

## Summary of Guidelines for the Use of Platelet Transfusions in a Platelet Shortage

### British Society for Haematology Guideline (2016) Adults

### British Society for Haematology Guideline (2016 & 2020 addendum) Children, Neonates

#### 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

#### Prior to prescribing a platelet transfusion consider:

What is the indication for transfusion in this patient?

Are there any alternatives which could be used instead?

Is the patient aware of the benefits, harms and alternatives to a platelet transfusion?

#### 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 (or >10% blood volume in children), use tranexamic acid (TXA) unless contraindicated
- Trauma patients who are bleeding or 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 pre-procedure – dialyse, correct anaemia, consider desmopressin
- Inherited platelet function disorders - specialist haematology advice required. Consider desmopressin
- Chronic Bone Marrow Failure (BMF) with bleeding – consider TXA

#### Indications for use of platelet transfusions in adults and children (AMBER ALERT)

Indication	Transfusion indicated (threshold)/ not indicated
Prophylactic use (No bleeding or WHO grade 1) - one adult dose required (or weight-based paediatric equivalent)	
<ul style="list-style-type: none"> <li>• Reversible Bone Marrow Failure, including allogeneic stem cell transplant</li> <li>• Critical illness</li> <li>• Reversible and Chronic Bone Marrow Failure receiving intensive therapy</li> <li>• Chronic Bone Marrow Failure to prevent persistent bleeding of grade &gt; 2</li> </ul>	10 x 10 <sup>9</sup> /L 10 x 10 <sup>9</sup> /L 10 x 10 <sup>9</sup> /L Count variable
<ul style="list-style-type: none"> <li>• Chronic stable Bone Marrow Failure, abnormal platelet function, platelet consumption/ destruction (e.g. DIC, TTP) or immune thrombocytopenia (ITP, HIT, PTP)</li> <li>• Reversible BMF with autologous stem cell transplant (patient stable)</li> </ul>	NOT INDICATED  NOT INDICATED
Prophylactic use in presence of risk factors for bleeding (e.g. sepsis, abnormalities of haemostasis)	
<ul style="list-style-type: none"> <li>• Reversible/chronic bone marrow failure, or critical care</li> </ul>	10 to 20 x 10 <sup>9</sup> /L
<ul style="list-style-type: none"> <li>• Abnormal platelet function, platelet consumption/destruction (e.g. TTP), immune thrombocytopenia</li> </ul>	NOT INDICATED
Pre-procedure	
<ul style="list-style-type: none"> <li>• Central venous catheter (CVC) tunnelled or untunnelled (excluding PICC line)</li> <li>• Lumbar puncture*</li> <li>• Percutaneous liver biopsy</li> <li>• Major surgery</li> <li>• Epidural anaesthesia, insertion &amp; removal</li> <li>• Neurosurgery or ophthalmic surgery involving the posterior segment of the eye</li> </ul>	20 x 10 <sup>9</sup> /l 40 x 10 <sup>9</sup> /l 50 x 10 <sup>9</sup> /l 50 x 10 <sup>9</sup> /l 80 x 10 <sup>9</sup> /l 100 x 10 <sup>9</sup> /l
<ul style="list-style-type: none"> <li>• Bone marrow aspirate or trephine biopsies, PICC line insertion, traction removal of central venous catheters (CVCs), cataract surgery, other procedures with low-risk of bleeding</li> </ul>	NOT INDICATED
Therapeutic use (Bleeding WHO grade 2 or above)	
<ul style="list-style-type: none"> <li>• Severe bleeding</li> <li>• Multiple trauma, brain or eye injury, spontaneous intracerebral haemorrhage</li> <li>• Bleeding (WHO grade &gt;2) but not severe</li> <li>• Bleeding in specific clinical conditions – see table next page for indications</li> </ul>	50 x 10 <sup>9</sup> /L 100 x 10 <sup>9</sup> /L 30 x 10 <sup>9</sup> /L

## Indications for use of platelet transfusions in neonates (AMBER ALERT)

Indication	Transfusion indicated (threshold)/ not indicated
Prophylactic use (No bleeding or WHO grade 1)	
<ul style="list-style-type: none"> <li>• Neonate (including very pre-term)</li> <li>• Neonate with NAIT (no family history of ICH)</li> </ul>	25 x 10 <sup>9</sup> /L 25 x 10 <sup>9</sup> /L
Prophylactic use in presence of risk factors for bleeding (e.g. sepsis)	
<ul style="list-style-type: none"> <li>• Preterm neonate with sepsis</li> <li>• Neonate with NAIT (Family history of ICH)</li> </ul>	25 x 10 <sup>9</sup> /L 50 x 10 <sup>9</sup> /L
Pre-procedure	
<ul style="list-style-type: none"> <li>• Lumbar puncture*</li> <li>• Major surgery</li> <li>• Neurosurgery</li> </ul>	40 x 10 <sup>9</sup> /l 100 x 10 <sup>9</sup> /l 100 x 10 <sup>9</sup> /l
Procedures with low-risk of bleeding	NOT INDICATED
Therapeutic use (Bleeding WHO grade 2 or above)	
<ul style="list-style-type: none"> <li>• Severe bleeding</li> </ul>	100 x 10 <sup>9</sup> /L

### Specific clinical conditions

Platelet function defect	
<ul style="list-style-type: none"> <li>• Congenital – Pre-procedure or therapeutic use. When alternative therapy contraindicated or ineffective. Directed by specialist in haemostasis.</li> <li>• Acquired (anti-platelet agents, uraemia)- only indicated for severe bleeding</li> </ul>	Count Variable
Disseminated intravascular bleeding	
<ul style="list-style-type: none"> <li>• Pre-procedure or therapeutic use. Consider threshold counts above but may not be achievable and individual case review required</li> </ul>	Use pre-procedure or therapeutic threshold as guide
Thrombotic thrombocytopenic purpura	
<ul style="list-style-type: none"> <li>• Platelet transfusion contraindicated unless life-threatening bleeding</li> </ul>	
Immune thrombocytopenia (excluding NAIT)	
<ul style="list-style-type: none"> <li>• (ITP, HIT, PTP) Pre-procedure when other therapy ineffective or procedure urgent or to treat severe bleeding. Consider threshold counts above but may be unachievable or unnecessary and individual case review required</li> </ul>	Use pre-procedure or therapeutic threshold as guide

#### Footnotes

\*It is accepted that prior to lumbar puncture some clinicians will transfuse platelets at higher counts (e.g. 50 x 10<sup>9</sup> /L) in clinically unstable children, non ALL patients, or for the first LP in newly-diagnosed ALL patients to avoid haemorrhage and cerebrospinal fluid contamination with blasts, or at lower counts (≤ 20 x 10<sup>9</sup> /L) in stable patients with ALL, depending on the clinical situation. These practices emphasise the importance of considering the clinical setting and patient factors.

#### Abbreviations

ALL acute lymphocytic leukaemia; BMF bone marrow failure; DIC Disseminated intravascular coagulation; HIT heparin-induced thrombocytopenia; ICH intracranial haemorrhage; ITP primary immune thrombocytopenia; LP lumbar puncture; NAIT neonatal alloimmune thrombocytopenia; PICC peripherally inserted central catheter; PTP post-transfusion purpura; TTP thrombotic thrombocytopenic purpura: