injection site reaction 1 (1.6%)

Infections and infestations Cellulitis at IV site 1 (1.6%)

Nervous system disorders Headache 2 (3.1%)

Respiratory, thoracic and mediastinal disorders Dry cough 1 (1.6%)

Renal and urinary disorders Renal infarct\(^2\) 1 (1.6%)

Respiratory, thoracic and mediastinal disorders Dyspnea (respiratory distress) 2 (3.2%)

Skin and subcutaneous disorders Hives 1 (0.8%)

Vascular disorders Rash 1 (1.6%)

**Adverse reactions reported within 72 hours of an infusion of BeneFIX.**

\(^1\) Low-titer transient inhibitor formation.

\(^2\) The renal infarct developed in a hepatitis C antibody-positive patient 12 days after a dose of BeneFIX for a bleeding episode. The relationship of the infusion to the prior administration of BeneFIX is uncertain.

### 8.1 Pregnancy

Animal reproduction and lactation studies have not been conducted with BeneFIX. Coagulation Factor IX (Recombinant). It is not known whether BeneFIX can affect reproductive capacity or cause fetal harm when given to pregnant women. BeneFIX should be administered to pregnant women only if needed.

### 8.2 Labor and Delivery

There is no information available on the effect of factor IX replacement therapy on labor and delivery. Use only if needed.

### 8.3 Nursing Mothers

It is not known whether this drug is excreted into human milk. Because many drugs are excreted into human milk, caution should be exercised if BeneFIX is administered to nursing mothers.

### 8.4 Pediatric Use

Safety, efficacy, and pharmacokinetics of BeneFIX have been evaluated in previously treated (PTP) and previously untreated pediatric patients (PUP) [see Clinical Studies (14) and Adverse Reactions (6)]. On average, lower recovery has been observed in pediatric patients younger than 15 years of age [see Clinical Pharmacology (12.3)]. Dose adjustment may be needed [see Dosage and Administration (2.1)].

### 8.5 Geriatric Use

Clinical studies of BeneFIX did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Do not use over 70 years of age.

### 11 DESCRIPTION

BeneFIX, Coagulation Factor IX (Recombinant), is a purified protein produced by recombinant DNA technology. The product is formulated as a sterile, non-pyrogenic, lyophilized powder preparation intended to be reconstituted for intravenous injection. It is available in single-use vials containing the labeled amount of factor IX activity, expressed in International Units (IU). Each vial contains nominally 250, 500, 1000, 2000, or 3000 IU of recombinant coagulation factor IX. The potency (in IU) is determined using an in vitro one-stage clotting assay against the World Health Organization (WHO) International Standard for Factor IX concentrate. One IU is the amount of factor IX activity present in 1 mL of pooled, normal human plasma. After reconstitution of the lyophilized drug product, the concentrations of excipients are 0.234% sodium chloride, 8 mM L-histidine, 0.8% sucrose, 208 mM glycine, 0.004% polysorbate 80. The specific activity of BeneFIX is greater than or equal to 200 IU per milligram of protein. BeneFIX contains no preservatives and all dosage strengths yield a clear, colorless solution upon reconstitution.

Coagulation factor IX is the active ingredient in BeneFIX. It has a primary amino acid sequence that is identical to the Ala\(^{148}\) allelic form of human factor IX, and has structural and functional characteristics similar to those of endogenous factor IX.

BeneFIX is not derived from human blood. It is produced by a genetically engineered Chinese hamster ovary (CHO) cell line that is extensively characterized. No additives of animal or human origin are used during the cell culture, purification, and formulation processes of BeneFIX. The stored cell banks are free of human blood or plasma products. The CHO cell line secretes recombinant factor IX into a defined cell culture medium, and the recombinant factor IX is purified by a four-step chromatography purification process that does not require a monoclonal antibody step. The process also includes a membrane filtration step that has the ability to retain molecules with apparent molecular weights >70,000 Da (such as large proteins and viral particles). BeneFIX is a single component by SDS-polyacrylamide gel electrophoresis evaluation.

### 12 CLINICAL PHARMACOLOGY

#### 12.1 Mechanism of Action

BeneFIX temporarily replaces the missing clotting factor IX that is needed for effective hemostasis.

