Coagulation Factor IX (Human)  
AlphaNine® SD

Solvent Detergent Treated/Virus Filtered

DESCRIPTION
Coagulation Factor IX (Human), AlphaNine® SD, is a purified, solvent detergent treated, virus filtered preparation of Factor IX derived from human plasma. It contains a minimum of 150 IU Factor IX activity/mg protein. 

Thrombogenicity of AlphaNine SD in animals is markedly lower than that of Factor IX Complex, Prothrombine® Heat-Treated. Five lots of AlphaNine SD (three lots of virus filtered product and two lots of virus treated product) failed to show any evidence of thrombogenicity when tested directly in the Wessler rabbit stasis model for thrombogenicity at a dose of 200 IU Factor IX/kg body weight. When various lots of AlphaNine SD were further tested at doses between 300 and 650 IU Factor IX/kg, only 5 out of 40 animals (12.5%) showed evidence of disseminated intravascular coagulation (DIC) following infusion. This model exhibited no depletion of coagulation factors, a minimal increase in fibrin monomer, and no protamine test, a slight temporary decrease in platelet counts, and no evidence of intravascular coagulation upon gross autopsy. In contrast, Harrison, et al., report that all Factor IX Complex concentrates administered to dogs were thrombogenic at doses between 50 and 100 IU of Factor IX/kg animal weight.

A clinical evaluation of AlphaNine SD half-life and recovery characteristics was performed. A total of 18 patients with severe to moderate hemophilia B each received a single infusion of 40 to 50 IU Factor IX/kg body weight of AlphaNine SD. Following the administration of AlphaNine SD, the mean half-life of Factor IX observed was approximately 21 hours. This half-life value was computed using the biphasic linear regression model recommended by the International Society for Thrombosis and Haemostasis. The mean recovery observed in clinical trials was approximately 49% of the administered dose.

Coagulation Factor IX (Human), AlphaNine SD in surgery patients and following administration of Factor IX Complex concentrates which contain high amounts of Factor IX activity resulted in a lower level of activation of thrombin on fibrinogen compared to the non-treatment state. These results suggest that the infusion of a high purity Factor IX, such as AlphaNine, may result in a lower level of activation of the coagulation cascade than does Factor IX Complex. The ability of the manufacturing process to inactivate and eliminate virus from the Coagulation Factor IX (Human) product was evaluated at key stages in the process (see Table 1). Known amounts of different viruses were added to samples obtained prior to those steps most likely to reduce virus load (DEAE Chromatography, Solvent Detergent, Dual Affinity Chromatography, and Nanofiltration) in the AlphaNine and AlphaNine SD processes to determine the level of viral inactivation/elimination of these specific steps in the process.

CLINICAL PHARMACOLOGY
AlphaNine SD is a purified formulation of Factor IX containing not less than 150 IU Factor IX activity/mg of total protein.α AlphaNine SD contains non-therapeutic levels of Factor II, Factor VII and Factor X.

Thrombogenicity of AlphaNine SD in animals is markedly lower than that of Factor IX Complex, Prothrombine® Heat-Treated. Five lots of AlphaNine SD (three lots of virus filtered product and two lots of virus treated product) failed to show any evidence of thrombogenicity when tested directly in the Wessler rabbit stasis model for thrombogenicity at a dose of 200 IU Factor IX/kg body weight. When various lots of AlphaNine SD were further tested at doses between 300 and 650 IU Factor IX/kg, only 5 out of 40 animals (12.5%) showed evidence of disseminated intravascular coagulation (DIC) following infusion. This model exhibited no depletion of coagulation factors, a minimal increase in fibrin monomer, and no protamine test, a slight temporary decrease in platelet counts, and no evidence of intravascular coagulation upon gross autopsy. In contrast, Harrison, et al., report that all Factor IX Complex concentrates administered to dogs were thrombogenic at doses between 50 and 100 IU of Factor IX/kg animal weight.

