

This document is to help guide the use of the provided GRH IV Iron package. The documents included in the IV Iron Sucrose Package are:

### 1. Adult Outpatient Iron Sucrose Order set (page 2 and 3)

Use this document to help determine:

- (1) If the patient is a candidate for IV Iron therapy
- (2) If the IV Iron therapy will be paid for by GRH or by the patient

AND

to order IV Iron to be administered at GRH. (Note: if the patient is to pay for the IV Iron they will also require a separate outpatient prescription)

### 2. Form 1: Facilitating Patient Payment for IV Iron (page 4)

This document provides guidance for payment options for patients that have to pay for their own Iron Sucrose.

### 3. Form 2: IV Iron Sucrose EAP request form (page 5)

This document can be used for patients who are ODB eligible to request Exceptional Access Program Coverage.

We suggest keeping these documents for your records to help you with ordering IV Iron for patient's in the future, however if you need a new package or any of the forms listed above please contact Medical Day Unit at Grand River Hospital at 519-749-4300 ext 2126.

# ROUTINE ORDERS Adult Outpatient Intravenous Iron Order Set

| 2) An order with a black box       will be activated UNLESS the prescriber crosses out the complete order with a line and initials.         Date:       year/month/day       Time:       Weight (kg):       Allergies: □ None □Yes<br>Review electronic record       beget   |
|--|
| Date:       Time:       Weight (kg).       Review electronic record <sup>a</sup> g to the second t |
| Required Criteria for Outpatient Administration of Intravenous Iron at GRH – Must be complete and<br>attach laboratory reports for 1. and 2. to book appointment.         All of the following criteria must be met. Check all that apply.         1. Diagnosis of iron deficiency anemia: Hemoglobin (Hgb) level less than 120g/L in females or less than 130g/L in<br>males AND         2. Low iron stores as demonstrated by: transferrin saturation (TSAT) less than 20% (0.20) AND/OR ferritin less than<br>15 mcg/L AND         3. Insufficient time (4 weeks or less) to evaluate efficacy of oral therapy for upcoming procedure (e.g prior to surgery)<br>OR documented intolerance/inadequate response to appropriate trial of oral therapy OR inability to absorb oral iron         Eligibility for insured services - Must be complete to book appointment         GRH Pays – intravenous iron will be supplied at the appointment – does not apply to iron isomaltoside.<br>Iron isomaltoside is patient pays only. (IF administration of intravenous iron is an adjunct therapy for an INSURED<br>HOSPITAL SERVICE, such as a surgical procedure, diagnostic test or treatment) Provide details and<br>date:         Patient Pays - patient must bring intravenous iron to appointment (IF sole reason for outpatient visit is<br>intravenous iron administration, even if being treated for iron-deficiency anemia and meets criteria above).   |
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|  |
| RETENDED ON THE AUTOMIC FURTING VID 12/42-422/110 INVESTORE DATED TO DOTO DOTO IS  |
| □ Patient has been provided with outpatient prescription for intravenous iron  |
| □ EAP application submitted on (date)  |
| Lab work and Diagnostics   |
| CBC, Ferritin at final scheduled appointment   |
| Iron Studies at final scheduled appointment  |
| IV fluid   |
| ■ Peripheral saline lock, if needed  |
| ■ Sodium Chloride 0.9% 250mL at 150mL/hr Medication  |
| Premedication (consider if patient has had reaction during previous iron infusion)   |
| DiphenhydrAMINE 50mg IV x 1 $\blacksquare$ prn for reaction $\square$ pre infusion   |
| $\Box$ Hydrocortisone 100mg IV x 1 $\Box$ prn for reaction $\Box$ pre infusion   |
| □ DimenhyDRINATE 50mg IV x 1 □ prn for reaction □ pre infusion   |
| □ Acetaminophen 1000mg PO x 1 □ prn for reaction □ pre infusion  |
|  |
| Intravenous Iron – See reverse for dosing. If more than a course of therapy is needed (based on  |
| maximum dose), after completion of the course of therapy a new order must be submitted with new blood work meeting the above criteria  |
|  |
| □ Specify type of intravenous iron (iron sucrose or iron gluconate complex):   |
| Dosemg IV every (frequency) x doses (Maximum 6 doses/course) <b>OR</b><br>□ Iron isomaltoside (see reverse for dosing chart) total dose per coursemg (Maximum 1  |
| course per order.) Give iron isomaltosidemg IV x 1dose and (if divided dose required)  |
| give mg IV x 1 dose at least 7 days after the first dose (maximum 1000mg per single dose)  |
| giveing iv x i dose al least / days aller the inst dose (maximum looping per single dose)  |
| Monitoring   |
| Monitor for signs and symptoms of hypersensitivity reactions for at least 30 minutes post infusion and<br>until clinically stable.   |
| *Enter Order # and initial (by Nurse/Clerical)   |
| Prescriber Signature:  |
| Transcriber Signature:Date:Date:   |

