Hemovigilance Module: A set of standardized national surveillance procedures used in the monitoring of transfusion-associated adverse reactions aimed at improving patient safety, minimizing morbidity and mortality of transfusion recipients, and identifying emerging complications and infections associated with recipient blood transfusions.

CMS reporting requirements for Facilities

Reporting to the Hemovigilance Module is not required by CMS.

Optional Reporting for Facilities

Title	Link	Reporting Frequency
Annual Facility Survey	Acute Care Facility Annual Survey Non-Acute Care Facility Annual Survey	Yearly
Monthly Reporting Plan	Monthly Reporting Plan	Monthly
Adverse Reactions (forms 57.307-57.320)	Adverse Reactions Forms	Monthly
Incidents	Incident Form	When incident is discovered (all incidents associated with a reported adverse reaction must be reported to NHSN using the corresponding incident form)
Monthly Incident Summary	Monthly Incident Summary Form	Monthly
Monthly Reporting Denominators	Monthly Reporting Denominators Form	Monthly

Additional Resources

Title	Link(s)
Hemovigilance Module Resources*	• Guidance for reporting to the Hemovigilance Module • Hemovigilance Protocol • HV Discard Guidance and FAQ • Linking Incident Records to Adverse Reaction Records *See Quick Reference Guides for additional resources
NHSN Administrator Training Videos	• NHSN Administrator Training | NHSN (cdc.gov)