

Consensus Statement

International consensus statement on the peri-operative management of anaemia and iron deficiency

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Summary

Despite current recommendations on the management of pre-operative anaemia, there is no pragmatic guidance for the diagnosis and management of anaemia and iron deficiency in surgical patients. A number of experienced researchers and clinicians took part in an expert workshop and developed the following consensus statement. After presentation of our own research data and local policies and procedures, appropriate relevant literature was reviewed and discussed. We developed a series of best-practice and evidence-based statements to advise on patient care with respect to anaemia and iron deficiency in the peri-operative period. These statements include: a diagnostic approach for anaemia and iron deficiency in surgical patients; identification of patients appropriate for treatment; and advice on practical management and follow-up. We urge anaesthetists and peri-operative physicians to embrace these recommendations, and hospital administrators to enable implementation of these concepts by allocating adequate resources.

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Recommendations for best clinical practice

- 1 Physicians should consider pre-operative anaemia and iron deficiency as an indication for a peri-operative care pathway that stretches from the decision to operate until complete recovery from surgery.
- 2 The presence of anaemia should be investigated in all surgical procedures with expected moderate-to-high blood loss (> 500 ml).
- 3 Serum ferritin level < 30 $\mu\text{g.l}^{-1}$ is the most sensitive and specific test used for the identification of absolute iron deficiency. However, in the presence of inflammation (C-reactive protein > 5 mg.l^{-1}) and/or transferrin saturation < 20%, a serum ferritin level < 100 $\mu\text{g.l}^{-1}$ is indicative of iron deficiency.
- 4 Major, non-urgent surgery should be postponed to allow the diagnosis and treatment of anaemia and iron deficiency.
- 5 When treating anaemia pre-operatively, the target haemoglobin concentration should be $\geq 130 \text{ g.l}^{-1}$ in both sexes, to minimise the risk of transfusion-associated unfavourable outcomes
- 6 Oral iron replacement should be targeted to patients with iron deficiency with or without anaemia whose surgery is scheduled 6–8 weeks after diagnosis, preferably by the primary care physician (General Practitioner).
- 7 Daily (40–60 mg) or alternate-day (80–100 mg) treatment with oral iron and nutritional advice should be initiated immediately in patients with iron deficiency and no contra-indications.
- 8 Sufficient data exist to support intravenous iron as efficacious and safe. Intravenous iron should be used as front-line therapy in patients who do not respond to oral iron or are not able to tolerate it, or if surgery is planned for < 6 weeks after the diagnosis of iron deficiency.
- 9 The diagnosis and treatment of anaemia and iron deficiency should commence as early as possible in the peri-operative period, and ideally as soon as the decision to undertake surgery is made.

Why was this consensus statement developed?

Several guidelines for the management of peri-operative anaemia have been published over the last decade [1–11]. However, for anaemia and iron deficiency in adult surgical patients, there are a number of non-evidence based misconceptions regarding prevalence, consequences, diagnosis and treatment, as well as inconsistency of terminology and lack of clear guidance on clinical pathways. Throughout surgical practice, there has been increased emphasis on speed to operation, and many pre-assessment clinics focus on the ability to facilitate same-day hospital admissions, which may overlook potential opportunities to optimise patients, and improve fitness for surgery.

How does this consensus statement differ from other available guidelines?

There are a number of guidelines from professional associations recommending ‘what to do’ [1–11]. The aim of this document is to utilise the recommendations therein, and provide a working practice document on ‘how to’ feasibly introduce these guidelines into clinical practice; or better stated, how to do it. Therefore, our goal is to provide independent, collective guidance and a pragmatic, clear clinical pathway for the diagnosis and treatment of peri-operative iron deficiency and anaemia in adult surgical patients, in order to improve outcomes in a cost-effective manner.

Iron deficiency

Iron is the most common and widespread nutritional deficiency, even in industrialised countries, and affects

approximately two billion people worldwide [12, 13]. Iron deficiency, with or without anaemia, is associated with chronic conditions such as cancer (43% across different tumours), inflammatory bowel disease (45%), chronic kidney disease (24–85%), chronic heart failure (43–100%) and other chronic inflammatory diseases [14].

Apart from its well-recognised role in erythropoiesis, iron is also a critical component in many important cellular processes such as oxygen transport, electron transfer reactions, mitochondrial respiration, gene regulation and cellular immunity [15, 16]. Functional iron deficiency in critically ill patients is associated with an increased duration of systemic inflammatory response syndrome and prolonged ICU stay [17]. In congestive heart failure, iron deficiency has been independently associated with compromised physical performance and quality of life [18], as well as an increase in all-cause and cardiovascular mortality [19]. Treatment of iron deficiency with i.v. iron may improve functional status within 4 weeks, which has been shown to be maintained after 24 and 52 weeks [20, 21].

Importantly, in a large series of non-cardiac surgical patients (n = 2115; 48% women), the prevalence of anaemia was 34%, and absolute iron deficiency and iron sequestration (see below) was responsible for 75% of the cases of anaemia (541/715) [22]. In addition, before major orthopaedic or abdominal surgery, low ferritin levels have been associated with increased rates of postoperative infections [23, 24].

For a 70-kg man, total body iron is about 3500 mg (50 mg.kg⁻¹ body weight). Most of the iron in the body is distributed in haemoglobin (Hb) within red blood cell (65%; 2300 mg). Approximately 10% is found in muscle fibres (in myoglobin) and other tissues (in enzymes and cytochromes) (350 mg). The remaining iron is stored in the liver, macrophages and bone marrow (850 mg) [25]. The amount of iron required for the daily production of 300 billion red blood cells (20–30 mg) is provided mainly by macrophages recycling iron from senescent red blood cells (RBC), while daily iron absorption (1–2 mg) balances daily losses. Increased iron requirements, limited external supply and increased blood loss may lead to progressive iron deficiency

and, subsequently, iron deficiency anaemia [25] (Table 1). Thus, different types (stages) of iron deficiency are found in surgical patients (Table 2), classified as follows:

- Inadequate (low) iron stores; this refers to the body's inability to sustain erythropoiesis to recover from blood loss at operation. This may be indicated by the serum ferritin level; in a healthy adult, 1 ng.l⁻¹ ferritin reflects approximately 8 mg stored iron [26]. As an example, a patient with a pre-operative ferritin < 100 ng.l⁻¹ may not have enough iron reserves to recover from a 30–40 g.l⁻¹ Hb drop, while maintaining normal iron stores (ferritin > 30 ng.l⁻¹).
- True (absolute) iron deficiency without anaemia refers to a reduction in total body iron with normal Hb, as levels of erythroid iron are still sufficient for erythropoiesis. This condition is often seen in women of child-bearing age.
- Iron deficiency anaemia refers to a more severe condition in which decreased iron stores result in decreased Hb and usually microcytic hypochromic red cells (mean corpuscular volume < 80 fl; mean corpuscular haemoglobin < 27 pg). The corrected

Table 1 Main causes of iron deficiency.

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| A. Increased demand |
| <ul style="list-style-type: none"> • Growth during infancy and childhood • Treatment with erythropoiesis-stimulating agents |
| B. Limited supply |
| <ul style="list-style-type: none"> • Poor intake • Inappropriate diet with deficit in bioavailable iron and/or ascorbic acid • Malabsorption <ul style="list-style-type: none"> – Gastric resection – <i>Helicobacter pylori</i> infection (even without significant bleeding) – Malabsorption syndromes (Crohn's disease and celiac disease) • Drug interference (gastric anti-acid agents and antisecretory drugs) |
| C. Increased losses |
| <ul style="list-style-type: none"> • Phlebotomy <ul style="list-style-type: none"> – Blood donation – Dialysis (particularly haemodialysis) • Haemorrhage <ul style="list-style-type: none"> – Surgery – Trauma – Gastrointestinal bleeding – Genitourinary bleeding – Respiratory tract bleeding |

Table 2 Characteristics of the different stages of iron deficiency in surgical patients.

