## Summary of Guidelines for the Use of Platelet Transfusions

# A British Society for Haematology Guideline (2016) - Appendix 1

To read the full guideline please go to <u>https://b-s-h.org.uk/guidelines/guidelines/use-of-platelet-transfusions/</u>

### Platelet transfusion: principles, risks, alternatives and best practice

Platelet transfusions are an essential component in the management of selected patients with thrombocytopenia. However they need to be used judiciously as they are a limited resource and are not risk free.

| Classification of conditions which<br>may require platelet transfusion  | Platelet transfusion: Indication categories<br>and contraindications  |
|---|---|
| Bone marrow failure (BMF). Reversible associated with<br>treatable disease and/or chemotherapy and<br>occasionally chronic (irreversible) BMF e.g.<br>myelodysplastic syndromes | <ul> <li>Prophylactic (WHO bleeding grade 0 or 1) to prevent bleeding</li> <li>&gt; Routine use in non-bleeding patients</li> <li>&gt; In the presence of additional risk factors for bleeding e.g. sepsis or abnormalities of haemostasis</li> </ul> |
| Peripheral platelet consumption/destruction e.g.<br>disseminated intravascular coagulation and immune<br>thrombocytopenia   | <b>Pre-procedure</b> to prevent bleeding expected to occur during surgery/invasive procedure  |
| Thrombocytopenia in critical care   | Therapeutic (WHO bleeding grade >2) to treat active bleeding  |
| Abnormal platelet function. Inherited or acquired disorders e.g. anti-platelet agents, uraemia  | <b>Contraindications to platelet transfusion unless</b><br><b>life-threatening haemorrhage</b><br>Thrombotic Thrombocytopenic Purpura (TTP)   |

### **Risks associated with platelet transfusion**

#### **Reduced effectiveness of future platelet transfusion** Alloimmunisation

#### Adverse effects

Febrile non-haemolytic transfusion reactions (FNHTR) and allergic reactions (including mild), reported incidence up to 3%. May require investigation to exclude other causes and prolong hospital stay.

# Estimated risk of moderate/severe reactions and infection transmission:

| FNHTR                                    | • 1 in 6,000                          |
|--|---------------------------------------|
| Allergic                                 | • 1 in 6,000                          |
| Haemolysis                               | • 1 in 600,000                        |
| Bacterial sepsis                         | • Rare since bacterial screening 2010 |
| Transfusion Related<br>Acute Lung Injury | • Less than 1 in 1,000,000            |
| <b>Hepatitis B infection</b>             | • 1 in 1,000,000                      |
| Hepatitis C infection                    | • 1 in 30,000,000                     |
| HIV infection                            | • 1 in 7,000,000                      |

# Prior to prescribing a platelet transfusion consider:



## Possible alternatives to platelet transfusion:

- Apply surface pressure after superficial procedures and correct surgical causes for bleeding
- Surgical patients expected to have at least a 500 ml blood loss, use tranexamic acid (TXA) unless contraindicated
- Trauma patients who are bleeding/ at risk of bleeding, early use of TXA
- Severe bleeding replace fibrinogen if plasma concentration less than 1.5 g/L
- Anti-platelet agents discontinue or if urgent procedure/bleeding use TXA if risk/benefit would support Uraemia with bleeding or preprocedure dialyse, correct anaemia, consider desmopressin
- Inherited platelet function disorders specialist haematology advice required. Consider desmopressin
- Chronic BMF with bleeding consider TXA

## Indications for use of platelet transfusions in adults

| Indication   | Transfusion indicated<br>(threshold)/not indicated |
|--|--|
| Prophylactic use (No bleeding or WHO grade 1)<br>One adult dose required   |  |
| - Reversible bone marrow failure (BMF) including allogeneic stem cell transplant   | 10 x 10 <sup>9</sup> /L                            |
| - Reversible BMF with autologous stem cell transplant (consider no prophylaxis)  | 10 x 10 <sup>9</sup> /L                            |
| - Critical illness   | 10 x 10 <sup>9</sup> /L                            |
| - Chronic BMF receiving intensive therapy  | 10 x 10 <sup>9</sup> /L                            |
| <ul> <li>Chronic BMF to prevent persistent bleeding of grade &gt; 2</li> </ul>   | Count variable                                     |
| <ul> <li>Chronic stable BMF, abnormal platelet function, platelet consumption/ destruction<br/>(e.g. DIC, TTP) or immune thrombocytopenia (ITP, HIT, PTP)</li> </ul> | Not indicated                                      |
| Prophylactic use in the presence of risk factors for bleeding (e.g. sepsis, antibiotic treatment, abnormalities of haemostasis)                                      |  |
| - Reversible/chronic bone marrow failure or critical care  | 10 to 20 x 10 <sup>9</sup> /L                      |
| - Abnormal platelet function, platelet consumption/destruction, immune thrombocytopenia  | Not indicated                                      |
| Platelet transfusion preprocedure  |  |
| - Central venous catheter (CVC) excluding PICC line  | 20 x 10 <sup>9</sup> /L                            |
| - Lumbar puncture  | 40 x 10 <sup>9</sup> /L                            |
| - Percutaneous liver biopsy  | 50 x 10 <sup>9</sup> /L                            |
| - Major surgery  | 50 x 10 <sup>9</sup> /L                            |
| - Epidural anaesthesia, insertion & removal  | 80 x 10 <sup>9</sup> /L                            |
| - Neurosurgery or ophthalmic surgery involving the posterior segment of the eye  | 100 x 10º/L  |
| Bone marrow aspirate or trephine biopsies, PICC line insertion, traction removal of central venous catheters (CVCs), cataract surgery                                | Not indicated                                      |
| Specific clinical conditions preprocedure-see below for indications  |  |
| Therapeutic use (Bleeding WHO grade 2 or above)  |  |
| -Severe bleeding   | 50 x 10 <sup>9</sup> /L                            |
| -Multiple trauma, brain or eye injury, spontaneous intracerebral haemorrhage   | 100 x 10 <sup>9</sup> /L                           |
| -Bleeding (WHO grade >2) but not severe  | 30 x 10 <sup>9</sup> /L                            |
| -Bleeding in specific clinical conditions – see the next table for indications   |  |

| Conception of |  |
|---------------|--|
| Specific cl   |  |
|               |  |

| Platelet function defect  |  |  |
|---|--|--|
| - Congenital – Preprocedure or therapeutic use. When alternative therapy contraindicated or ineffective. Directed by specialist in haemostasis.   | Count Variable   |  |
| - Acquired (anti-platelet agents, uraemia)- only indicated for severe bleeding  |  |  |
| Disseminated intravascular bleeding   |  |  |
| Preprocedure or therapeutic use. Consider threshold counts above but may not be achievable and individual case review required  | Use preprocedure or<br>therapeutic threshold<br>as guide |  |
| Thrombotic thrombocytopenic purpura   |  |  |
| Platelet transfusion contraindicated unless life-threatening bleeding   | Count Variable   |  |
| Immune thrombocytopenia   |  |  |
| (ITP, HIT, PTP). Preprocedure when other therapy ineffective/procedure urgent or to treat severe bleeding. Consider threshold counts above but may be unachievable or unnecessary and individual case review required | Use preprocedure or<br>therapeutic threshold<br>as guide |  |
| Abbreviations   |  |  |
| Disseminated intravascular coagulation (DIC), peripherally inserted central catheter (PICC), thrombotic thrombocytopenic purpura (TTP), primary immune thrombocytopenia (ITP),  |  |  |

heparin-induced thrombocytopenia (HIT), post-transfusion purpura (PTP)