Organisation name Site or unit		FORM Adverse reaction report	
Document ID	BLD-FRM-003	**Version**	1.0
Status	Blank form, to adapt	**Issue date**	
Review date		**Replaces**	
Owner		**Approved by**	
Signature			

Form · BLD-FRM-003 · Version 1.0 · Approved by the ASF Expert Committee

Supports: 5.7

1. Purpose

Records a reaction in a donor or a recipient, its investigation and its report.

2. How to complete

- Complete at once.
- Record the investigation outcome.
- Record the report number.

3. Checks

Check	Entry
Donor or recipient	Yes
Unit number and time	Yes
Reaction described	Yes
Investigation outcome and action	Yes
Reported to the national system (number)	Yes or N/A

4. Evidence

Reference	Note
E-009	Based on the Council of Europe Guide to the Preparation, Use and Quality Assurance of Blood Components.
E-006	ASF Policy Framework v1.2. Adapted.
E-009	ASF Expert Committee consensus.

5. Change history

Version	Date	What changed	Approval
1.0	the date of approval	First version	set Expert Committee approval

6. References

- Accréditation Sans Frontières (2026). ASF Policy Framework v1.2. (E-006)
- Accréditation Sans Frontières (2026). ASF Expert Committee consensus on Standard 1. (E-009)

Download this document as PDF