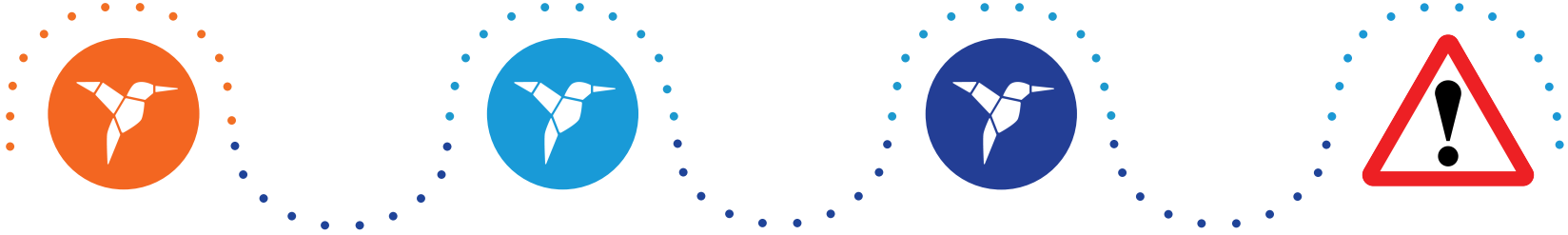
Using the CTIS System for EU Trial Applications

The Clinical Trial Information System (CTIS) is an electronic portal launched by the European Medicines Agency (EMA) as a part of the new EU regulation (536/2014).



From 31 January 2022 until January 2023

All initial clinical trial applications can be submitted under either the Directive or the Clinical Trial Regulation.

Before 31 January 2023

Trials approved under EU-CTD can continue to be regulated under EU-CTD until 31 January 2025.

Starting on 31 January 2023

All initial clinical trial applications need to be submitted under the Clinical Trial Regulation.

From 31 January 2025

All ongoing and new clinical trials must ultilize the CTIS System.

QPS

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