

# Key Information for Sponsors on CTIS

The Clinical Trials Regulation (CTR) ensures consistent rules for clinical trials throughout Europe and harmonises assessment and supervision via the Clinical Trials Information System (CTIS).

[euclinicaltrials.eu](https://euclinicaltrials.eu)



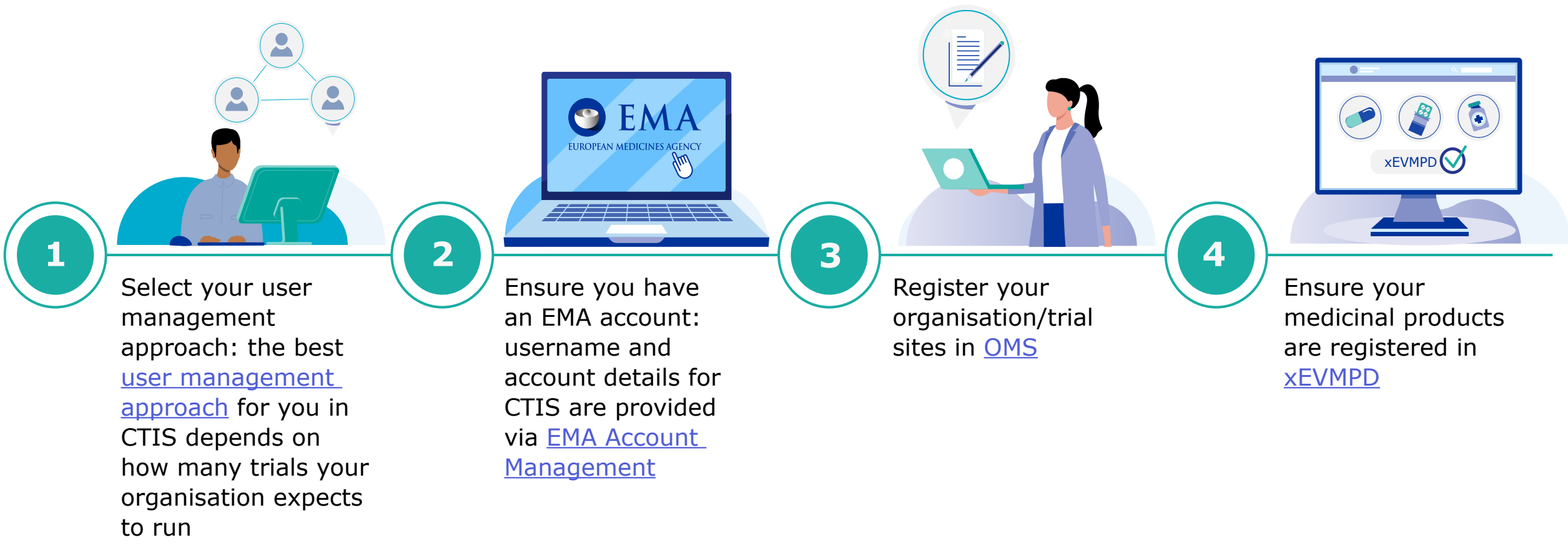
## Transition period

There will be a transition period from 2022 to 2025:



## Getting started with CTIS

To get started with CTIS, sponsors must decide their user management approach and complete registrations:



Find more information about getting started with CTIS in [the Getting started quick guide](#).

## CTIS Go-Live

Go to [euclinicaltrials.eu](https://euclinicaltrials.eu) to learn more and to access the CTIS secure Sponsor workspace



## Key links for clinical trials sponsors

 CTIS training and support <a href="#">LEARN MORE</a>	 Online modular training on CTIS functionalities <a href="#">LEARN MORE</a>	 Guide to CTIS training catalogue <a href="#">LEARN MORE</a>
 CTIS Sponsor Handbook <a href="#">LEARN MORE</a>	 CTIS Newsletter <a href="#">LEARN MORE</a>	 Information on the Clinical Trials Regulation <a href="#">LEARN MORE</a>