Multisite Clinical Trials: What about the Experimental Drug Circuit? Failure Modes, Effects and Criticality Analysis

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PHARMACIE
A USAGE INTERIEUR

RECHERCHE CLINIQUE
ET INNOVATION

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BACKGROUND AND IMPORTANCE

>Multisite coordination of clinical trials —— Key lever to promote patient recruitment, facilitate access to innovation and enhance the attractiveness of sites

> Requirement for coordination at both clinical research and pharmaceutical levels

>Involvement of multiple pharmacies in the distribution of the experimental product — Differences from usual management — New risks

AIM AND OBJECTIVES



The TRINITI project aims to identify, assess, and mitigate the risks associated with this pharmaceutical activity to secure the circuit of the experimental product.

MATERIALS AND METHODS

Establishment of a Multidisciplinary Working Group

- Coordinating pharmacy: 1 TRINITI project manager, 1 quality project manager, 2 pharmacists
- Coordinating clinical research unit: 1 project manager
- Pharmacy of a satellite center: 1 pharmacist





1) Mapping of device-specific processes and sub-processes



2) Identification of risks associated with these processes



Risks with a potential severity of 5

satellite center

3) Rating of each identified risk in terms of Severity, Frequency, and Detectability -> Criticality

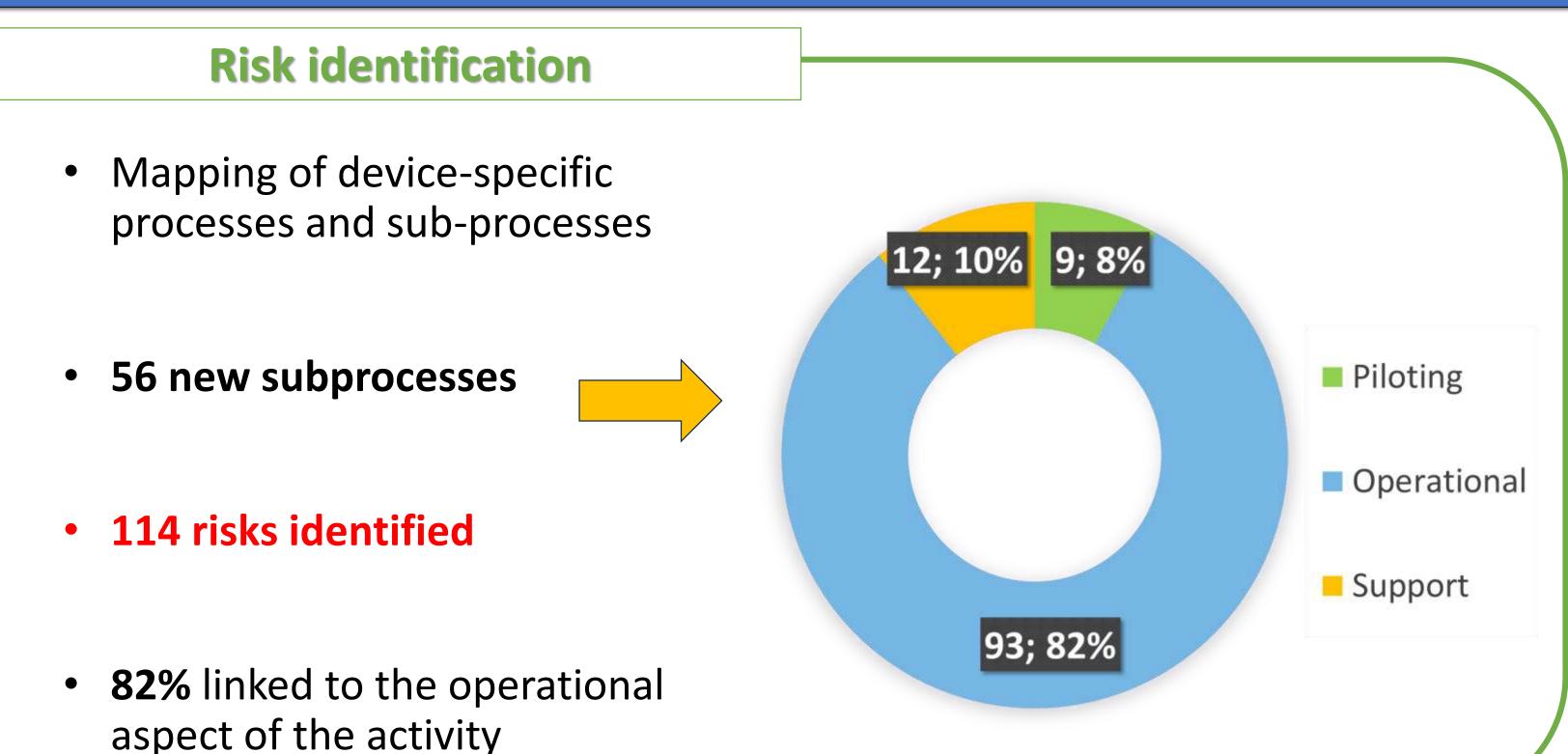


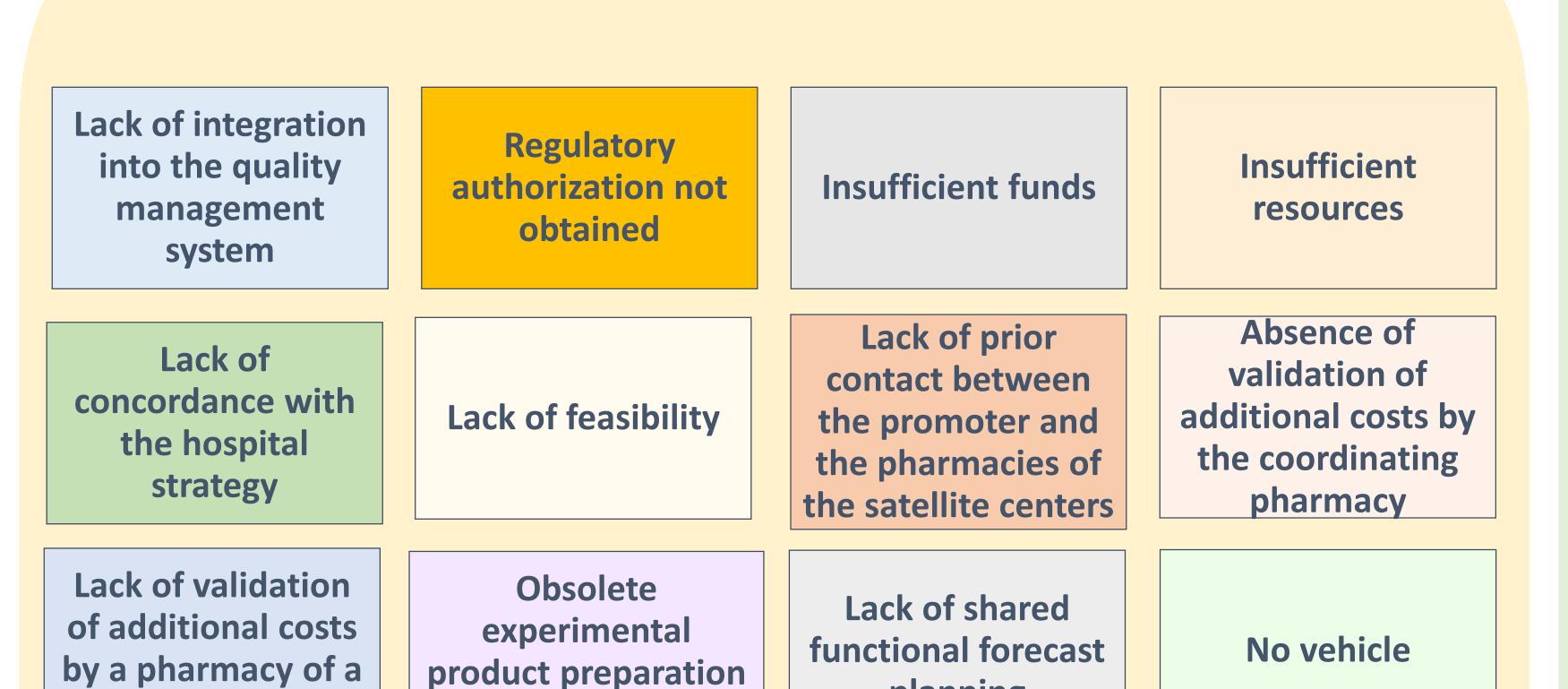
4) Risk prioritization



5) Existing and/or To-Be-Implemented Preventive Barriers

RESULTS