Iron status	Laboratory findings	Initial treatment	Adjuvant treatment
Normal	Ferritin 30–300 $\mu\text{g.l}^{-1}$ TSAT 20–50% CRP < 5 mg.l^{-1}	None	None
Low iron stores (for surgery with moderate-to-high blood losses)	Ferritin < 100 $\mu\text{g.l}^{-1}$	Low dose oral iron (40–60 mg iron per day or 80–100 mg alternate days, 6–8 weeks)	Intravenous iron, if intolerance, contraindication or no response to oral iron or short time to surgery
Iron deficiency	Ferritin < 30 $\mu\text{g.l}^{-1}$	Low dose oral iron (40–60 mg iron per day or 80–100 mg alternate days, 6–8 weeks)	Gastro-intestinal/urological investigations Intravenous iron, if intolerance, contraindication or no response to oral iron or short time to surgery
	*Ferritin 30–100 $\mu\text{g.l}^{-1}$ TSAT < 20% and/or CRP > 5 mg.l^{-1}	Intravenous iron (Calculated dose)†	Treat underlying chronic disease if possible Recombinant erythropoietin in the selected case if anaemia does not respond to i.v. iron alone
Functional iron deficiency*	Ferritin 100–500 $\mu\text{g.l}^{-1}$ TSAT < 20%	Intravenous iron (Calculated dose)†	Revise erythropoietin dose Check folic acid and vitamin B ₁₂
Iron sequestration*	Ferritin > 100 $\mu\text{g.l}^{-1}$ TSAT < 20% and/or CRP > 5 mg.l^{-1}	Recombinant erythropoietin, if anaemia	Intravenous iron, if functional iron deficiency develops

CRP, C-reactive protein; Fe, elemental iron; TSAT, transferrin saturation index.

*Low reticulocyte Hb content (CHr < 28 pg), high proportion of hypochromic red cells (HYPO > 5%), low RBC size factor (RBCSF < 88 fl) or ferritin index > 2 will confirm a component of iron deficiency.

†Total iron deficiency [mg] = (Target Hb – Baseline Hb [g.l^{-1}]) · 0.24 · Body weight [kg] + 500. A simplified calculation is 200 mg per each 10 g.l^{-1} of Hb deficit respect to target Hb + 500 mg for stores.

reticulocyte count provides an estimate of the rate of effective marrow production compared with normal (set as a production index of 1). A production index greater than 2 is not compatible with iron deficiency anaemia.

- Functional iron deficiency: this refers to insufficient mobilisation of iron from stores due to increased demands, in the presence of normal or elevated iron stores. It is clearly defined in renal patients following treatment with erythropoiesis-stimulating agents [15].
- Iron sequestration: this refers to decreased iron mobilisation from the stores in the liver and the macrophages to meet daily bone marrow requirements. This is caused by inflammation, which up-regulates hepatic hepcidin synthesis and may decrease the synthesis and activity of erythropoietin. Hpcidin binds to ferroportin, the only known iron-exporting protein in humans, and promotes its internalisation and degradation, resulting in inhibition of intestinal iron absorption and iron

sequestration at liver and macrophages [25] (Fig. 1). This may evolve to true iron deficiency in patients with inflammation and chronic blood losses.

Diagnosis of iron deficiency

Serum ferritin and transferrin saturation (TSAT) are commonly used for an initial evaluation of iron status. Serum ferritin level is the most widely-used test for evaluating iron stores, while TSAT reflects iron availability for erythropoiesis (TSAT < 20% indicates insufficient iron supply to support normal erythropoiesis). C-reactive protein (CRP) is a marker of inflammation which is useful when iron-restricted erythropoiesis due to iron sequestration is suspected [25, 26]. Thus:

- A serum ferritin level < 30 $\mu\text{g.l}^{-1}$ is the most sensitive (92%) and specific (98%) cut-off level for the identification of true iron deficiency (with or without anaemia); no further laboratory work-up is needed (Fig. 2) [15, 25, 27].

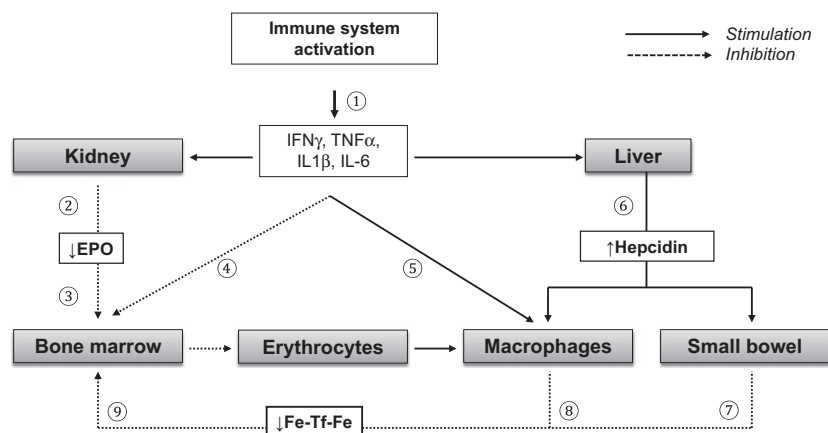


Figure 1 Effects of inflammation on iron metabolism and erythropoiesis. 1, release of immune and inflammatory cytokines; 2, decreased erythropoietin (EPO) production; 3, decreased erythropoietic stimulation; 4, inhibition of erythroid cell proliferation; 5, augmented erythrophagocytosis; 6, IL-6 induced hepcidin release; 7 & 8, decreased ferroportin-mediated release of iron from enterocytes (inhibition of iron absorption) and macrophages (iron sequestration), leading to decreased transferrin-bound iron; 9, decreased iron availability for erythropoiesis.

- In the presence of inflammation (CRP > 5 mg.l⁻¹), and/or TSAT < 20%, ferritin < 100 µg.l⁻¹ strongly suggests iron deficiency (see below) (Fig. 2). It also indicates inadequate iron stores for surgery during which moderate-to-high blood loss is expected.
- In contrast, TSAT < 20% and ferritin > 100 µg.l⁻¹ usually indicates iron sequestration, as seen in inflammatory conditions (CRP > 5 mg.l⁻¹), or functional iron deficiency, as seen during treatment with erythropoiesis-stimulating agents (Table 1; Fig. 2) [25].

However, although ferritin values > 100 µg.l⁻¹ argue against concurrent true iron deficiency in the setting of inflammation, due to its acute phase reactivity, its diagnostic value is imperfect. Other tests, such as low reticulocyte Hb content (CHr < 28 pg) or an increase in hypochromic red cells (Hypo > 5%), can be utilised to evaluate for a component of iron deficiency, which if present suggests iron supplementation may be beneficial [25, 28]. The ratio of serum transferrin receptor level to the log of ferritin (sTfR/log Ft), the so-called ferritin index, may be used when these automated red cell parameters are not routinely available [25, 28]. A ferritin index > 2 indicates true iron deficiency, and is a strong predictor of response to i.v. iron [28, 29]. Detailed information regarding the advantages, disadvantages, limitations and diagnostic

values of these parameters can be found elsewhere [30].

Anaemia

Anaemia is defined by the World Health Organization as an Hb concentration < 130 g.l⁻¹ for men, < 120 g.l⁻¹ for non-pregnant women and < 110 g.l⁻¹ for pregnant women [31]. These definitions are widely quoted and accepted, and are used in most epidemiological studies. They are also adopted in most blood conservation guidelines, where no further laboratory work-up is deemed necessary for non-anaemic patients.