#### 12.2 Pharmacodynamics

The activated partial thromboplastin time (aPTT) is prolonged in people with hemophilia B. Treatment with factor IX concentrate may normalize the aPTT by temporarily replacing the factor IX. The administration of BeneFIX increases plasma levels of factor IX, and can temporarily correct the coagulation defect in these patients.
12.3 Pharmacokinetics

After single intravenous doses of 50 IU/kg of previously marketed BeneFIX (reconstituted with Sterile Water for Injection), in 37 previously treated adult patients (>15 years), each given as a 10-minute infusion, the mean increase from pre-infusion level in circulating factor IX activity was 0.8 ± 0.2 IU/dL per IU/kg infused (range 0.4 to 1.4 IU/dL per IU/kg) and the mean biologic half-life was 18.8 ± 5.4 hours (range 11 to 36 hours).

In the original randomized, cross-over pharmacokinetic study in previously treated patients (PTPs), the in vivo recovery using previously marketed BeneFIX was statistically significantly less (28% lower, p<0.05) than the recovery using a plasma-derived factor IX product (pdFIX). A summary of pharmacokinetic data for BeneFIX and pdFIX are presented in Table 4.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>BeneFIX, n = 11</th>
<th>pdFIX, n = 11</th>
</tr>
</thead>
<tbody>
<tr>
<td>AUC∞(IU/hr/dL)</td>
<td>548 ± 92</td>
<td>928 ± 191</td>
</tr>
<tr>
<td>t1/2 (hr)</td>
<td>18.1 ± 5.1</td>
<td>17.7 ± 5.3</td>
</tr>
<tr>
<td>CL (mL/hr/kg)</td>
<td>8.62 ± 1.7</td>
<td>6.00 ± 1.4</td>
</tr>
<tr>
<td>K-value (IU/dL per IU/kg)</td>
<td>0.84 ± 0.30</td>
<td>1.17 ± 0.26</td>
</tr>
<tr>
<td>In vivo Recovery (%)</td>
<td>37.8 ± 14.0</td>
<td>52.6 ± 12.4</td>
</tr>
</tbody>
</table>

Abbreviations: AUC∞ = area under the plasma concentration-time curve from zero to infinity; K-value = incremental recovery; t1/2 = plasma elimination half-life; CL = clearance; SD = standard deviation.

There was no significant difference in biological half-life. Structural differences of the BeneFIX molecule compared with pdFIX were shown to contribute to the lower recovery. In subsequent evaluations for up to 24 months, the pharmacokinetic parameters were similar to the initial results.

In a subsequent randomized, cross-over pharmacokinetic study, BeneFIX reconstituted in 0.234% sodium chloride diluent was shown to be pharmacokinetically equivalent to the previously marketed BeneFIX (reconstituted with Sterile Water for Injection) in 24 previously treated patients (≥12 years) at a dose of 75 IU/kg. In addition, pharmacokinetic parameters were followed up in 23 previously treated patients after repeated administration of BeneFIX for six months and found to be unchanged compared with those obtained at the initial evaluation. A summary of pharmacokinetic data are presented in Table 5.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Parameters at Initial Visit</th>
<th>Parameters at Month 6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cmax (IU/dL)</td>
<td>54.5 ± 15.0</td>
<td>57.3 ± 13.2</td>
</tr>
<tr>
<td>AUC∞(IU/hr/dL)</td>
<td>940 ± 237</td>
<td>923 ± 205</td>
</tr>
<tr>
<td>t1/2 (hr)</td>
<td>22.4 ± 5.3</td>
<td>23.8 ± 6.5</td>
</tr>
<tr>
<td>CL (mL/hr/kg)</td>
<td>8.47 ± 2.12</td>
<td>8.54 ± 2.04</td>
</tr>
<tr>
<td>K-value (IU/dL per IU/kg)</td>
<td>0.73 ± 0.20</td>
<td>0.76 ± 0.18</td>
</tr>
<tr>
<td>In vivo Recovery (%)</td>
<td>34.5 ± 9.3</td>
<td>36.8 ± 8.7</td>
</tr>
</tbody>
</table>

Abbreviations: Cmax = peak concentration; K-value = incremental recovery; t1/2 = plasma elimination half-life; CL = clearance; SD = standard deviation.