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INSTRUCTIONS
Individuals who receive infusions of blood or blood products may develop signs and/or symptoms of some viral infections. Scientific opinion encourages hepatitis B and hepatitis A vaccinations at birth or diagnosis for patients with hemophilia. Incidences of thrombosis or disseminated intravascular coagulation (DIC), have been reported following administration of Factor IX Complex concentrates which contain high amounts of Factor II, VII and X. Therefore, it is not indicated for the reversal of coumarin anticoagulant-induced hemorrhage, nor in the treatment of hemophilia A patients with inhibitors to Factor VIII.

INDICATIONS AND USE
AlphaNine SD is indicated for the prevention and control of bleeding in patients with Factor IX deficiency due to hemophilia B. AlphaNine SD contains low, non-therapeutic levels of Factors II, VII, X, and Xa, therefore, is not indicated for the treatment of Factor II, VII or X deficiencies. This product is also not indicated for the reversal of coumarin anticoagulant-induced hemorrhage, nor in the treatment of hemophilia A patients with inhibitors to Factor VIII.

CONTRAINDICATIONS
None known.

WARNINGS
Because Coagulation Factor IX (Human), AlphaNine SD is made from pooled human plasma, it may carry a risk of transmitting infectious agents, e.g., viruses, and theoretically, the Creutzfeldt-Jakob disease (CJD) agent. Stringent procedures designed to reduce the risk of adventitious agent transmission have been employed in the manufacture of this product, from the screening of plasma donors and the collection and testing of plasma to the application of viral inactivation/reduction steps such as column chromatography, solvent detergent treatment and nanofiltration in the manufacturing process. Despite these measures, such product can potentially transmit disease, therefore the risk of infectious agents cannot be totally eliminated. The physician should weigh the risks and benefits of the use of this product and should discuss these with the patient. Individuals who receive infusions of blood or plasma products may develop signs and/or symptoms of some viral infections. Scientific opinion encourages hepatitis B and hepatitis A vaccinations at birth or diagnosis for patients with hemophilia. Incidences of thrombosis or disseminated intravascular coagulation (DIC), have been reported following administration of Factor IX Complex concentrates which contain high amounts of Factor II, VII and X. Following administration of Coagulation Factor IX (Human), AlphaNine SD in surgery patients and following administration of Factor IX Complex concentrates which contain high amounts of Factor II, VII and X, always, therefore, is not indicated for the reversal of coumarin anticoagulant-induced hemorrhage, nor in the treatment of hemophilia A patients with inhibitors to Factor VIII. This product is also not indicated for the reversal of coumarin anticoagulant-induced hemorrhage, nor in the treatment of hemophilia A patients with inhibitors to Factor VIII.
Allergic type hypersensitivity reactions, including anaphylaxis, have been reported for all Factor IX products. Frequently these events have occurred in close temporal association with the development of factor IX inhibitors. Patients should be informed of the early symptoms and signs of hypersensitivity reactions, including hives, generalized urticaria, angioedema, chest tightness, dyspnea, wheezing, faintness, hypotension, tachycardia and anaphylaxis. Patients should be advised to discontinue use of the product and contact physician and/or seek immediate emergency care, depending on the severity of the reactions, if any of these symptoms occur.

Nephrotic syndrome has been reported following attempted immune tolerance induction with Factor IX products in Hemophilia B patients with factor IX inhibitors and a history of severe allergic reactions to Factor IX. The safety and efficacy of using AlphaNine SD in attempted immune tolerance induction has not been established.

In Previously Untreated Patients (PUPs), it is possible that anaphylaxis may occur after a median exposure of eleven (11) days. It is recommended that these patients are monitored closely between the tenth and twentieth exposure day.

PRECAUTIONS

General
In order to minimize the possibility of thrombogenic complications, dosing guidelines should be strictly followed. Refer to "Dosage and Administration" section for recommended amount of product to be administered. AlphaNine SD should not be administered at a rate exceeding 10 mL/minute. Rapid administration may result in vasomotor reactions. Nursing personnel and others who administer this material should exercise appropriate caution in handling due to the risk of exposure to viral infection. Discard any unused contents into the appropriate safety container. Discard administration equipment after single use into the appropriate safety container. Do not resterilize containers.