However, we consider that this may not be reliable for the classification of non-pregnant women undergoing surgical procedures in which moderate-to-high blood loss is expected. Women have lower circulating blood volumes than men, but the same procedures performed in either sex often result in comparable amounts of blood loss. Therefore, when measured as a proportion of circulating blood volume, blood losses are proportionally higher in women and may result in higher transfusion rates, as corroborated by a study in orthopaedic surgery [32] and the first Austrian benchmark study [33]. As women with a pre-operative Hb of 120 g.l⁻¹ are twice as likely to require a transfusion as men with an Hb of 130 g.l⁻¹, this Hb level should be considered as suboptimal in surgical settings. In

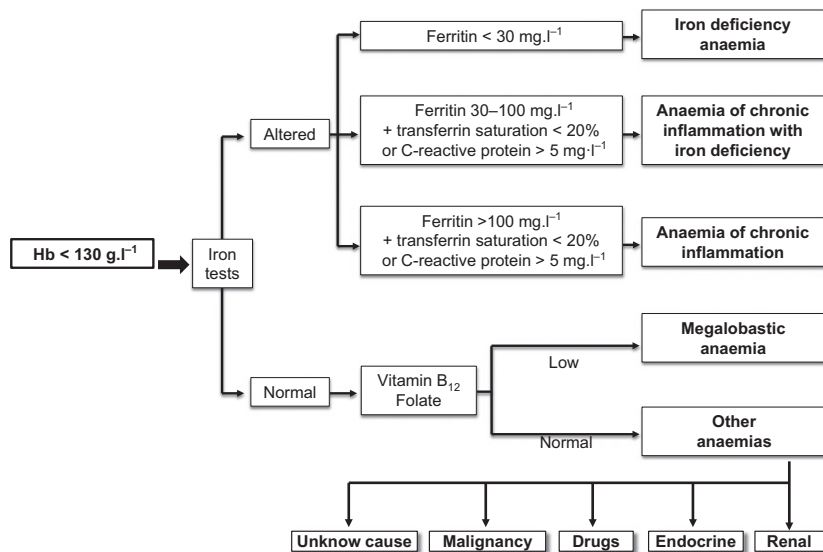


Figure 2 Algorithm for classification of peri-operative anaemia.

cardiac surgery, a 10 g.l⁻¹ decrease in Hb has been shown to be independently associated with increased transfusion requirements, increased mortality and prolonged hospital stay [34].

Consequently, apart from replenishing iron stores, pre-operative treatment of iron deficiency anaemia should be aimed at producing a target Hb level ≥ 130 g.l⁻¹ in both sexes, to minimise the risk of transfusion-associated unfavourable outcomes [35].

Patients who are anaemic and require major surgery (including the obstetrics and gynaecology population), especially if moderate to high blood loss (> 500 ml) is likely and/or if there is a ≥ 10% statistical probability for red blood cell (RBC) transfusion, should be investigated (Fig 3). Laboratory investigations should be performed as early as possible in the pre-operative period in order to implement the most appropriate treatment, if needed. They should initially include at least full blood count, serum ferritin, TSAT, a marker of inflammation (e.g. serum CRP) and a marker of renal function (e.g. serum creatinine), which can be easily requested by a surgeon (at the time of listing for surgery), an anaesthetist (during the pre-anaesthesia assessment) or a primary care physician/general practitioner. In some hospitals, there is also the possibility of automatically ordering additional laboratory tests when an Hb < 130 g.l⁻¹ is detected by pre-operative analysis.

Intervening to treat pre-operative iron deficiency anaemia in the surgical patient can be expected to reduce the number of transfusions given and thereby improve outcomes, even in those for whose anaemia, surgery is the cure. There is good evidence that attempts to correct iron deficiency anaemia pre-operatively will improve Hb before surgery, but there is little good quality evidence that it can modify the excess of risk for postoperative complications, other than those associated with transfusion [28, 36–42]. However, moderate to severe anaemia (Hb < 110 g.l⁻¹) results in worse outcomes than milder anaemia [43]. In contrast, there is also good evidence that correcting anaemia by transfusing blood can be detrimental to the outcomes of surgery [37, 44, 45].

Despite the lack of level-one evidence for improved outcomes, it is still recommended as good clinical practice to treat all surgical patients with pre-operative iron deficiency anaemia, but with a particular emphasis on treating those undergoing major surgery. Patients undergoing more minor surgical procedures are unlikely to see wide fluctuations in Hb postoperatively, and surgery can proceed while anaemia evaluation and treatment is ongoing.

Patients who have absolute iron deficiency should also be investigated for gastro-intestinal pathology (Table 1). The prevalence of other haematinic deficiencies before elective major orthopaedic surgery is

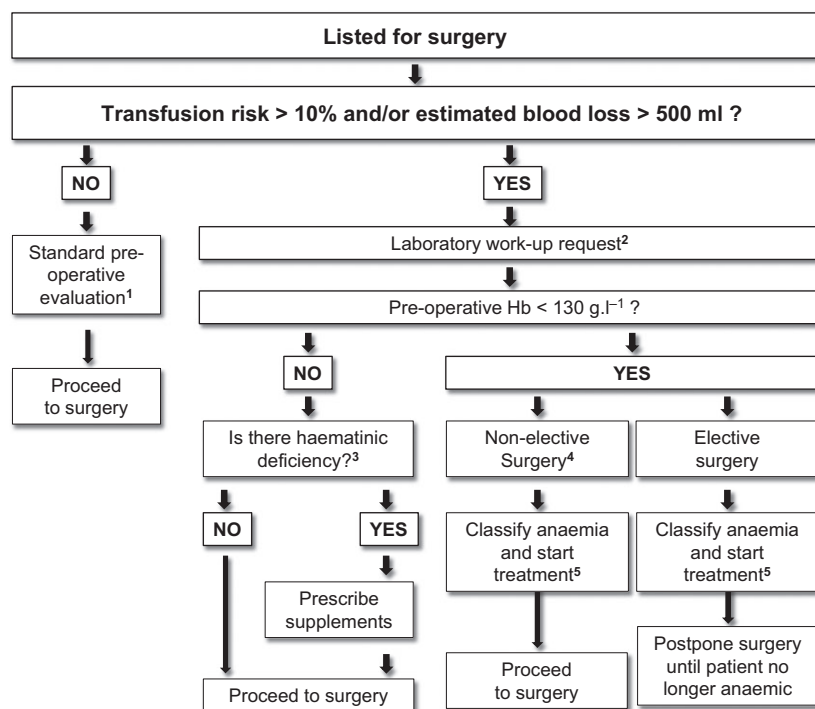


Figure 3 Algorithm for the management of a surgical patients. 1, According to centre protocol for each surgical procedure; 2, It should initially include at least full blood counts, serum ferritin, transferrin saturation, a marker of inflammation (e.g. serum C-reactive protein) and a marker of renal function (e.g. serum creatinine); 3, Haematinic deficiencies are defined by ferritin $< 100 \mu\text{g}\cdot\text{l}^{-1}$, vitamin $\text{B}_{12} < 270 \text{ pg}\cdot\text{ml}^{-1}$ and/or folate $< 3 \text{ pg}\cdot\text{ml}^{-1}$; 4, Including patients presenting late before surgery and non-deferrable surgery (e.g. cancer surgery); 5, According to algorithm depicted in Fig. 2.

approximately 12% for vitamin B_{12} [defined by serum concentration $< 270 \text{ pg}\cdot\text{ml}^{-1}$ ($200 \text{ pmol}\cdot\text{l}^{-1}$)], and 3% for folate [defined by serum concentration $< 3 \text{ ng}\cdot\text{ml}^{-1}$ ($5 \text{ nmol}\cdot\text{l}^{-1}$)] [46]. If none are found, other causes of anaemia should be ruled out (Fig. 2).

