Shortening development and review timelines, thinking more like a regulator

**EUAs during COVID** – 600

**EUAs** amidst other public health emergencies – 65

Increased collaboration between regulatory agencies and industry enabled shorter review timelines for COVID-19 drugs and treatments

Regulators around the world worked hand-in-hand with industry rather than just having an endpoint type of review, and it is anticipated that these run-on trials will be the case post-COVID

The future of regulatory is digital and collaborative

“Think like a regulator”

- Patient safety first
- Risk based approach
- Objective evidence
- A controlled process

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