Information for Patients
Patients should be informed of the early symptoms and signs of hypersensitivity reaction, including hives, generalized urticaria, chest tightness, dyspnea, wheezing, faintness, hypotension, and anaphylaxis. Patients should be advised to discontinue use of the product and contact their physician and/or seek immediate emergency care, depending on the severity of the reaction, if any of these symptoms occur.

Some viruses, such as parvovirus B19 or hepatitis A, are particularly difficult to remove or inactivate at this time. Parvovirus B19 may most seriously affect sero-negative pregnant women, or immunocompromised individuals. The majority of parvovirus B19 and hepatitis A infections are acquired by environmental (natural) sources.

Preliminary information suggests a relationship may exist between the presence of major deletion mutations in the Factor IX gene and an increased risk of inhibitor formation and of acute hypersensitivity reactions. Patients known to have major deletion mutations of the Factor IX gene should be observed closely for signs and symptoms of acute hypersensitivity reactions, particularly during the early phases of initial exposure to product.

Pregnancy Category C
Animal reproduction studies have not been conducted with AlphaNine SD. It is also not known whether AlphaNine SD can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. AlphaNine SD should be given to a pregnant woman only if clearly indicated.

Pediatric Use
Clinical trials for safety and effectiveness in pediatric patients 16 years of age and younger have not been conducted. Across a well controlled half-life and recovery clinical trial in patients previously treated with Factor IX concentrates of Hemophilia B, the three pediatric patients receiving AlphaNine SD (solvent detergent treated) responded similarly when compared with 15 adult patients. In an ongoing safety and efficacy clinical trial in patients not previously treated with Factor IX concentrates for Hemophilia B, 21 pediatric patients received AlphaNine SD (solvent detergent treated) similarly when compared with the five adult patients above the age of 16 years. Adverse events were similar in this group compared to the patients above the age of 16 years. Anecdotical evaluation of the results indicates no safety and efficacy differences between pediatric and adult populations.

ADVERSE REACTIONS

The administration of plasma preparations may cause allergic reactions, mild chills, nausea or stingling at the infusion site. For most reactive individuals, slowing the infusion rate relieves the symptoms. For those highly reactive individuals, a different lot may be satisfactory.

Adverse reactions, characterized by either thrombosis or disseminated intravascular coagulation (DIC), have been reported following administration of Factor IX Complex concentrates. Patients who receive Coagulation Factor IX (Human), AlphaNine SD, following operation, or those with known liver disease, should be kept under close observation for potential signs or symptoms of intravascular coagulation. Continued administration should be left to the discretion of the physician. In the clinical study that compared the in vivo half-life and recovery of AlphaNine SD and HT FIX products, no adverse events were associated with 18 infusions of AlphaNine SD administered to 18 individuals with severe to moderate hemophilia B. Short term safety of the earlier version of this product, AlphaNine, was threatened by an absence of adverse events after 225 infusions of this product were received by 31 patients participating in three clinical trials. In the clinical trial to evaluate efficacy of AlphaNine in providing hemostatic protection during and after surgery, 13 patients received a total of 370,655 IU of AlphaNine. In 258 total infusions, each patient received approximately 15,000 IU (range 2,290 to 52,290 IU Factor IX) in an average of 16 infusions (range 2 to 26 infusions). Results from this study showed no bleeding episodes during the course of surgery therapy. There was no hematological evidence (measured by hematocrit, partial thromboplastin time, prothrombin time, fibrinogen/fibrin degradation products, fibrin monomers, D-dimers and platelet counts) of thrombogenicity.

To report SUSPECTED ADVERSE REACTIONS, contact Gelifols at 1-888-GRIFOLS (1-888-474-3657) or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DOSEAGE AND ADMINISTRATION

For adult usage:
AlphaNine SD should be administered intravenously promptly following reconstitution. Administration of AlphaNine SD within three hours after reconstitution is recommended to avoid the potential ill effect of any inadvertent bacterial contamination occurring during reconstitution.

