

| Guideline Number | Guideline Title  |
|------------------|--|
| 1                | <a href="#"><u>Informed Consent for Administration of Blood, Blood Components, and/or Plasma Protein Products</u></a>    |
| 2                | <a href="#"><u>Patient Identification in Specimen Collection for Pre-Transfusion Testing</u></a>                         |
| 3                | <a href="#"><u>Patient Identification in Blood, Blood Components and/or Plasma Protein Products</u></a>                  |
| 4                | <a href="#"><u>Receipt of Blood, Blood Components and/or Plasma Protein Products</u></a>                                 |
| 5                | <a href="#"><u>Monitoring of Patients Receiving Transfusion</u></a>  |
| 6                | <a href="#"><u>Patient Required Health Record Documentation of Blood, Blood Products and Plasma Protein Products</u></a> |
| 7                | <a href="#"><u>Transfusion Reaction - Identification, Management, and Reporting</u></a>                                  |
| 8                | <a href="#"><u>Administration of Blood and Blood Components</u></a>  |
| 9                | <a href="#"><u>Administration of Plasma Protein Products (Derivatives)</u></a>   |
| 10               | <a href="#"><u>Education Requirements for Patients Receiving Transfusion</u></a>   |
| 11               | <a href="#"><u>Nurses Performing Laboratory Duties</u></a>   |