Organisation name Site or unit		FORM Adverse reaction record	
Document ID	PRO-FRM-006	**Version**	1.0
Status	Blank form, to adapt	**Issue date**	
Review date		**Replaces**	
Owner		**Approved by**	
Signature			

Form · PRO-FRM-006 · Version 1.0 · Approved by the ASF Expert Committee

Supports: 11.9

1. Purpose

Records each adverse reaction to a product, the batch and the report made.

2. How to complete

- Record the reaction and the batch at once.
- Record the treatment.
- Record the report made.

3. Checks

Check	Entry
Product and batch	Yes
Patient ID and procedure	Yes
Reaction and time	Yes
Treatment given	Yes
Reported (yes or no) and date	Yes

4. Evidence

Reference	Note
E-009	Based on the ASF Expert Committee consensus and the WHO Guidelines for Safe Surgical Care and Safe Procedures (2009).
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)

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