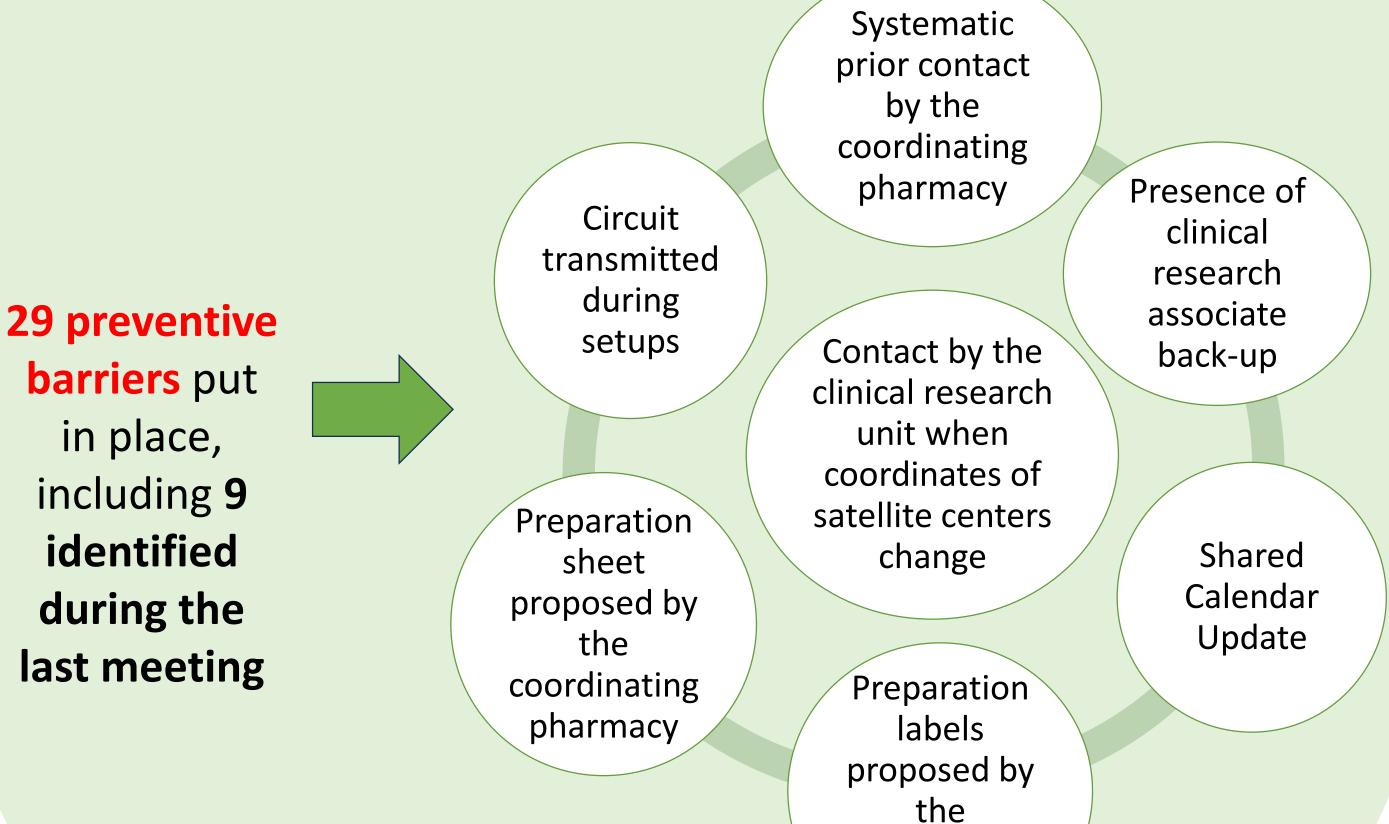




Analysis of the criticalities obtained

Acceptable risks
Tolerable risks
Unacceptable risks

- Majority acceptable risks (78%)
- 30% of tolerable risks related to the preparation of clinical trials in satellite center pharmacies
- Two unacceptable risks identified related to the financial viability of the project and the availability of the transport vehicle



83% of these risks with barriers already in place

sheet

CONCLUSION AND RELEVANCE

- >The most critical steps in the pharmaceutical circuit are identified, facilitating the implementation of preventive measures
- The need to assess residual criticality after implementing these measures
- In continuation of this work, a post-analysis of risks should be conducted based on initial experiences

planning



coordinating

pharmacy