IV Iron Prescribing Checklist	PATIENT LABEL
	UR No.:
INDICATION	Name:
*Confirmed Iron Deficiency Anaemia AND:	D.O.B: Sex:
*See Clinical Update on Iron Deficiency Anaemia MJA 2010	Doctor:Ward:
 □ Short time to non-deferrable surgery associated with substantial blood loss □ Rapid iron repletion clinically important to prevent decompensation or transfusion □ Demonstrated intolerance to oral iron despite modification of dose & frequency (eg. to alternate daily) □ Demonstrated non-compliance with oral iron □ Demonstrated lack of efficacy with therapeutic doses of oral iron: 100 – 200 mg of elemental iron per 	
day eg. 1 or 2 tablets a day of Ferro-tab, Ferro-f-tab, Ferrograd, Fefol, FGF or Maltofer or equivalent) Ongoing iron (blood) losses exceeding absorption Malabsorption of iron Absolute or functional iron deficiency in chronic heart failure (as per local or national guidelines)	
☐ Absolute or functional iron deficiency in chronic kidney disease (as per local renal unit guidelines) Details re indication:	
Contraindications	
seek advice regarding interpretation and if cause of anaemia is unclear)	
Evidence of iron overload or disturbances of iron utilisation including haemochromatosis	
Known hypersensitivity to IV or IM iron: discuss indication, alternatives & choice of IV iron preparation (if indicated) with an expert such as haematologist, nephrologist, gastroenterologist or other specialist	
Previous IM or IV iron NONE	
Precautions	
Significant liver dysfunction (discuss risks / benefits with gastroenterologist), avoid in patients with hepatic dysfunction where iron overload is a precipitating factor, in particular porphyria cutanea tarda	
☐ Use with caution in acute or chronic infection after assessing risks / benefits & seek expert advice. Avoid during active systemic infection / bacteraemia.	
Use with caution in asthma, eczema or atopic allergies, consider in hospital use – seek expert advice	
In pregnancy seek expert advice (risks, benefits, timing, fetal monitoring); avoid in 1st trimester; give in hospital	
 Not recommended in children: refer to product's PI, health service guidelines & seek expert paediatric advice Avoid paravenous leakage which may cause irritation and potentially permanent brown skin staining. Distant skin discolouration has also been reported. 	
☐ IV iron (particularly ferric carboxymaltose) can cause hypophosphataemia - see PI for precautions / risks See product's PI for other precautions such as lactation, fertility, inflammatory disease, effects on lab tests	
 IV iron can cause hypersensitivity reactions (including anaphylactoid, fetal bradycardia & acute allergic coronary arteriospasm with infarction), which may be fatal & can occur after previous uneventful doses. Cardiopulmonary resuscitation facilities & trained staff MUST be available. STOP immediately if signs of allergy, intolerance or paravenous leakage. Observe for at least 30 min after each administration. Regular monitoring of FBE & iron studies for recurrent iron deficiency & for iron overload is required. Assess underlying cause in ALL patients – refer to Clinical Update on Iron Deficiency Anaemia MJA 2010. ALWAYS consult the product's full PI for further details & updates, seek expert advice when required. 	
Patient IV iron LEAFLET given www.sahealth.sa.gov.au/bloodsafe or other	
Completing MO	hile/Dager:
Name:	_
BloodSafe Resource Version 1.2 26/6/21© Department for Health an	_