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



A.Mobarek¹, C. Metz¹, N. Zeggagh¹, A.Jacob², A.Touati³, M. Antignac¹, F. Charbonnier-Beaupel¹, M. Hinterlang¹

1: Pharmacy, REQPHARM Unit, Pitié-Salpêtrière Hospital

2: Pharmacy, Lariboisière Hospital

3: Clinical research unit, Saint-Antoine Hospital

BACKGROUND AND IMPORTANCE

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

 \geq 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

Use of the FMECA method : Failure Mode, Effects and Criticality Analysis (FMECA)



Contact by the back-up



CONCLUSION AND RELEVANCE

>The most critical steps in the pharmaceutical circuit are identified, facilitating the implementation of preventive measures

 \succ 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