Discard any unused contents into the appropriate safety container.

Each vial of AlphaNine SD is labeled with the total units expressed as International Units (IU) of Factor IX, which is referenced to the WHO International Standard. One unit approximates the activity in one ml of pooled normal plasma.

The amount of AlphaNine SD required to establish hemostasis will vary with each patient and depend upon the circumstances. The following formula may be used as a guide in determining the number of units to be administered:

\[ \text{Body weight (in kg)} \times \text{Desired increase in Plasma Factor IX (percent)} \times 1.0 \text{ IU/kg} = \text{Number of Factor IX IU Required} \]

Example:

- Body weight: 70 kg
- Desired increase in Plasma Factor IX: 40 (% increase)
- Number of Factor IX IU Required: 2,800 IU

In clinical practice there is variability between patients and their clinical response. Therefore, the Factor IX level of each patient should be monitored frequently during replacement therapy.

For pediatric usage: See PRECAUTIONS

Treatment Guidelines for Hemorrhagic Events and Surgery in Patients Diagnosed with Hemophilia B

<table>
<thead>
<tr>
<th>Type of Hemorrhage or Surgical Procedure</th>
<th>Examples</th>
<th>Treatment Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Minor Hemorrhages</td>
<td>Bruises, cuts or scrapes, uncomplicated joint hemorrhage</td>
<td>Fix levels should be brought to at least 20-30% (20-30 IU FIX/kg/twice daily) until hemostasis stops and healing has been achieved (1-2 days).16,17,18,19,20,21</td>
</tr>
<tr>
<td>Moderate Hemorrhages</td>
<td>Nose bleeds, mouth and gum bleeds, dental extractions, hematuria</td>
<td>Fix levels should be brought to 25-50% (25-50 IU FIX/kg/twice daily) until hemostasis has been achieved (2-7 days, on average).16,17,18,19,20</td>
</tr>
<tr>
<td>Major Hemorrhages</td>
<td>Joint and muscle hemorrhages (especially in the large muscles), major trauma, hematuria, intracranial and intraarticular bleeding</td>
<td>Fix levels should be brought to 50% for at least 3-5 days (30-50 IU FIX/kg/twice daily). Following this treatment period, FIX levels should be maintained at 20% (20 IU FIX/kg/twice daily) until healing has been achieved. Major hemorrhages may require treatment for up to 10 days.16,17,18,19,20,21</td>
</tr>
<tr>
<td>Surgery</td>
<td></td>
<td>Prior to surgery, FIX should be brought to 50-100% of normal (50-100 IU FIX/kg/twice daily). For the next 7 to 10 days, or until healing has been achieved, the patient should be maintained at 50-100% FIX levels (50-100 IU FIX/kg/twice daily).16,17,18,19,20,21</td>
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</tbody>
</table>

Dosing requirements and frequency of dosing is calculated on the basis of an initial response of 1% FIX increase achieved per IU of FIX infused per kg body weight and an average half-life of FIX of 18 hours. If dosing studies have revealed that a particular patient exhibits a lower response, the dose should be adjusted accordingly.

