Organisation name Site or unit		REGISTER Implant and prosthetic component traceability register	
Document ID	DEN-REG-005	**Version**	1.0
Status	Blank form, to adapt	**Issue date**	
Review date		**Replaces**	
Owner		**Approved by**	
Signature			

Register · DEN-REG-005 · Version 1.0 · Approved by the ASF Expert Committee

Supports: 31.20

1. Purpose

Records each implant or prosthetic component placed, with its lot, supplier and patient, so that any lot can be traced.

2. How to complete

- Record each component at the time it is placed.
- Link the component to the patient record.
- Test the trace of one lot at the set interval.

3. Checks

Check	Entry
Patient and date placed	Yes
Component, lot and supplier	Yes
Patient record reference	Yes
Trace test (date and time to trace)	Yes

4. Evidence

Reference	Note
E-009	Based on the CDC guidelines for infection control in dental health-care settings (Kohn et al., 2003), ISO 17665 for moist-heat sterilisation, the Minamata Convention on Mercury (2013), IAEA Specific Safety Guide SSG-46 (2018), Council Directive 2013/59/Euratom, and the ADA guidelines on sedation (2016).
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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