Date:



| Calculating Iron Replacement Requirements                |  |  |  |
|--|--|--|--|
| Normal Hgb; Women: Greater than 120g/L Men: Gre          | ater than 130g/L   |  |  |
| Hgb deficit (g/L) = target Hgb – actual Hgb              | Deficit =  |  |  |
| Total iron dose required (mg) = (Hgb deficit x 20) + 500 | <b>Total Iron requirements=</b><br>Divide total iron requirement by intravenous iron dose to<br>determine number of infusions. |  |  |

| Intravenous Iron P                        | rescribing Guidelin   | es (See GRH IV manual  | or Product Monograph for  | more information)  |  |  |
|---|---|--|---|--|--|--|
| IV Iron Sucrose<br>(Venofer®)             | and maximum dos   | ed doses with a<br>n daily dose of 300mg<br>e of 1000mg in 14 days<br>of therapy per order – | Dosage regimen once per week but can give<br>multiple doses within a week in certain<br>circumstances (preferable 2 to 3 days between<br>doses)<br>Consider initiating at lower doses for special patient<br>populations such as elderly, pregnant women and<br>renal patients to reduce infusion reactions |  |  |  |
| IV Iron Gluconate<br>Complex (Ferrlecit®) | single dose: 250mg<br>Maximum course<br>6 doses                                 | ximum recommended<br>g<br>of therapy per order –   |   |  |  |  |
| Iron Isomaltoside                         | Hemoglobin (g/L) Total Iron Dose – Maximum dose per course of therapy per order |  |   |  |  |  |
| (Monoferric®)                             |   | Body weight less than<br>50kg  | Body weight 50 to 69 kg   | Body weight 70kg or<br>greater   |  |  |
|   | 100 or greater  | 500mg  | 1000mg  | 1500mg<br>(given in 2 divided<br>doses of 1000mg +<br>500mg 7 days apart)  |  |  |
|   | Less than 100   | 1000mg<br>(given in 2 divided<br>doses of 500mg +<br>500mg 7 days apart)                     | 1500mg<br>(given in 2 divided doses<br>of 1000mg + 500mg 7<br>days apart)   | 2000mg<br>(given in 2 divided<br>doses of 1000mg +<br>1000mg 7 days apart) |  |  |

| Guidance for outpatient p  | rescription    |                           |               |                   |    |
|----------------------------|----------------|---------------------------|---------------|-------------------|----|
| When providing an outpatie | nt prescriptio | n please include as follo | ows:          |                   |    |
| Intravenous type/brand     | Dose           | mg (dose) every           | (frequency) x | (number of doses) | 11 |

# Form 1: Facilitating Patient Payment for IV Iron

For patients obtaining their own supply of IV Iron for administration at the Medical Day Unit at Grand River Hospital there are 4 potential options. See below for pricing and information regarding Health Care Centre Pharmacy dispensing.

#### 1. Private insurance

Patients contact their private insurance provider to determine if they are eligible to have IV Iron dispensed through their plan. The patient must do this on their own, but may need the drug identification number listed here **Iron Sucrose DIN: 02243716** 

Iron Isomaltoside DIN: 02477777

### 2. Exceptional Access Coverage – Only avaible for IV Iron Sucrose

Physicians can apply for exceptional access for all ODB patients (including those on Trillium) for **IV iron sucrose therapy**. The Exceptional Access Form (Form 2) has been attached or can also be accessed from the Medical Day Unit at Grand River Hospital.

#### 3. Patients pay cash

Patients can pay cash at their own community pharmacy or Health Care Centre Pharmacy at the hospital for their IV iron and pick the dose up prior to their scheduled appointment

### 4. Special considerations

For patients that don't have private or EAP coverage but who are unable to afford their IV iron, we will discuss these cases on an individual basis to determine the best course of action. Please contact the Clinical Manager, Medical Day Unit; 519-749-4300 ext 3956.

#### **Health Care Centre Pharmacy Information**

#### Cost per dose of medication for cash paying patients (prices are subject to change)

| Iron Sucrose (Venofer) | Approximate |  |  |
|------------------------|-------------|--|--|
| Dose                   | Cost        |  |  |
| 100mg                  | \$53        |  |  |
| 200mg                  | \$96        |  |  |
| 300mg                  | \$140       |  |  |
| 400mg                  | \$183       |  |  |
| 500mg                  | \$227       |  |  |

| Iron Isomaltoside<br>(Monoferric) | Approximate<br>Cost |  |  |
|-----------------------------------|---------------------|--|--|
| 500mg                             | \$274               |  |  |
| 1000mg                            | \$535               |  |  |

- Monoferric requires fewer visits to infuse the same amount of iron.
- Most private drug plans currently cover Monoferric without any prior authorization requirement

#### Reasons to use HCCP for IV Iron

- Convenience pick up your Iron on the way to your appointment
- Supply HCCP will always have supply of IV Iron available for our Medical Day Unit Patients
- Quick and friendly service HCCP will only need 30 minute notice to fill your IV Iron prescription

# FORM 2: IV iron sucrose EAP request form

To be completed and submitted for Ontario Drug Benefit (ODB) patients (e.g. over 65 years, on social assistance, or covered through Trillium Drug Program)