Treatment

Treatment of pre-operative iron deficiency anaemia should be implemented as early as possible before the scheduled surgical procedure. Most major surgery is elective. In the UK, the recent NHS Blood and Transplant audit showed there was a median of 60 days from listing the patient for surgery to the operation itself (<http://hospital.blood.co.uk/audits/national-comparative-audit/>). Thus, treatment of iron deficiency anaemia should be attempted while the patient is on the surgical waiting list, although this is currently rarely practised [47].

Regarding oral iron, in non-anaemic iron-deficient women, high doses of iron sulphate stimulate hepcidin

release, which results in lower iron absorption from the next daily dose [48]. In contrast, it appears that low-dose ($\leq 60 \text{ mg}$) oral iron given on alternate days may maximise fractional iron absorption, increase dosage efficacy, reduce gastro-intestinal exposure to unabsorbed iron and ultimately improve tolerance and adherence to treatment [48]. As anaemia-induced hypoxia and erythropoietin production downregulate the expression of hepcidin [15, 25], this may counterbalance the stimulatory effect of iron. In octogenarian patients with iron deficiency anaemia, 50 mg or 150 mg iron sulphate daily for 2 months both improved Hb and ferritin levels equally, but fewer side-effects were observed with the lower dose [49]. Therefore, when the interval before surgery is sufficient (at least 6–8 weeks) and no contra-indications are present, daily (40–60 mg) or alternate-day (80–100 mg) supplementation with oral iron and nutritional advice may be appropriate. The actual dose may depend upon the elemental iron content of the available oral iron

formulation, which differs depending on country of manufacture.

However, many patients will not respond to oral iron, especially those with functional iron deficiency and chronic illness or infection and those with ongoing blood loss [15, 25]. Others will not tolerate oral iron due to gastro-intestinal side-effects. Once oral iron has been commenced, the Hb should be measured again, at least 4 weeks before surgery. In the absence of an increased Hb or if the patient is intolerant, i.v. iron is the preferred replacement route. If surgery is planned in less than 6 weeks time, i.v. iron may also be the most effective option.

Intravenous iron is highly efficacious at replenishing iron stores and increasing Hb in anaemia due to iron deficiency with or without inflammation (Fig. 2). The dose of i.v. iron may be calculated from the baseline and target Hb and patient's body weight, adding 500 mg for iron stores (in patients > 35 kg) [25]. In practice, a dose of 1000–1500 mg is sufficient in most surgical patients and can usually be given by slow infusion over less than 1 h in one sitting or in two divided doses, depending on the preparation used. Most patients feel better in 3 days, with a rapid Hb response (50% at 5 days, 75% at 10–14 days, maximal at 3 weeks) [50].

In a small series, i.v. iron sucrose before orthopaedic surgery has been shown to be useful for treating iron deficiency anaemia, with a maximum effect on Hb after 2 weeks [51]. In abdominal surgery, i.v. iron carboxymaltose (1000 mg) 2–4 weeks [40, 42] or 8–10 days pre-operatively [41] has been shown to decrease RBC transfusion and hospital stay.

Intravenous iron treatment < 2 weeks pre-operatively has also been shown to be successful in decreasing the need for RBC transfusion, acute kidney injury, hospital stay and infections in orthopaedic and cardiac surgery [52, 53]. A quicker recovery in Hb postoperatively was also observed in iron-deficient patients treated with i.v. iron sucrose [54] or carboxymaltose [41, 42]. Administration of i.v. iron isomaltoside (1000 mg) 1 day before or even on the day of surgery can improve postoperative Hb recovery even in non-anaemic patients without iron deficiency [55].

Presently, there is a lack of high-quality data about effect on outcomes, but a number of large randomised, controlled trials are in progress [56].

Iron deficiency without anaemia

A recent meta-analysis of 21 trials in different clinical settings concluded that there is emerging evidence that non-anaemic iron deficiency is a disease in its own right, deserving of further research in the development of strategies for detection and treatment [57]. Non-anaemic patients with iron deficiency or inadequate iron stores undergoing surgical procedures with expected moderate-to-high blood loss may benefit from pre-operative iron administration, to boost recovery from postoperative anaemia [1, 3] (Fig. 2). In those with absolute iron deficiency (ferritin < 30 $\mu\text{g.l}^{-1}$), the need for gastro-intestinal investigations should be considered [3]. Microcytosis due to iron deficiency without anaemia should be evaluated for chronic hypoxaemia, myeloproliferative disease (e.g. polycythaemia rubra vera), or another cause of increased red cell production. Iron replacement may cause a surge in Hb that could induce hyperviscosity in this subgroup.

In non-anaemic orthopaedic surgical patients, the administration of iron ferrous sulphate and vitamins during the weeks preceding surgery has been shown to increase serum ferritin and transferrin saturation [58] and reduce transfusion [59]. However, adherence to iron supplementation was 67%, and adverse drug reactions were present in 52% of patients [60].

In women undergoing major non-cardiac surgical procedures, the prevalence of true iron deficiency (ferritin < 30 $\mu\text{g.l}^{-1}$) for Hb 120–129 g.l^{-1} was similar to that for Hb < 120 g.l^{-1} , and significantly higher than for Hb \geq 130 g.l^{-1} (42%, 51% and 24%, respectively; $p < 0.05$). This also applies to insufficient iron supply for erythropoiesis, as defined by TSAT < 20% (58%, 69% and 34%, respectively; $p < 0.05$) [22]. Thus, although considered as non-anaemic according to WHO definitions [31], most women with Hb 120–129 g.l^{-1} would benefit from iron replacement therapy, for optimising pre-operative Hb levels and facilitating recovery from postoperative anaemia.

For non-anaemic patients undergoing surgery with a high risk for developing severe postoperative anaemia, we suggest the administration of oral iron. Intravenous iron administration (e.g. 500 mg) should be considered if surgery is scheduled to take place in < 4 weeks or if the patient cannot tolerate oral iron [1].

Delaying surgery

As stated above, uncorrected pre-operative anaemia is a risk factor for poor outcome [61–66]. Although there is little evidence currently that treatment of iron deficiency anaemia improves outcome [60, 67], it does reduce transfusion and may reduce hospital stay. Consequently, there is considerable uncertainty as to whether surgery should be delayed in patients with undiagnosed/untreated anaemia.

The problem may be divided into patients undergoing surgery for benign vs. malignant disease. In major surgery for benign disease, especially when blood loss > 500 ml is expected or possible, patients should be informed about the relationship between anaemia, morbidity and mortality, and should be given the opportunity to postpone non-urgent surgery until their anaemia is investigated and treated [4, 35]. Several guidelines recommend postponing the operation for about 4 weeks to allow appropriate management of anaemia [1, 2, 4, 8].

In some European countries, the allowable time between diagnosis and surgery for malignant disease is subjected to legal regulations. Nevertheless, many receive pre-operative adjuvant therapy, which allows sufficient time for concurrent treatment of pre-operative iron deficiency anaemia.

Postoperative anaemia

Anaemia is present in up to 90% of patients in the immediate postoperative period after major surgery [68]. The main causes are: presence of pre-operative anaemia; peri-operative blood loss; poor nutritional intake in the postoperative period; and frequent blood sampling for laboratory tests. In addition, increased hepcidin due to the inflammatory response to surgery can lead to inhibition of iron absorption from the small bowel and reduced iron release from stores (iron sequestration) [25] (Fig. 1). These effects can last for a few weeks after major surgery and aggravate postoperative iron deficiency anaemia.