For pediatric usage: See PRECAUTIONS

RECONSTITUTION

Use Aseptic Technique

1. Warm diluent (Sterile Water for Injection, USP) and concentrate (AlphaNine SD) to at least room temperature (but not above 37 °C).
2. Remove the plastic flip off cap from the diluent vial.
3. Gently swab the exposed stopper surface with a cleansing agent such as alcohol trying to avoid leaving any excess cleansing agent on the stopper.
4. Open the Mix2Vial® package by peeling away the lid (Figure 1). Leave the Mix2Vial in the clear set, ensuring the Mix2Vial remains attached to the diluent vial.
5. Place the diluent vial upright on an even surface and hold the vial tight and pick up the Mix2Vial in its clear outer packaging. Holding the diluent vial securely, push the blue end of the Mix2Vial vertically down through the diluent vial stopper (Figure 2).
6. While holding onto the diluent vial, carefully remove the clear outer packaging from the Mix2Vial set, ensuring the Mix2Vial remains attached to the diluent vial (Figure 3).
7. Place the product vial upright on an even surface, invert the diluent vial with the Mix2Vial attached.
8. While holding the product vial securely on a flat surface, push the clear end of the Mix2Vial set vertically down through the product vial stopper (Figure 4). The diluent will automatically transfer out of its vial into the product vial. (Note: If the Mix2Vial is connected at an angle, the vacuum may be released from the product vial and the diluent will not transfer into the product vial.)
9. With the diluent and product vials still attached to the Mix2Vial, gently swirl the product vial to ensure the product is fully dissolved (Figure 5). Reconstitution requires less than 5 minutes. Do not shake the vial.

REFERENCES

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10. Disconnect the Mix2Vial into two separate pieces (Figure 6) by holding each vial adapter and twisting counterclockwise. After separating, discard the diluent vial with the blue end of the Mix2Vial.

11. Draw air into an empty, sterile syringe. Keeping the product vial upright with the clear end of the Mix2Vial attached, screw the disposable syringe onto the luer lock portion of the Mix2Vial device by pressing and twisting clockwise. Inject air into the product vial.

12. While keeping the syringe plunger depressed, invert the system upside down and draw the reconstituted product into the syringe by pulling the plunger back slowly (Figure 7).

13. When the reconstituted product has been transferred into the syringe, firmly hold the barrel of the syringe and the clear vial adapter (keeping the syringe plunger facing down) and unscrew the syringe from the Mix2Vial (Figure 8). Hold the syringe upright and push the plunger until no air is left in the syringe. Attach the syringe to a venipuncture set.

14. NOTE: If the same patient is to receive more than one vial of concentrate, the contents of two vials may be drawn into the same syringe through a separate unused Mix2Vial set before attaching to the venipuncture set.

15. Use the prepared drug as soon as possible within three hours after reconstitution.

16. After reconstitution, parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. When reconstitution procedure is strictly followed, a few small particles may occasionally remain. The Mix2Vial set will remove particles and the labeled potency will not be reduced.

17. Discard all administration equipment after use into the appropriate safety container. Do not reuse.

HOW SUPPLIED

AlphaNine SD is supplied in sterile, lyophilized form in single dose vials accompanied by 10 mL diluent (Sterile Water for Injection, USP). Factor IX activity, expressed in International Units (IU) which is referenced to WHO International Standard, is stated on the label of each concentrate vial. AlphaNine SD is packaged with a Mix2Vial filter transfer set for use in administration.

It is available in the following potencies, and the product is also color coded based upon assay on the carton and vial label as follows:

<table>
<thead>
<tr>
<th>Potency</th>
<th>NDC</th>
<th>Assay Color Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>500 IU FIX/10 mL single dose vial</td>
<td>68516-3601-2 or 68516-3607-2</td>
<td>500 IU FIX Range - blue box</td>
</tr>
<tr>
<td>1000 IU FIX/10 mL single dose vial</td>
<td>68516-3602-2 or 68516-3608-2</td>
<td>1000 IU FIX Range - red box</td>
</tr>
<tr>
<td>1500 IU FIX/10 mL single dose vial</td>
<td>68516-3603-2 or 68516-3609-2</td>
<td>1500 IU FIX Range - black box</td>
</tr>
</tbody>
</table>

STORAGE

AlphaNine SD is stable for three years, up to the expiration date printed on its label, provided that the storage temperature is between 2 and 8 °C (36 and 46 °F). Do not freeze to prevent damage to diluent vial. May be stored at room temperature not to exceed 30 °C for 1 month. When removed from refrigeration, record the date removed on the space provided on the carton.

REFERENCES

2. Data on file at Grifols Biologicals Inc.

Manufactured by:
Grifols Biologicals Inc.
5555 Valley Boulevard
Los Angeles, CA 90032, U.S.A.
U.S. License No. 1634
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