Pediatric Patients (≤15 years)

Nineteen (19) previously treated pediatric patients (range 4 to ≤15 years) underwent pharmacokinetic evaluations for up to 24 months. Fifty-eight previously untreated patients (PUPs) less than 15 years of age at baseline underwent at least one recovery assessment within 30 minutes post-infusion in the presence or absence of hemorrhage during the study. A total of 202 recovery assessments collected during the 60-month period from these 58 PUPs are combined with 19 recovery assessments from PTPs and were summarized by age group in Table 6. There was one recovery assessment in a neonate, which had a value of 0.46 IU/dL per IU/kg. The overall mean recovery and FIX elimination half-lives were 0.7 ± 0.3 IU/dL per IU/kg and 20.2 ± 4.0 hours, respectively.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>K-value</th>
<th>t1/2(b)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Infants (≥1 month to &lt;2 years)</td>
<td>33</td>
<td>0.7 ± 0.4 (0.2, 2.1)</td>
</tr>
<tr>
<td>Children (≥2 years to &lt;12 years)</td>
<td>61</td>
<td>0.7 ± 0.2 (0.2, 1.5)</td>
</tr>
<tr>
<td>Adolescents (≥12 years to ≤15 years)</td>
<td>9</td>
<td>0.8 ± 0.3 (0.4, 1.4)</td>
</tr>
</tbody>
</table>

^a n = 13
^b n = 6

Data presented are mean ± standard deviation (min, max). Abbreviations: ND = not determined; K-value = incremental recovery; t1/2 = terminal phase elimination half-life.

Note: The columns are not mutually exclusive; individual patients may be listed under more than 1 age category.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

BeneFIX has been shown to be nonmutagenic in the Ames assay and non-clastogenic in a chromosomal aberrations assay. No investigations on carcinogenesis or impairment of fertility have been conducted.

14 CLINICAL STUDIES

Efficacy of BeneFIX has been evaluated in clinical studies in which a total of 128 subjects received BeneFIX either for the treatment of bleeding episodes on an on-demand basis, for the prevention of bleeds (prophylaxis) or for management of hemostasis in the surgical setting (surgical prophylaxis).

Fifty-six PTPs and sixty-three PUPs were treated for bleeding episodes on an on-demand basis or for the prevention of bleeds (see Tables 7 and 8). The PTPs were followed over a median interval of 24 months (mean 23.4 ± 6.3 months) and for a median of 83.5. The PUPs were followed over a median interval of 37 months (mean 38.1 ± 16.4 months) and for a median of 89 exposure days.

Fifty-five PTPs and forty-four PUPs received BeneFIX for the treatment of bleeding episodes (see Table 7). Bleeding episodes that were managed successfully included hemarthrosis and bleeding in soft tissue and muscle. Data concerning the severity of bleeding episodes were not reported. In the PTPs, 88% of total infusions administered for on-demand treatment were rated as an “excellent” or “good” response.

<table>
<thead>
<tr>
<th>Median dose: IU/kg (range)</th>
<th>Rate of bleeds resolved with 1 infusion</th>
<th>Response to 1st infusion Rating</th>
</tr>
</thead>
<tbody>
<tr>
<td>PTPs</td>
<td>N=55^a</td>
<td>42.8 (6.5 - 224.6)</td>
</tr>
<tr>
<td>PUPs</td>
<td>N=54^a</td>
<td>62.7 (8.2 - 292)</td>
</tr>
</tbody>
</table>

^a One subject discontinued the study after one month of treatment due to bleeding episodes that were difficult to control; he did not have a detectable inhibitor.

^b Three subjects were not successfully treated during one episode in a subject due to delayed time to infusion and insufficient dosing and in 2 subjects due to inhibitor formation.

Response ratings not provided for 1.3% and 2% of 1st infusions for PTPs and PUPs, respectively.

A total of 20 PTPs were treated with BeneFIX for secondary prophylaxis (the regular administration of FIX replacement therapy to prevent bleeding in patients who may have already demonstrated clinical evidence of hemophilic arthropathy or joint disease) at some regular interval during the study with a mean of 2 infusions per week (see Table 8). Thirty-two PUPs were administered BeneFIX for routine (primary and secondary) prophylaxis (see Table 8). Twenty-four PUPs were administered BeneFIX at least twice weekly, and eight PUPs were administered BeneFIX once weekly. Seven PTPs experienced a total of 26 spontaneous bleeding episodes within 48 hours after an infusion. Six spontaneous bleeds within 48 hours after an infusion were reported in 5 PUPs. Prophylaxis therapy was rated as “excellent” or “effective” in 93% of PTPs receiving prophylaxis one to two times per week.

Data from 57 PUP subjects who underwent repeat recovery testing for up to 60 months demonstrated that the average incremental FIX recovery was consistent over time, as shown in Figure 1.