Recently, the National Institute for Clinical Excellence [69] and other international guidelines [4, 5, 70] have reduced the transfusion thresholds for surgical patients to 70 g.l⁻¹ (80 g.l⁻¹ for patients with a history of ischaemic heart disease) in the absence of active

bleeding. This change in practice has led to more patients being discharged with significant anaemia after major surgery. Therefore, it is important to implement strategies for postoperative anaemia management.

Management of iron deficiency with oral iron in the immediate postoperative period has a very limited role due to poor absorption and considerable side-effects, and is not recommended [4]. In contrast, i.v. iron has been successfully used to treat iron deficiency anaemia after surgery for lower limb arthroplasty [71], gastrectomy [72] and postpartum haemorrhage [73]. The administration of a single 1000-mg dose of ferric carboxymaltose after major orthopaedic surgery, abdominal and genito-urinary surgery, and others, has been shown to correct anaemia and improve quality of life compared with oral iron [74, 75].

Transfused packed RBC contains haem iron (200–250 mg per unit), plus a small amount of labile iron which is immediately available for erythropoiesis; in part, dependent on the storage time of the transfused RBC unit [76]. After transfusion, the patient's Hb is usually < 10 g.l⁻¹. Correcting postoperative iron deficiency, in addition to RBC transfusion to raise Hb, may be important to improve function, as shown in animal models [77]. Further research is required as to whether supplemental iron given to correct iron deficiency in patients who are transfused peri-operatively is beneficial in humans.

Safety of i.v. iron therapy

Currently, six i.v. iron formulations are available in the USA and/or Europe (Table 3), and serious adverse events (SAEs) are very rare (38 per 10⁶ administrations; deaths, 0.4 per 10⁶ administrations) [78]. In contrast, the most recent Serious Hazards of Transfusion Report in the UK quotes the risk of death related to transfusion as 1 in 100,000, and the risk of major morbidity as 1 in 16,000 [79]. The European Medicines Agency concluded that the benefits of i.v. iron exceed the risks when used appropriately (correct indication and dose), without any difference in safety profile among available formulations [80]. In line with these recommendations, guidance for risk minimisation and management of hypersensitivity reactions to i.v. iron has been recently published [81].

Table 3 Characteristics of different i.v. iron formulations.

	Iron gluconate**	Iron sucrose††	Low molecular weight iron dextran (LMWID)‡‡	Ferric carboxymaltose§§	Iron isomaltoside 1000¶¶	Ferumoxytol***
Brand name	Ferrlecit®	Venofer®	Cosmofer® INFeD®	Ferinject® Injectafer®	Monofer® Monoferro®	FeraHeme® Rienso®
Carbohydrate shell	Gluconate (monosaccharide)	Sucrose (disaccharide)	Dextran (branched polysaccharide)	Carboxymaltose (branched polysaccharide)	Isomaltoside (linear oligosaccharide)	Polyglucose sorbitol carboxy-methylether
Complex type*	Type II	Type II	Type I	Type I	Type I	Type I
Molecular weight; kD	289–440	30–60	165	150	150	750
Initial distribution volume; l	6	3.4	3.5	3.5	3.4	3.16
Plasma half-life; h	1	6	20	16	20	15
Labile iron (% injected dose)†	3.3	3.5	2.0	0.6	1.0	0.8
Iron content; mg.ml ⁻¹	12.5	20	50	50	100	30
Maximal single dose; mg	125	200	20 mg.kg ⁻¹	20 mg.kg ⁻¹ (max 1000 mg)	20 mg.kg ⁻¹	510
Infusion time for 1000 mg; min.‡	720	300	90–150§	≥ 15	≥ 15	≥ 15
Product cost per 1000 mg; €¶	–	128	100	227	212	162

*Type I, robust and strong; Type II, semirobust and moderately strong; Type III, labile and weak (Crichton RR, Danielson BG, Geisser P. Iron therapy with special emphasis on i.v. administration, 4th Ed. UNI-MED Verlag AG, Bremen, 2008).
 †Jahn MR, et al. Eur J Pharm Biopharm 2011; 78:480–91.
 ‡Includes 30-min postinfusion observation.
 §Safe administration of 1000 mg LMWID in 1 h has been described (Auerbach M et al. Am J Hematol 2011; 86:860–2).
 ¶British National Formulary BNF68, September 2014 – March 2015. BMJ Group and the Royal Pharmaceutical Society of Great Britain, London, 2014.
 **Ferrlecit summary of product characteristics. <http://www.products.sanofi-aventis.us/ferrlecit/ferrlecit.pdf> (accessed 01/08/2016).
 ††Venofer summary of product characteristics. <http://www.luitpold.com/documents/22.pdf> (accessed 01/08/2016).
 ‡‡Cosmofer summary of product characteristics. <http://www.cosmofer.com/product/cosmofer-spc/cosmofer-spc.aspx> (accessed 01/08/2016).
 §§Ferinject summary of product characteristics. <http://www.ferinject.co.uk/smpc/> (accessed 01/08/2016).
 ¶¶Monofer summary of product characteristics. <http://www.monofer.com/spc.aspx> (accessed 01/08/2016).
 ***FeraHeme summary of product characteristics. http://www.feraheme.com/pdfs/Feraheme_Prescribing_Information.pdf (accessed 01/08/2016).

Some patients who receive i.v. iron may exhibit complement activation-related pseudo-allergy, which should not be misinterpreted as hypersensitivity. This occurs in approximately 1:200 iron-treated patients, and can consist of arthralgia, myalgia or flushing, but without associated hypotension, tachycardia, tachypnoea, wheezing, stridor or peri-orbital oedema [82]. Symptoms abate without intervention, and the patient may be rechallenged or another i.v. iron formulation tried [82].

A recent meta-analysis of 103 trials published from 1965 to 2013 (including 19,253 patients) concluded that i.v. iron therapy was not associated with an increased risk of serious adverse events or infection when compared with oral or intramuscular iron, no iron or placebo [83]. In large observational studies, peri-operative i.v. iron did not negatively impact on rates of transfusion, infection and 30-day mortality in surgical patients [51, 84]. In contrast, red cell transfusion delivers haem and labile iron which supports bacterial growth more readily [85].

The preponderance of published evidence indicates that i.v. iron is safe, supporting a greater and earlier role for treating iron deficiency and raising the question of whether parenteral iron should be considered for front-line therapy in conditions where oral iron will predictably fail. However, further prospective efficacy and safety trials in various surgical settings are required to make evidenced-based conclusions [82].

Cost implications of iron therapy

There are few studies that have undertaken formal cost–benefit analysis of iron treatment. Bhandari et al. [86] performed a comparative analysis of the costs of administering i.v. iron formulations against RBC transfusion, by considering acquisition costs plus administration costs (nursing time and giving sets) and transportation to hospital, across three dosage levels (600 mg, 1000 mg and 1600 mg). At all doses, i.v. iron resulted in a net saving when compared with RBC transfusion, with the biggest differences observed for ferric carboxymaltose and iron isomaltoside-1000.

Calvet et al. [87] compared the cost implications of ferric carboxymaltose vs. iron sucrose vs. oral iron for avoiding red cell transfusion in 282 patients with

colorectal cancer and anaemia, using cost-minimisation analysis. Direct and indirect costs for acquisition and administration of iron product and RBC concentrates, as well as hospitalisation costs, were included in the cost model. Ferric carboxymaltose reduced hospital stay by 2.3 days compared with iron sucrose and by 2.6 days compared with oral iron, resulting in cost savings of £437 (485€, \$532) and £245 (274€, \$300) per patient, respectively.

