

# TRACE AROUND THE BLOCK!

## VALIDATION OF AUTOMATED INTEGRATION OF IMPLANTABLE MEDICAL DEVICE TRACEABILITY DATA INTO AN ELECTRONIC PATIENT RECORD

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### BACKGROUND AND IMPORTANCE

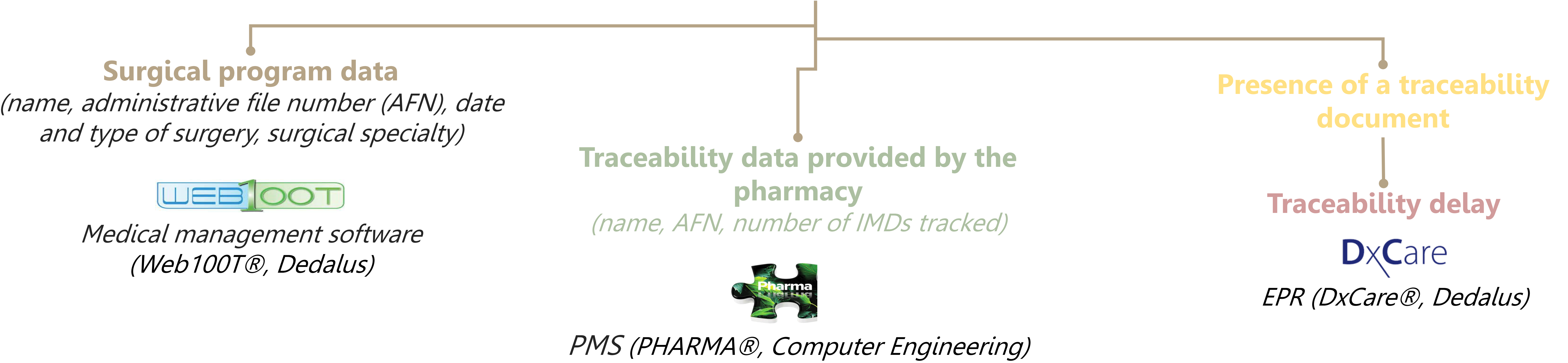
- French regulations require **traceability** of Implantable Medical Devices (IMDs) to be recorded on **discharge documents** and in the **Electronic Patient Record (EPR)**
- 2023** Audit showing that only **69,5% of patients** EPR (Electronic Patient Record) mentioned the type of IMD used
- 2024** Development of an **HL7 interface** between our Pharmaceutical Management Software (PMS) and our EPR that **automatically** uploads to the EPR a file specifying the **traceability data of IMDs entered in the PMS**

### AIM AND OBJECTIVES

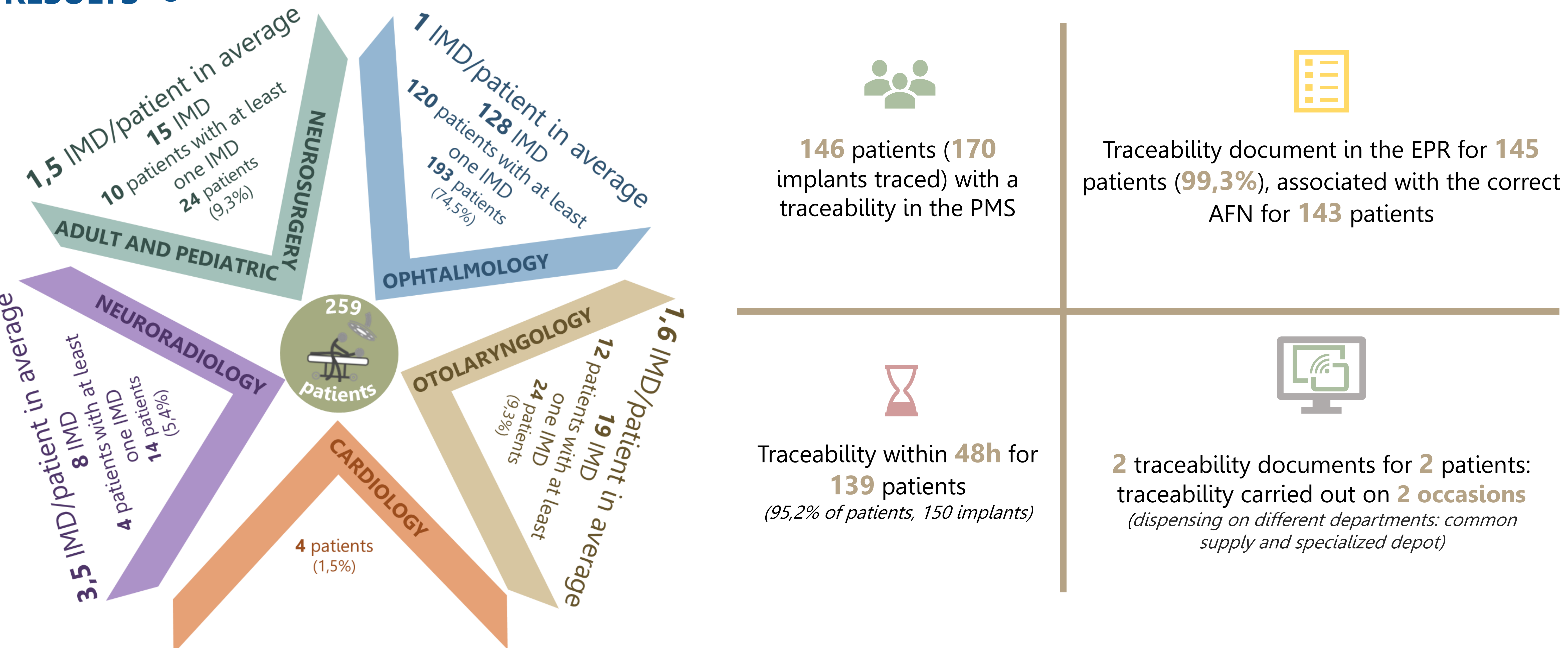
Validate the **data transfer automation** from the Pharmaceutical Management Software (PMS) to the EPR via an interface

### MATERIALS AND METHODS

« Single day » audit (single observer) on 3 independent days (August and September 2024)



### RESULTS



### CONCLUSION AND RELEVANCE

- Automation of traceability data transfer (**99.3%**)
- Complete EPR with exhaustive health traceability
- Toward an implementation of the final stage in the data transfer automation (from EPR to shared digital medical record)