Figure 1. Average Incremental rFIX Recovery over Time
Management of hemostasis was evaluated in the surgical setting in both PTPs and PUPs (see Table 9). Thirty-six surgical procedures have been performed in 28 PTPs with 23 major surgical procedures performed (including 6 complicated dental extractions). Thirty surgical procedures have been performed in 23 PUPs. Twenty-eight of these procedures were considered minor. Hemostasis was maintained throughout the surgical period; however, one PTP subject required evacuation of a surgical wound-site hematoma, and another PTP subject who received BeneFIX after a tooth extraction required further surgical intervention due to oozing at the extraction site. There was no clinical evidence of thrombotic complications in any of the subjects.

Among the PTP surgery subjects, the median increase in circulating factor IX activity was 0.7 IU/dL per IU/kg infused (range 0.3 – 1.2 IU/dL; mean 0.8 ± 0.2 IU/dL per IU/kg). The median elimination half-life for the PTP surgery subjects was 19.4 hours (range 10 – 37 hours; mean 21.3 ± 8.1 hours).

Table 9: Efficacy of BeneFIX for Surgical Procedures in PTPs and PUPs

<table>
<thead>
<tr>
<th>Surgery Type</th>
<th>Number of Procedures</th>
<th>Excellent/ Good</th>
<th>Moderate</th>
<th>No Response</th>
</tr>
</thead>
<tbody>
<tr>
<td>Previously Treated Patients</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ankle surgery</td>
<td>2 (2)</td>
<td>2 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Hip prosthesis implant (right)</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Knee arthroplasty (2 bilateral, 1 right)</td>
<td>3 (3)</td>
<td>3 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Knee arthroscopic synovectomy</td>
<td>2 (2)a</td>
<td>1 (50%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Liver transplantation (orthotopic)</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Splenectomy</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>External fixation device removal (wrist)</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Hernia repair</td>
<td>3 (2)</td>
<td>3 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Subacromial decompression (left)</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Calf debridement, dental extractionb</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lymph node removal, dental extractionb</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Left heel cord lengthening</td>
<td>1 (1)</td>
<td>1 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Dental proceduresc</td>
<td>12 (11)</td>
<td>11 (92%)</td>
<td>1 (8%)</td>
<td>-</td>
</tr>
<tr>
<td>Minor procedures</td>
<td>6 (6)</td>
<td>6 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Previously Untreated Patients</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hernia repair</td>
<td>2 (2)</td>
<td>2 (100%)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Minor procedures</td>
<td>28 (21)d</td>
<td>27 (96%)</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

- Response assessment not provided for 1 procedure.
- Includes pulse and continuous-infusion regimens; GI counted as 1 procedure in this summary.
- Includes complicated extractions (6), clearance, and fillings.

Nine of the major surgical procedures were performed in 8 PUPs using a continuous-infusion regime. Five of the surgical procedures were performed in PUPs using a continuous-infusion regimen over 3 to 5 days. Although circulating factor IX levels targeted to restore and maintain hemostasis were achieved with both pulse replacement and continuous infusion regimens, clinical trial experience with continuous infusion of BeneFIX for surgical prophylaxis in hemophilia B has been too limited to establish the safety and clinical efficacy of administration of the product by continuous infusion.

15 REFERENCES

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied
BeneFIX, Coagulation Factor IX (Recombinant), is supplied in kits that include single-use vials which contain nominally 250, 500, 1000, 2000, or 3000 IU per vial (NDC 58394-633-03, 58394-634-03, 58394-635-03, 58394-636-03 and 58394-637-03, respectively) with sterile pre-filled diluent syringe, vial adapter reconstitution device, sterile infusion set, and two (2) alcohol swabs, one bandage, and one gauze pad. Actual factor IX activity in IU is stated on the label of each vial.

Storage and Handling
Product kit as packaged for sale
• Store BeneFIX at room temperature or under refrigeration, at a temperature of 2 to 30°C (36 to 86°F).
• Do not freeze to prevent damage to the diluent syringe.
• Do not use BeneFIX after the expiration date on the label.

Product after reconstitution
The product does not contain a preservative and should be used within 3 hours.

17 PATIENT COUNSELING INFORMATION
• Advise patients to read the FDA-approved patient labeling (Patient Information and Instructions for Use)
• Allergic-type hypersensitivity reactions are possible. Inform patients of the early signs of hypersensitivity reactions [including hives (rash with itching), generalized urticaria, tightness of the chest, wheezing, hypotension] and anaphylaxis. Advise patients to discontinue use of the product and contact their physicians if these symptoms occur.
• Advise patients to contact their physician or treatment facility for further treatment and/or assessment if they experience a lack of a clinical response to factor IX replacement therapy, as in some cases this may be a manifestation of an inhibitor.
BeneFIX is an injectable medicine that is used to help control and prevent bleeding in people with hemophilia B. Hemophilia B is also called congenital factor IX deficiency or Christmas disease. BeneFIX is NOT used to treat hemophilia A.