Data from 182 matched pairs of total lower limb arthroplasty patients, managed with a restrictive transfusion protocol and without (control) or with postoperative i.v. iron, were retrospectively reviewed [88]. Acquisition and administration costs of iron (600 mg iron sucrose or ferric carboxymaltose) and RBC concentrates, Hb, and prolonged stay in hospital were used. Patients receiving RBC transfusion stayed in hospital longer (1.9 days [95% CI 1.2–2.6]). Compared with control, i.v. iron reduced RBC transfusion rate (11.5% vs. 26.4%) without incremental cost [88].

Thus, the use of newer i.v. iron formulations (e.g. iron isomaltoside 1000 or ferric carboxymaltose) allows the rapid administration of larger single doses (≥ 1000 mg), and are more convenient both for the patient (e.g. fewer hospital visits, less time off work) and the health system (e.g. less visits to day hospital, less ambulance transfers, less total cost) [40, 86–89].

When comparing i.v. iron with other interventions implemented to reduce blood transfusion and/or placebo, it would be interesting to include the costs of postoperative complications and other clinical endpoints as well. Anaesthetists and surgeons should consider pre-operative iron deficiency not as a single ‘point’, although its treatment is essential. Instead, this should trigger a peri-operative bundle of care, known as Patient Blood Management, started at the time that surgery is booked and carried on until full recovery from the surgical intervention has occurred. This should be planned and documented pre-operatively, both in urgent and elective patients [90–92]. We believe there is ample evidence to support peri-operative blood management within the context of an evidence-based enhanced recovery programme to secure optimal interpretation of the intervention [93].

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References

- Beris P, Muñoz M, García-Erce JA, Thomas D, Maniatis A, Van der Linden P. Perioperative anaemia management: consensus statement on the role of intravenous iron. *British Journal of Anaesthesia* 2008; **100**: 599–604.
- Goodnough LT, Maniatis A, Earnshaw P, et al. Detection, evaluation, and management of preoperative anaemia in the elective orthopaedic surgical patient: NATA guidelines. *British Journal of Anaesthesia* 2011; **106**: 13–22.
- National Blood Authority. Patient Blood Management Guidelines: Module 2 – Peri-operative. <https://www.blood.gov.au/pbm-module-2> (accessed 17/08/2016).
- Leal-Noval SR, Muñoz M, Asuero M, et al. Spanish Consensus Statement on alternatives to allogeneic blood transfusion: the 2013 update of the "Seville Document". *Blood Transfusion* 2013; **11**: 585–610.
- Kozek-Langenecker SA, Afshari A, Albaladejo P, et al. Management of severe perioperative bleeding: guidelines from the European Society of Anaesthesiology. *European Journal of Anaesthesiology* 2013; **30**: 270–382.
- Goodnough LT, Shander A, Spivak JL, et al. Detection, evaluation, and management of anemia in the elective surgical patient. *Anesthesia and Analgesia* 2005; **101**: 1858–6.

7. SABM. Anemia prevention and management program implementation guide, 2015. https://www.sabm.org/sites/default/files/anemia_prevention_management_program_implementation_guide.pdf (accessed 01/08/2016).
8. Kotzé A, Harris A, Baker C, et al. British Committee for Standards in Haematology guidelines on the identification and management of pre-operative anaemia. *British Journal of Haematology* 2015; **171**: 322–31.
9. Practice guidelines for perioperative blood management: an updated report by the American Society of Anesthesiologists Task Force on Perioperative Blood Management. *Anesthesiology* 2015; **122**: 241–75.
10. Vaglio S, Prisco D, Biancofiore G, et al. Recommendations for the implementation of a Patient Blood Management programme. Application to elective major orthopaedic surgery in adults. *Blood Transfusion* 2016; **14**: 23–65.
11. Klein AA, Arnold P, Bingham RM, et al. AAGBI guidelines: the use of blood components and their alternatives 2016. *Anaesthesia* 2016; **71**: 829–42.
12. WHO. *Iron Deficiency Anaemia: Assessment, Prevention, and Control. A Guide for Programme Managers*. Geneva: WHO, 2001.
13. Pasricha SR, Drakesmith H, Black J, Hipgrave D, Biggs BA. Control of iron deficiency anemia in low- and middle-income countries. *Blood* 2013; **121**: 2607–17.
14. Peyrin-Biroulet L, Williet N, Cacoub P. Guidelines on the diagnosis and treatment of iron deficiency across indications: a systematic review. *American Journal of Clinical Nutrition* 2015; **102**: 1585–94.
15. Camaschella C. Iron-deficiency anemia. *New England Journal of Medicine* 2015; **372**: 1832–43.
16. Wong CC, Ng AC, Kriharides L, Sindone AP. Iron deficiency in heart failure: looking beyond anaemia. *Heart Lung Circulation* 2016; **25**: 209–16.
17. Patteril MV, Davey-Quinn AP, Gedney JA, Murdoch SD, Bellamy MC. Functional iron deficiency, infection and systemic inflammatory response syndrome in critical illness. *Anaesthesia and Intensive Care* 2001; **29**: 473–8.
18. Enjuanes C, Bruguera J, Grau M, et al. Iron status in chronic heart failure: impact on symptoms, functional class and sub-maximal exercise capacity. *Revista Española de Cardiología (Engl Ed)* 2016; **69**: 247–55.
19. Cleland JG, Zhang J, Pellicori P, et al. Prevalence and outcomes of anemia and hematinic deficiencies in patients with chronic heart failure. *JAMA Cardiology* 2016; **1**: 539–47.
20. Anker SD, Comin Colet J, Filippatos G, et al. Ferric carboxymaltose in patients with heart failure and iron deficiency. *New England Journal of Medicine* 2009; **361**: 2436–48.
21. Ponikowski P, van Veldhuisen DJ, Comin-Colet J, et al. CONFIRM-HF Investigators. Beneficial effects of long-term intravenous iron therapy with ferric carboxymaltose in patients with symptomatic heart failure and iron deficiency. *European Heart Journal* 2015; **36**: 657–68.
22. García-Erce JA, Laso-Morales MJ, Gómez-Ramírez S, Núñez-Matas MJ, Muñoz M. Analysis of the prevalence and causes of low preoperative haemoglobin levels in a large multicentre cohort of patients undergoing major non-cardiac surgery. *Transfusion Medicine* 2016; **26**(Suppl. 1): 48.
23. Harju E. Empty iron stores as a significant risk factor in abdominal surgery. *Journal of Parenteral and Enteral Nutrition* 1988; **12**: 282–5.
24. Izuel-Ramí M, García-Erce JA, Gómez-Barrera M, et al. Relación entre la transfusión de sangre alogénica, la deficiencia de hierro y la infección nosocomial en pacientes con fractura de cadera. *Medicina Clínica (Barcelona)* 2008; **131**: 647–52.
25. Muñoz M, García-Erce JA, Remacha AF. Disorders of iron metabolism. Part II: iron deficiency and iron overload. *Journal of Clinical Pathology* 2011; **64**: 287–96.
26. Walters GO, Miller FM, Worwood M. Serum ferritin concentration and iron stores in normal subjects. *Journal of Clinical Pathology* 1973; **26**: 770–2.
27. Mast AE, Blinder MA, Gronowski AM, Chumley C, Scott MG. Clinical utility of the soluble transferrin receptor and comparison with serum ferritin in several populations. *Clinical Chemistry* 1998; **44**: 45–51.
28. Basora M, Colomina MJ, Tio M, Mora L, Salazar F, Ciercoles E. Optimizing preoperative haemoglobin with intravenous iron. *British Journal of Anaesthesia* 2013; **110**: 488–90.
29. Thomas DW, Hinchliffe RF, Briggs C, Macdougall IC, Littlewood T, Cavill I; British Committee for Standards in Haematology. Guideline for the laboratory diagnosis of functional iron deficiency. *British Journal of Haematology* 2013; **161**: 639–48.
30. Muñoz M, Gómez-Ramírez S, Kozek-Langenecker S. Pre-operative haematological assessment in patients scheduled for major surgery. *Anaesthesia* 2016; **71**(Suppl. 1): s19–28.
31. World Health Organization. Haemoglobin concentrations for the diagnosis of anaemia and assessment of severity. WHO/NMH/NHD/MNM/11.1. <http://www.who.int/vmnis/indicators/haemoglobin.pdf> (accessed 28/04/2016).
32. Gombotz H, Rehak PH, Shander A, Hofmann A. Blood use in elective surgery: the Austrian benchmark study. *Transfusion* 2007; **47**: 1468–80.
33. Rosencher N, Kerckamp HE, Macheras G, et al. Orthopedic surgery transfusion hemoglobin European overview (OSTHEO) study: blood management in elective knee and hip arthroplasty in Europe. *Transfusion* 2003; **43**: 459–69.
34. Klein AA, Collier TJ, Brar MS, et al. The incidence and importance of anaemia in patients undergoing cardiac surgery in the UK – the first Association of Cardiothoracic Anaesthetists national audit. *Anaesthesia* 2016; **71**: 627–35.
35. Muñoz M, Gomez-Ramirez S, Kozek-Langenecker S, et al. 'Fit to fly': overcoming barriers to preoperative haemoglobin optimization in surgical patients. *British Journal of Anaesthesia* 2015; **115**: 15–24.
36. Keeler B, Simpson J, Ng S, et al. The feasibility and clinical efficacy of intravenous iron administration for preoperative anaemia in patients with colorectal cancer. *Colorectal Diseases* 2014; **16**: 794–800.
37. Hogan M, Klein AA, Richards T. The impact of anaemia and intravenous iron replacement therapy on outcomes in cardiac surgery. *European Journal of Cardiothoracic Surgery* 2015; **4**: 218–26.
38. Okuyama M, Ikeda K, Shibata T, Tsukahara Y, Kitada M, Shimano T. Preoperative iron supplementation and intraoperative transfusion during colorectal cancer surgery. *Surgery Today* 2005; **35**: 36–40.
39. Bisbe E, García-Erce JA, Díez-Lobo AI, Muñoz M; Anaemia Working Group España. A multicentre comparative study on the efficacy of intravenous ferric carboxymaltose and iron sucrose for correcting preoperative anaemia in patients undergoing major elective surgery. *British Journal of Anaesthesia* 2011; **107**: 477–8.
40. Laso-Morales MJ, Gómez-Ramírez S, Pontes-García C, Díaz-espallardo C, Muñoz M. Intravenous versus oral iron for treating iron deficiency anaemia in colorectal cancer patients: a single-centre, observational cohort study. *Transfusion Medicine* 2016; **26**(Suppl. 1): 53.