# Exceptional Access Program (EAP) Request for Iron Sucrose (Venofer) for the Treatment of Iron-Deficiency Anemia

Fax the completed form and/or any additional relevant information to (416) 327-7526 or toll free to 1-866-811-9908; OR send to EAPB Ontario Public Drug Programs, Exceptional Access Program Branch, 3<sup>rd</sup> Floor, 5700 Yonge Street, Toronto, ON, M2M 4K5

| ection 1 – Prescriber Information               |   |   | Sect                                    | Section 2 – Patient Information |  |                                  |   |  |
|---|---|---|---|---------------------------------|--|----------------------------------|---|--|
| rst name  | Initial   | Last name                                     |   | First n                         | ame  | Initial                          | Last name   |  |
| ailing Address                                  |   |   |   | Health                          | Number   |                                  |   |  |
| reet no. Street name                            |   |   |   |                                 |  |                                  |   |  |
|   |   |   |   |                                 |  |                                  |   |  |
| ty  |   | Pos   | stal code                               |                                 |  |                                  |   |  |
| IX NO.  |   | Telephone no.                                 |   | Date o                          | f birth (yyyy/mm/dd)   |                                  |   |  |
| )   |   | ( )   |   |                                 |  |                                  |   |  |
| New request                                     |   | enewal of existin                             | g EAP approv                            | al (specify                     | EAP#)  |                                  |   |  |
| Section 3 – Drug,                               | Dose and Regin  | nen Requested                                 |   |                                 |  |                                  |   |  |
| Drug product: Iror                              | sucrose (Veno   | fer) 100mg/5mL                                | vial(s)                                 |                                 |  |                                  |   |  |
| Dose:   |   |   |   |                                 |  |                                  |   |  |
| Frequency:                                      |   |   |   |                                 |  |                                  |   |  |
| Number of doses:                                |   |   |   |                                 |  |                                  |   |  |
|   |   |   |   |                                 |  |                                  |   |  |
|   |   |   |   |                                 | e following results indi   | cated belo                       | w)  |  |
| Diagnosis of iro                                | n-deficiency and  | emia has been c                               | onfirmed with                           | n docume                        | nted bloodwork   |                                  |   |  |
| Hemoglobin:                                     | g/L MCV:  | fl  | L Date collec                           | ted:                            |  |                                  |   |  |
| If Hemoglobin less                              | than 120 g/L in   | females or less                               | than 130 g/l                            | in males o                      | or MCV less than 75fL or   | greater th                       | an 120fl , provide the                                      |  |
| following:                                      | 2008/201  |   |   |                                 |  | 8. eater th                      | an 11011) provide the                                       |  |
| Date Drawn                                      | L   | evel  | Date Dra                                | awn                             |  | Level                            |   |  |
|   | Ferritin  | mcg   | /L                                      |                                 | Serum Iron Levels  |                                  | mcg/dL  |  |
|   | TSAT  | -   | %                                       |                                 | Total iron binding   |                                  | mcg/dL  |  |
|   |   |   |   |                                 | capacity (TIBC)  |                                  |   |  |
| Section 5 – Medic                               | ation: Current a  | and/or Previous                               |   |                                 |  |                                  |   |  |
| Patient has alread                              | ady been treate   | d with at least o                             | ne iron produ                           | ict as sum                      | marized below:   |                                  |   |  |
| Medication and                                  | Name  | Dose  | Dose                                    |                                 | rt Date  | Dura                             | tion  |  |
| Oral Iron                                       |   |   |   |                                 |  |                                  |   |  |
| Oral Iron                                       |   |   |   |                                 |  |                                  |   |  |
| IV Iron   |   |   |   |                                 |  |                                  |   |  |
| AND   |   |   |   |                                 |  |                                  |   |  |
| <ul> <li>Patient has dem</li> <li>OR</li> </ul> | onstrated intole  | erance to oral in                             | on therapy                              |                                 |  |                                  |   |  |
| -   | corporded to an   | a adaguata thar                               | ny with oral                            | iron                            |  |                                  |   |  |
| Patient has not<br>The information on the       |   |   | 1.7                                     |                                 | ation Protection $\Delta ct = 2001 \le 0$  | 2001 63 50                       | hed. A (PHIPA) and Section 13 of                            |  |
| the Ontario Drug Ben<br>Information Practices   | <i>efit Act,</i> R. S. O. 199<br>", which may be acc<br>esk at 1800-668-664 | 0c.O.10 and will be u<br>essed at ww.health.g | sed in accordance<br>gov.on.ca If you h | e with PHIPA<br>ave any ques    | , as set out in the Ministry of H<br>tions about the collection or u<br>ram Branch (EAPB), Ministry of | ealth and Lon<br>se of this info | g-Term Care "Statement of<br>rmation, call the Ontario Drug |  |
|   |   |   |   |                                 |  | _                                |   |  |
| Prescriber signature (                          | mandatory)  |   |   | (PSO numb                       | CPSO number  |                                  | Date  |  |
|   |   |   |   | ci so numb                      | er   | Date                             |   |  |