What should I tell my doctor before using BeneFIX?
Tell your doctor about all of the medicines you take, including prescription and non-prescription medicines, such as over-the-counter medicines, supplements, or herbal medicines. Tell your doctor about all of your medical conditions, including if you:
- are pregnant or planning to become pregnant. It is not known if BeneFIX may harm your unborn baby.
- are breastfeeding. It is not known if BeneFIX passes into the milk and if it can harm your baby.

How should I store BeneFIX?
The initial administrations of BeneFIX should be administered under proper medical supervision, where proper medical care for severe allergic reactions could be provided.

See the step-by-step instructions for infusing BeneFIX at the end of this leaflet. You should always follow the specific instructions given by your doctor. The steps listed below are general guidelines for using BeneFIX. If you are unsure of the procedures, please call your doctor or pharmacist before using.

Call your doctor right away if bleeding is not controlled after using BeneFIX.

Your doctor will prescribe the dose that you should take. Your doctor may need to test your blood from time to time. BeneFIX should not be administered by continuous infusion.

What if I take too much BeneFIX?
Call your doctor if you take too much BeneFIX.

What are the possible side effects of BeneFIX?
Allergic reactions may occur with BeneFIX. Call your doctor or get emergency treatment right away if you have any of the following symptoms:
- wheezing
- swelling of the face
- difficulty breathing
- faintness
- chest tightness
- rash
- turning blue (look at lips and gums)
- hives
- fast heartbeat

Your body can also make antibodies, called “inhibitors,” against BeneFIX, which may stop BeneFIX from working properly.

Some common side effects of BeneFIX are nausea, injection site reaction, injection site pain, headache, dizziness and rash. BeneFIX may increase the risk of thromboembolism (abnormal blood clots) in your body if you have risk factors for developing blood clots, including an indwelling venous catheter through which BeneFIX is given by continuous infusion. There have been reports of severe blood clotting events, including life-threatening blood clots in critically ill neonates, while receiving continuous-infusion BeneFIX through a central venous catheter. The safety and efficacy of BeneFIX administration by continuous infusion have not been established.

These are not all the possible side effects of BeneFIX.

Tell your doctor about any side effect that bothers you or that does not go away.

How should I store BeneFIX?
DO NOT FREEZE BeneFIX kit.

BeneFIX kit can be stored at room temperature (below 86°F) or under refrigeration.

Throw away any unused BeneFIX and diluent after the expiration date indicated on the label.

Freezing should be avoided to prevent damage to the pre-filled diluent syringe.

BeneFIX does not contain a preservative. After reconstituting BeneFIX, you can store it at room temperature for up to 3 hours. If you have not used it in 3 hours, throw it away.

Do not use BeneFIX if the reconstituted solution is not clear and colorless.

What else should I know about BeneFIX?
Medicines are sometimes prescribed for purposes other than those listed here. Do not use BeneFIX for a condition for which it was not prescribed. Do not share BeneFIX with other people, even if they have the same symptoms that you have.

This Patient Leaflet summarizes the most important information about BeneFIX. If you would like more information, talk with your doctor. You can ask your doctor or pharmacist for information about BeneFIX that was written for healthcare professionals.

Instructions for Use
BeneFIX® / BEN-uh-fiks/
[Coagulation Factor IX (Recombinant)]

BeneFIX is supplied as a powder. Before it can be infused in your vein (intravenous injection), you must reconstitute the powder by mixing it with the liquid diluent supplied. The liquid diluent is 0.234% sodium chloride. BeneFIX should be reconstituted and infused using the infusion set, diluent, syringe, and adapter provided in this kit, and by following the directions below.

RECONSTITUTION
Always wash your hands before performing the following steps. Try to keep everything clean and germ-free while you are reconstituting BeneFIX. Once you open the vials, you should finish reconstituting BeneFIX as soon as possible. This will help keep the infusion set materials germ-free.

Note: If you use more than one vial of BeneFIX per infusion, reconstitute each vial according to steps 1 through 13.