41. Froessler B, Palm P, Weber I, Hodyl NA, Singh R, Murphy EM. The important role for intravenous iron in perioperative patient blood management in major abdominal surgery: a randomized controlled trial. *Annals of Surgery* 2016; **264**: 41–6.
42. Calleja JL, Delgado S, del Val A, et al. Colon Cancer Study Group. Ferric carboxymaltose reduces transfusions and hospital stay in patients with colon cancer and anemia. *International Journal of Colorectal Diseases* 2016; **31**: 543–51.
43. Acheson AG, Brookes MJ, Spahn DR. Effects of allogeneic red blood cell transfusions on clinical outcomes in patients undergoing colorectal cancer surgery - systematic review and meta-analysis. *Annals of Surgery* 2012; **256**: 235–44.
44. Carson JL, Duff A, Poses RM, et al. Effect of anaemia and cardiovascular disease on surgical mortality and morbidity. *Lancet* 1996; **348**: 1055–60.
45. Ferraris VA, Davenport DL, Saha SP, Austin PC, Zwischenberger JB. Surgical outcomes and transfusion of minimal amounts of blood in the operating room. *Archives of Surgery* 2012; **147**: 49–55.
46. Bisbe E, Castillo J, Sáez M, et al. Prevalence of preoperative anemia and hematinic deficiencies in patients scheduled for major orthopedic surgery. *Transfusion Alternatives in Transfusion Medicine* 2008; **4**: 166–73.
47. Rogers BA, Cowie A, Alcock C, Rosson JW. Identification and treatment of anaemia in patients awaiting hip replacement. *Annals of the Royal College of Surgeons of England* 2008; **90**: 504–7.
48. Moretti D, Goede JS, Zeder C, et al. Oral iron supplements increase hepcidin and decrease iron absorption from daily or twice-daily doses in iron-depleted young women. *Blood* 2015; **126**: 1981–9.
49. Rimon E, Kagansky N, Kagansky M, et al. Are we giving too much iron? Low-dose iron therapy is effective in octogenarians. *American Journal of Medicine* 2005; **118**: 1142–7.
50. Goodnough LT, Skikne B, Brugnara C. Erythropoietin, iron, and erythropoiesis. *Blood* 2000; **96**: 823–33.
51. Theussinger OM, Leyvraz PF, Schanz U, et al. Treatment of iron deficiency anemia in orthopedic surgery with intravenous iron: efficacy and limits: a prospective study. *Anesthesiology* 2007; **107**: 923–27.
52. Yoo YC, Shim JK, Kim JC, Jo YY, Lee JH, Kwak YL. Effect of single recombinant human erythropoietin injection on transfusion requirements in preoperatively anemic patients undergoing valvular heart surgery. *Anesthesiology* 2011; **115**: 929–37.
53. Muñoz M, Gómez-Ramírez S, Cuenca J, et al. Very-short-term perioperative intravenous iron administration and postoperative outcome in major orthopedic surgery: a pooled analysis of observational data from 2547 patients. *Transfusion* 2014; **54**: 289–99.
54. García-Erce JA, Cuenca J, Martínez F, Cardona R, Pérez-Serrano L, Muñoz M. Perioperative intravenous iron preserves iron stores and may hasten the recovery from post-operative anaemia after knee replacement surgery. *Transfusion Medicine* 2006; **16**: 335–41.
55. Johansson PI, Rasmussen AS, Thomsen LL. Intravenous iron isomaltoside 1000 (Monofer®) reduces postoperative anaemia in preoperatively non-anaemic patients undergoing elective or subacute coronary artery bypass graft, valve replacement or a combination thereof: a randomized double-blind placebo-controlled clinical trial (the PROTECT trial). *Vox Sanguinis* 2015; **109**: 257–66.
56. Richards T, Clevenger B, Keidan J, et al. PREVENTT: preoperative intravenous iron to treat anaemia in major surgery: study protocol for a randomised controlled trial. *Trials* 2015; **16**: 254.
57. Pratt JJ, Khan KS. Non-anaemic iron deficiency – a disease looking for recognition of diagnosis: a systematic review. *European Journal of Haematology* 2016; **96**: 618–28.
58. Cuenca J, García-Erce JA, Martínez F, Cardona R, Pérez-Serrano L, Muñoz M. Preoperative haematinics and transfusion protocol reduce the need for transfusion after total knee replacement. *International Journal of Surgery* 2007; **5**: 89–94.
59. Lachance K, Savoie M, Bernard M, et al. Oral ferrous sulfate does not increase preoperative hemoglobin in patients scheduled for hip or knee arthroplasty. *Annals of Pharmacotherapy* 2011; **45**: 764–70.
60. Kotze A, Carter LA, Scally AJ. Effect of a patient blood management programme on preoperative anaemia, transfusion rate, and outcome after primary hip or knee arthroplasty: a quality improvement cycle. *British Journal of Anaesthesia* 2012; **108**: 943–52.
61. Beattie WS, Karkouti K, Wijeyesundera DN, Tait G. Risk associated with preoperative anemia in noncardiac surgery: a single-center cohort study. *Anesthesiology* 2009; **110**: 574–81.
62. Musallam KM, Tamim HM, Richards T, et al. Preoperative anaemia and postoperative outcomes in noncardiac surgery: a retrospective cohort study. *Lancet* 2011; **378**: 1396–407.
63. Leichtle SW, Mouawad NJ, Lampman R, Singal B, Cleary RK. Does preoperative anemia adversely affect colon and rectal surgery outcomes? *Journal of the American College of Surgeons* 2011; **212**: 187–94.
64. Ranucci M, Di Dedda U, Castelveccchio S, et al. Impact of preoperative anemia on outcome in adult cardiac surgery: a propensity-matched analysis. *Annals of Thoracic Surgery* 2012; **94**: 1134–41.
65. Jans Ø, Jørgensen C, Kehlet H, Johansson PI; Lundbeck Foundation Centre for Fast-track Hip and Knee Replacement Collaborative Group. Role of preoperative anemia for risk of transfusion and postoperative morbidity in fast-track hip and knee arthroplasty. *Transfusion* 2014; **54**: 717–26.
66. Fowler AJ, Ahmad T, Phull MK, et al. Meta-analysis of the association between preoperative anaemia and mortality after surgery. *British Journal of Surgery* 2015; **102**: 1314–24.
67. Elhenawy AM, Meyer SR, Bagshaw SM, MacArthur RG, Carroll LJ. Role of preoperative intravenous iron therapy to correct anemia before major surgery: study protocol for systematic review and meta-analysis. *Systematic Reviews* 2015; **4**: 29.
68. Shander A, Knight K, Thurer R, Adamson J, Spence R. Prevalence and outcomes of anaemia in surgery: a systematic review of the literature. *American Journal of Medicine* 2004; **116**(Suppl. 7A): 58–69.
69. National Institute for Clinical Excellence guidance. Blood Transfusion. <https://www.nice.org.uk/guidance/ng24/resources/blood-transfusion-1837331897029> (accessed 01/08/2016).
70. Carson JL, Grossman BJ, Kleinman S, et al. Red blood cell transfusion: a clinical practice guideline from the AABB. *Annals of Internal Medicine* 2012; **157**: 49–58.
71. Muñoz M, Naviera E, Seara J, Cordero J. Effects of postoperative intravenous iron on transfusion requirements after lower limb arthroplasty. *British Journal of Anaesthesia* 2012; **108**: 532–4.
72. Yoon HM, Kim Y-W, Nam BH, et al. Intravenous iron supplementation may be superior to observation in acute isovolemic anemia after gastrectomy for cancer. *World Journal of Gastroenterology* 2014; **20**: 1852–57.

