How can your stakeholder group contribute to the successful implementation of the CTR, and what does success look like?

- Successfull transition of all running clinical trials in CTIS
- Reduction of administrative burden (CTIS)

>> highly demanding in human resources

- Adaptation of CTIS to master protocols especially platform trials
- Harmonisation of ethical review >> voluntary harmonisation
- Low-intervention trials under-used: review of risk categories and corresponding requirements in EU CTR