Organisation name Site or unit		REGISTER Product sourcing and labelling register	
Document ID	TRM-REG-002	**Version**	1.0
Status	Blank form, to adapt	**Issue date**	
Review date		**Replaces**	
Owner		**Approved by**	
Signature			

Standards › Standard 41 › Registers and logs

Register · TRM-REG-002 · Version 1.0 · Approved by the ASF Expert Committee

Supports: 41.4

1. Purpose

Records each product supplied, its source, its label content and its claims check.

2. How to complete

- Record each product and its licensed source.
- Record the label content checked.
- Record the claims check.

3. Checks

Check	Entry
Product and licensed source	Yes
Ingredients and warnings on label	Yes
Unlicensed claim found (yes or no)	Yes
Check date and checker	Yes

4. Evidence

Reference	Note
E-009	Based on the WHO Traditional Medicine Strategy 2014–2023, the WHO guidance on substandard and falsified medical products, and the ASF Hospital standard for informed consent and incident reporting.
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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