1. If refrigerated, let the vial of BeneFIX and the pre-filled diluent syringe reach room temperature.
2. Remove the plastic flip-top cap from the BeneFIX vial to show the center part of the rubber stopper.
3. Wipe the top of the vial with the alcohol swab provided, or use another antiseptic solution, and allow to dry. After cleaning, do not touch the rubber stopper with your hand or allow it to touch any surface.
4. Peel back the cover from the clear plastic vial adapter package. Do not remove the adapter from the package.
5. Place the vial on a flat surface. While holding the adapter in the package, place the vial adapter over the vial. Press down firmly on the package until the adapter snaps into place on top of the vial, with the adapter spike penetrating the vial stopper.

These are not all the possible side effects of BeneFIX.

Tell your doctor about any side effect that bothers you or that does not go away.

How should I store BeneFIX?
DO NOT FREEZE BeneFIX kit.

BeneFIX kit can be stored at room temperature (below 86°F) or under refrigeration.
6. Grasp the plunger rod as shown in the picture below. Do not touch the shaft of the plunger rod. Attach the threaded end of the plunger rod to the diluent syringe plunger by pushing and turning firmly.

7. Break off the tamper-resistant, plastic-tip cap from the diluent syringe by snapping the perforation of the cap. Do not touch the inside of the cap or the syringe tip. The diluent syringe may need to be recapped (if reconstituted BeneFIX is not used immediately), so place the cap on its tip on a clean surface in a spot where it will stay clean.

8. Lift the package away from the adapter and discard the package.

9. Place the vial on a flat surface. Connect the diluent syringe to the vial adapter by inserting the tip of the syringe into the adapter opening while firmly pushing and turning the syringe clockwise until the connection is secured.

10. Slowly push the plunger rod to inject all the diluent into the BeneFIX vial.

11. With the syringe still connected to the adapter, gently swirl the contents of the vial until the powder is dissolved. Look at the final solution before infusing it. The solution should be clear to colorless. If it is not, throw away the solution and use a new kit.

12. Make sure the syringe plunger rod is still fully pressed down, then turn over the vial. Slowly pull the solution into the syringe. Turn the syringe upward again and remove any air bubbles by gently tapping the syringe with your finger and slowly pushing air out of the syringe. If you reconstituted more than one vial of BeneFIX, remove the diluent syringe from the vial adapter and leave the vial adapter attached to the vial. Quickly attach a separate large luer lock syringe and pull the reconstituted solution as instructed above. Repeat this procedure with each vial in turn. Do not detach the diluent syringes or the large luer lock syringe until you are ready to attach the large luer lock syringe to the next vial adapter.

13. Remove the syringe from the vial adapter by gently pulling and turning the syringe counter-clockwise. Throw away the vial with the adapter attached.

If you are not using the solution right away, you should carefully replace the syringe cap. Do not touch the syringe tip or the inside of the cap.

BeneFIX should be infused within 3 hours after reconstitution. The reconstituted solution may be stored at room temperature prior to infusion.

**INFUSION (Intravenous Injection)**

Continuous infusion is not an approved way to administer BeneFIX. Your doctor or healthcare professional should teach you how to infuse BeneFIX. Once you learn how to self-infuse, you can follow the instructions in this insert.

1. Attach the syringe to the luer end of the provided infusion set tubing.

2. Apply a tourniquet and prepare the injection site by wiping the skin well with an alcohol swab provided in the kit.

3. Insert the butterfly needle of the infusion set tubing into your vein as instructed by your doctor or healthcare provider. Remove the tourniquet. Infuse the reconstituted BeneFIX product over several minutes. Your comfort level should determine the rate of infusion.

Clumping of red blood cells in the tubing/syringe has been reported with the administration of BeneFIX. No adverse events have been reported in association with this observation. To minimize the possibility of clumping it is important to limit the amount of blood entering the tubing. Blood should not enter the syringe.

Note: If red blood cell clumping is observed in the tubing or syringe, discard all material (tubing, syringe and BeneFIX solution) and continue administration with a new package.

4. After infusing BeneFIX, remove the infusion set and discard. The amount of drug product left in the infusion set will not affect your treatment. Dispose of all unused solution, the empty vial(s), and the used needles and syringes in an appropriate container used for throwing away waste that might hurt others if not handled properly.

It is a good idea to record the lot number from the BeneFIX vial label every time you use BeneFIX. You can use the peel-off label found on the vial to record the lot number.

If you have any questions or concerns about BeneFIX, ask your doctor or healthcare provider.

Manufactured by

**Wyeth Pharmaceuticals LLC**
A subsidiary of Pfizer Inc, Philadelphia, PA 19101

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