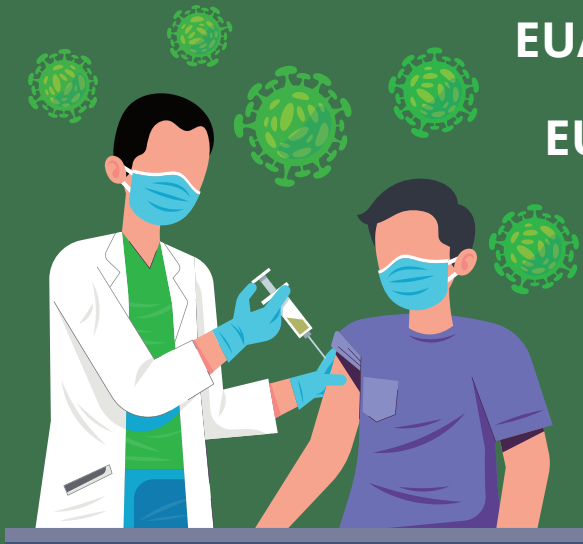


Shortening development and review timelines, thinking more like a regulator



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

“Think like a regulator”



The future of regulatory is **digital** and **collaborative**



Patient safety first

Risk based approach



Objective evidence

A controlled process



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