Organisation name Site or unit		FORM Result authorisation record	
Document ID	LBT-FRM-005	**Version**	1.0
Status	Blank form, to adapt	**Issue date**	
Review date		**Replaces**	
Owner		**Approved by**	
Signature			

Form · LBT-FRM-005 · Version 1.0 · Approved by the ASF Expert Committee

Supports: 33.12

1. Purpose

Records the authorisation of each result, the authoriser and the time, before release.

2. How to complete

- Check the result before authorising it.
- Record the authoriser's name and the time.
- Confirm the authoriser is competent for the test.

3. Checks

Check	Entry
Result identifier and test	Yes
Authorised by (name, competence confirmed)	Yes
Authorisation time before release	Yes
Release time	Yes

4. Evidence

Reference	Note
E-009	Based on ISO 15189:2022 Medical laboratories: requirements for quality and competence, ISO 9001:2015 for supplier control, and the Global Patient Safety Action Plan 2021–2030 for critical results.
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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