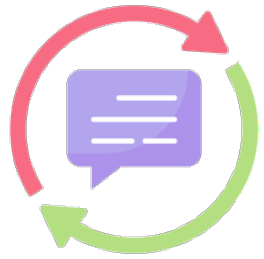


From your stakeholder group perspective, what needs to be put in place to create an invigorated clinical trials environment?

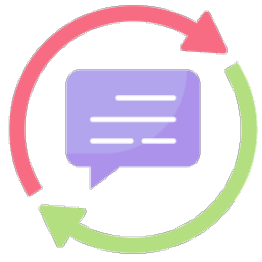
2 main ideas



Feedback loop



Share data by default



Patient-centric feedback Loop

- Patient organizations can facilitate a continuous **feedback loop** by partnering with regulatory bodies and clinical trial sponsors
- They could implement simple surveys or feedback forms at **different stages** of clinical trials
- This data should be **regularly reviewed** and used to make incremental changes to the clinical trial regulations patients' perspectives on the process



Share data by default after CT

- Patient organizations can advocate for and help implement a “**Share-by-Default**” approach to clinical trial data
- After a trial concludes, anonymized patient data can be made **available** through a secure, accessible online portal
- This platform would allow patients, researchers, and healthcare professionals to review and analyze data, contributing to a more **transparent and collaborative** environment
- Patients could opt-in to receive **updates** on how their data is being used and what discoveries or advancements have been made as a result
- Additionally, patients should have the ability to **withdraw** their data if they change their mind