

See dosage and indications for Rho(D) Immune globulin administration on reverse

## Indications for administration of Rho(D) Immune globulin (WinRho®SDF)1

Always confirm RhD negative status and draw antibody screen BEFORE administering WinRho®SDF. Testing is required within the previous 14 days.

- NOTE: 1. WinRho®SDF not needed if paternal/sperm donor testing is documented to be RhD negative and weak D negative.
  - 2. Administer within 72 hours of event to ensure effectiveness (if omitted, give as soon as possible, up to 28 days later).
  - 3. WinRho®SDF is a *blood product* that requires a written order and consent form. Recipients should be informed of the risks and benefits of the product and informed consent should be obtained. Refer to pamphlet "The Rh Factor and Pregnancy". All forms are available on the website below or through contacting the Rh Program.
  - 4. Due to the possibility of a reaction to WinRho®SDF, vital signs should be taken pre-administration and recipients advised to stay for 15 to 30 minutes post-injection.
  - 5. Administer by **DEEP IM or IV route**, to ensure adequate absorption. <u>Note:</u> Volumes of 2 mL or less can be given in the deltoid muscle. Volumes greater than 2 mL can be given in the ventrogluteal or vastus lateralus muscles.<sup>2</sup>
  - 6. Injection reporting forms are available from the Rh Program or our website. Please mail or fax completed copy to the Rh Program.
- 28 29 weeks gestation: give 300 mcg. If WinRho®SDF was given within prior 3 weeks, may delay injection for up to 6 weeks later.
- Postpartum when infant is RhD positive, Rh indeterminate or Rh unknown: obtain Kleihauer and give minimum of 120 mcg. May withhold injection if WinRho®SDF has been given within 3 weeks of delivery provided Kleihauer is negative AND passive anti-D antibodies (due to Rho(D) Immune globulin) are detected at delivery.
- Surgical abortion, surgical ectopic or partial molar pregnancy management: up to 12 weeks gestation: minimum 120 mcg; after 12 weeks gestation: 300 mcg.
- Threatened abortion, spontaneous abortion, medical abortion: Less than 8 weeks (56 days) gestation with confident and reliable pregnancy dating may safely withhold blood testing (ABO/Rh and antibody screen) and WinRho®SDF.³ Less than 8 weeks gestation with unreliable dating give minimum of 120 mcg; 8 to 12 weeks gestation give minimum of 120 mcg; after 12 weeks gestation give 300 mcg.
- Antepartum bleeding, placental abruption, placenta previa with bleeding, abdominal trauma, amniocentesis, cordocentesis, chorionic villus sampling (CVS): up to 12 weeks gestation: minimum 120 mcg; after 12 weeks gestation: 300 mcg. For repeat events 6 or more weeks later obtain Kleihauer, antibody screen and give an additional 300 mcg. For repeat events less than 6 weeks later: may withhold WinRho®SDF when Kleihauer is negative AND passive anti-D antibodies (due to Rho(D) Immune globulin) are detected.
- External versions: obtain Kleihauer and give minimum of 120 mcg
- Platelet transfusion if platelet donors are RhD positive: 120 mcg covers up to 6 full buffy coat or apheresed transfused platelet units and protects for up to 4-6 weeks. WinRho®SDF should be administered within 72 hours of the transfusion. Rationale: Platelets from RhD positive donors contain a small amount of red blood cells.
- Transfusion of RhD positive red blood cells (RBC) to RhD negative recipient: 24 mcg per mL red blood cells (RBC's). Caution: see product insert for limitations or consult with the Rh Program or your local blood transfusion service.

## KLEIHAUER TEST DOSING for fetomaternal hemorrhage (FMH) of RhD positive whole blood:

Maternal circulation estimated whole blood volume = 5000 mL. Administer 12 mcg WinRho®SDF per mL of fetal whole blood.

120 mcg protects for FMH of 0% to 0.2% of maternal whole blood volume (0.002 x 5000 mL = 10 mL fetal whole blood x 12 = 120 mcg required)

300 mcg protects for FMH of 0% to 0.5% of maternal whole blood volume (0.005 x 5000 mL = 25 mL fetal whole blood x 12 = 300 mcg required)

Depending on dose calculated above: (1) IM route up to **1200 mcg** every 12 hours or (2) IV route administer up to **600 mcg** every 8 hours until the total dose has been administered. Consult with the Rh Program for further assistance or refer to the product insert under "Dosage and Administration".

Rh Program of Nova Scotia, 5850/5980 University Avenue, PO Box 9700, Halifax, NS B3K 6R8 Tel: (902) 470-6458 Fax: (902) 470-7468 Website: http://rcp.nshealth.ca/rh REV AUG2022

 $<sup>^{\</sup>rm 1}$  Prevention of Rh Alloimmunization. No. 133, Reaffirmed Guidelines. JOGC January 2018.

<sup>&</sup>lt;sup>2</sup> Perry & Potter. Clinical Nursing Skills & Techniques. Elsevier Mosby 10<sup>th</sup> edition, 2021

<sup>&</sup>lt;sup>3</sup> Guideline on Rh Prophylaxis before 8 weeks (56 days) gestation for Early Pregnancy Complications and Medical Abortions. Rh Program of NS June 2022.