73. Seid MH, Derman RJ, Baker JB, Banach W, Goldberg C, Rogers R. Ferric carboxymaltose injection in the treatment of postpartum iron deficiency anemia: a randomized controlled clinical trial. *American Journal of Obstetrics and Gynecology* 2008; **199**: e1–435.
74. Bisbe E, Molto L, Arroyo R, Muniesa JM, Tejero M. Randomised trial comparing ferric carboxymaltose vs oral ferrous glycine sulphate for postoperative anaemia after total knee arthroplasty. *British Journal of Anaesthesia* 2014; **113**: 402–9.
75. Khalafallah AA, Yan C, Al-Badri R, et al. Intravenous ferric carboxymaltose versus standard care in the management of postoperative anaemia: a prospective, open-label, randomised controlled trial. *Lancet Haematology* 2016; **3**: e415–25.
76. Hod EA, Brittenham GM, Billote GB, et al. Transfusion of human volunteers with older, stored red blood cells produces extravascular hemolysis and circulating non-transferrin-bound iron. *Blood* 2011; **118**: 6675–82.
77. Finch CA, Miller LR, Inamdar AR, Person R, Seiler K, Mackler B. Iron deficiency in the rat. Physiological and biochemical studies of muscle dysfunction. *Journal of Clinical Investigation* 1976; **58**: 447–53.
78. Chertow GM, Mason PD, Vaage-Nilsen O, Ahlmén J. Update on adverse drug events associated with parenteral iron. *Nephrology Dialysis and Transplantation Journal* 2006; **21**: 378–82.
79. Serious Hazards of Transfusion Report, 2015. <http://www.shotuk.org/report-summary-supplement-2015/> (accessed 01/08/2016).
80. European Medicines Agency. New recommendations to manage risk of allergic reactions with intravenous iron-containing medicines. http://www.ema.europa.eu/ema/index.jsp?curl=pages/news_and_events/news/2013/06/newsdetail_001833.jsp&mid=WC0b01ac058004d5c1 (accessed 01/08/2016).
81. Rampton D, Folkersen J, Fishbane S, et al. Hypersensitivity reactions to intravenous iron: guidance for risk minimization and management. *Haematologica* 2014; **99**: 1671–6.
82. Auerbach M, Adamson JW. How we diagnose and treat iron deficiency anemia. *American Journal of Hematology* 2016; **91**: 31–8.
83. Avni T, Bieber A, Grossman A, Green H, Leibovici L, Gafter-Gvili A. The safety of intravenous iron preparations: systematic review and meta-analysis. *Mayo Clinic Proceedings* 2015; **90**: 12–23.
84. Torres S, Kuo YH, Morris K, Neibart R, Holtz JB, Davis JM. Intravenous iron following cardiac surgery does not increase the infection rate. *Surgical Infections (Larchmt)* 2006; **7**: 361–6.
85. Andrews SC, Robinson AK, Rodríguez-Quiñones F. Bacterial iron homeostasis. *FEMS Microbiology Reviews* 2003; **27**: 215–37.
86. Bhandari S. Update of a comparative analysis of cost minimization following the introduction of newly available intravenous iron therapies in hospital practice. *Therapeutics and Clinical Risk Management* 2011; **7**: 501–9.
87. Calvet X, Gené E, Ruiz MA, et al. Cost-minimization analysis favours intravenous ferric carboxymaltose over ferric sucrose or oral iron as preoperative treatment in patients with colon cancer and iron deficiency anaemia. *Technology and Health Care* 2016; **24**: 111–20.
88. Muñoz M, Gómez-Ramírez S, Martín-Montañez E, Naveira E, Seara J, Pavia J. Cost of post-operative intravenous iron therapy in total lower limb arthroplasty: a retrospective, matched cohort study. *Blood Transfusion* 2014; **12**: 40–9.
89. Reinisch W, Staun M, Bhandari S, Muñoz M. State of the iron: how to diagnose and efficiently treat iron deficiency anemia in inflammatory bowel disease. *Journal of Crohn's and Colitis* 2013; **7**: 429–40.
90. Clevenger B, Richards T. Pre-operative anaemia. *Anaesthesia* 2015; **70**(Suppl. 1): 20–8, e6–8.
91. Clevenger B, Mallett SV, Klein AA, Richards T. Patient blood management to reduce surgical risk. *British Journal of Surgery* 2015; **102**: 1325–37.
92. Meybohm P, Richards T, Isbister J, et al. Patient blood management bundles to facilitate implementation. *Transfusion Medicine Reviews* 2017; **31**: 62–71.
93. Kehlet H, Jørgensen CC. Advancing surgical outcomes research and quality improvement within an enhanced recovery program framework. *Annals of Surgery* 2016; **264**: 237–8.