

**Patient Demographics**

_____ Last Name	_____ First Name	_____ Middle Initial	_____ Social Security Number
_____ Date of Birth	_____ Phone Number	_____ Insurance Provider	_____ ID Number

**Provider Information**

_____ Referring Provider Name	_____ Phone Number	_____ Practice Contact (Name)
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**Clinical Information**

_____ Diagnosis	_____ ICD-10 code	_____ Allergies:
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**Prescriber's Orders: orders expire after 12 months**

<input type="checkbox"/>	<b>Darbepoetin (ARANESP)</b> , ____ mcg/kg, Subcutaneous, Every ____ Weeks, for ____ treatments
<input type="checkbox"/>	<b>Epoetin alfa</b> ____ Units/kg, Subcutaneous, 3 times weekly, for ____ treatments Or
<input type="checkbox"/>	<b>Epoetin alfa</b> ____ Units, Subcutaneous, Every ____ Weeks, for ____ treatments Note: orders for epoetin alfa will be automatically interchanged to the preferred biosimilar
<input type="checkbox"/>	<b>Ferric carboxymaltose (INJECTAFER)</b> 750 mg, Intravenously over 15 minutes, Weekly, For 2 doses
<input type="checkbox"/>	<b>Ferric carboxymaltose (INJECTAFER)</b> 1000 mg, Intravenously over 15 minutes, Once
<input type="checkbox"/>	<b>Ferric derisomaltose (MONOFERRIC)</b> 1000 mg, Intravenously over 30 minutes, Once
<input type="checkbox"/>	<b>Iron sucrose (VENOFER)</b> ____ mg <b>total</b> dose; space doses over ____ day(s)
<input type="checkbox"/>	<input type="checkbox"/> 200 mg, Intravenous over 15 minutes for ____ doses <input type="checkbox"/> 400 mg, Intravenous over 2.5 hours for ____ doses <input type="checkbox"/> 300 mg, Intravenous over 90 minutes for ____ doses <input type="checkbox"/> 500 mg, Intravenous over 3.5 hours for ____ doses
<input type="checkbox"/>	<b>Romiplostim (NPLATE)</b> 1 mcg/kg, Subcutaneous, Weekly, for ____ treatments (max 6 months)
<input type="checkbox"/>	<b>Sutimlimab (ENJAYMO)</b> ____ g, Intravenously over 60 minutes, Weekly, for 2 doses, followed by ____ g, Intravenously over 15 minutes, Every 2 Weeks, for ____ doses

<b>Additional Order Comments:</b>	
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_____ Physician Signature	_____ Printed name	_____ Date/time
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In order to refer a patient to one of the Baptist Infusion Centers, please send the following information:

- Patient's complete demographics, including insurance information
  - A copy of the patient's diagnosis with the appropriate ICD-10 diagnosis code, preferably in the provider's note
  - A copy of the most recent labs and the most recent provider's progress note
  - A copy of the Baptist Cancer Center order sheet with the correct drug selected and the order signed, dated, and timed
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- **Iron Agents:** if referring for iron replacement, a progress note must include failure of, or intolerance to, oral iron products (oral iron failure documentation is not required for CHF or CKD patients).
    - Labs- iron studies completed within 90 days; hematocrit (Hct) and hemoglobin (Hgb) completed within 30 days.
  - **ESAs:** if referring for Erythropoiesis-Stimulating Agents, we will need documentation of the patient's diagnosis of anemia and chronic renal insufficiency.
    - Labs- hematocrit (Hct) and hemoglobin (Hgb) completed within 30 days of initiation
  - **Romplostim (Nplate):** will need documentation of patient's immune thrombocytopenia diagnosis
    - Labs- platelet levels completed within 30 days of initiation
  - We will not proceed with prior authorization or scheduling the patient for treatment until all information requested is provided
  - Please ensure that we have a valid contact name and number should we need to call for additional information or to clarify orders
  - Please note that orders are only valid for 12 months. After that time, new orders and new documentation must be provided.

Thank you for allowing Baptist Cancer Center to care for your patients.