



ACT EU workplan and key priorities

Expected contribution from MSP AG

Inaugural meeting of the ACT EU Multi-stakeholder Platform Advisory Group
20 March 2024



ACT EU is delivering benefits to clinical trial stakeholders across key areas:



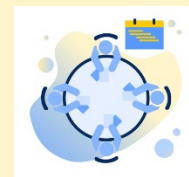
Mapping &
governance



Implementation of
the Clinical Trials
Regulation*



Multinational clinical
trials by non-
commercial
sponsors*



Multi-stakeholder
platform*



Good clinical
practice
modernisation



Clinical trials
analytics



Scientific advice*



Clinical Trials
methodologies



Clinical trials
safety



Clinical trials
training
curriculum



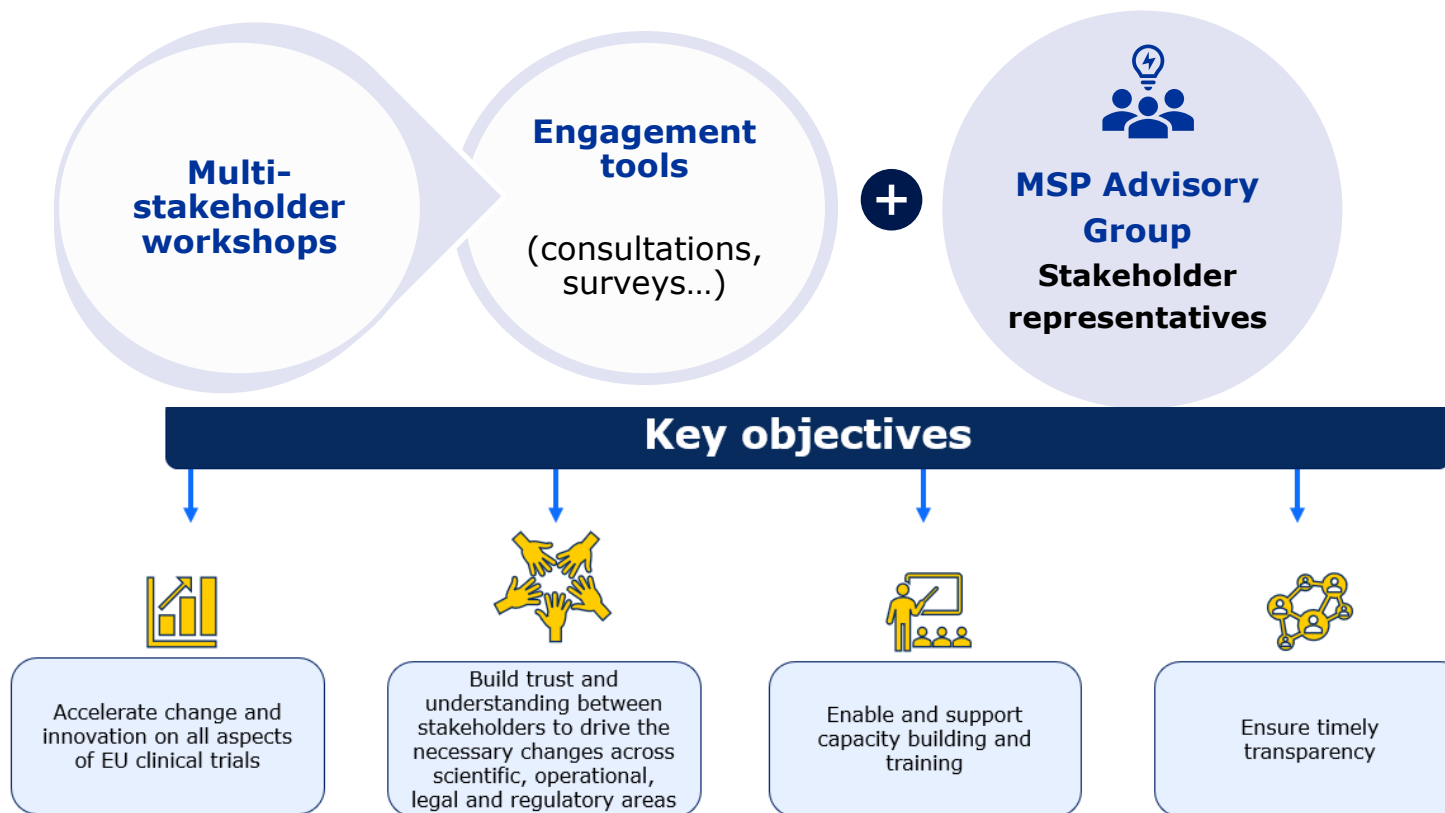
Clinical trials in
public health
emergencies*

Over the next 4 years ACT EU aims to

- Establish the **Multi-Stakeholder Platform** (MSP) and its **Advisory Group** as a vehicle for dialogue and collaboration with stakeholders
- Ensure **effective operation** of the clinical trials (CT) regulation
- **Simplify governance** and **align CT approval** with scientific advice
- Support academic sponsors to conduct **impactful clinical trials**
- Establish the place of **novel methodologies**
- Enable **decentralised approaches**
- Training: create an **educational “ecosystem”** (leveraging existing initiatives)
- **Align internationally** (including Good Clinical Practice, GCP)
- **Optimise the use of data** about clinical trials for better research and decision-making
- Enable CTs in **public health emergencies (PHEs)**



Multi-stakeholder platform



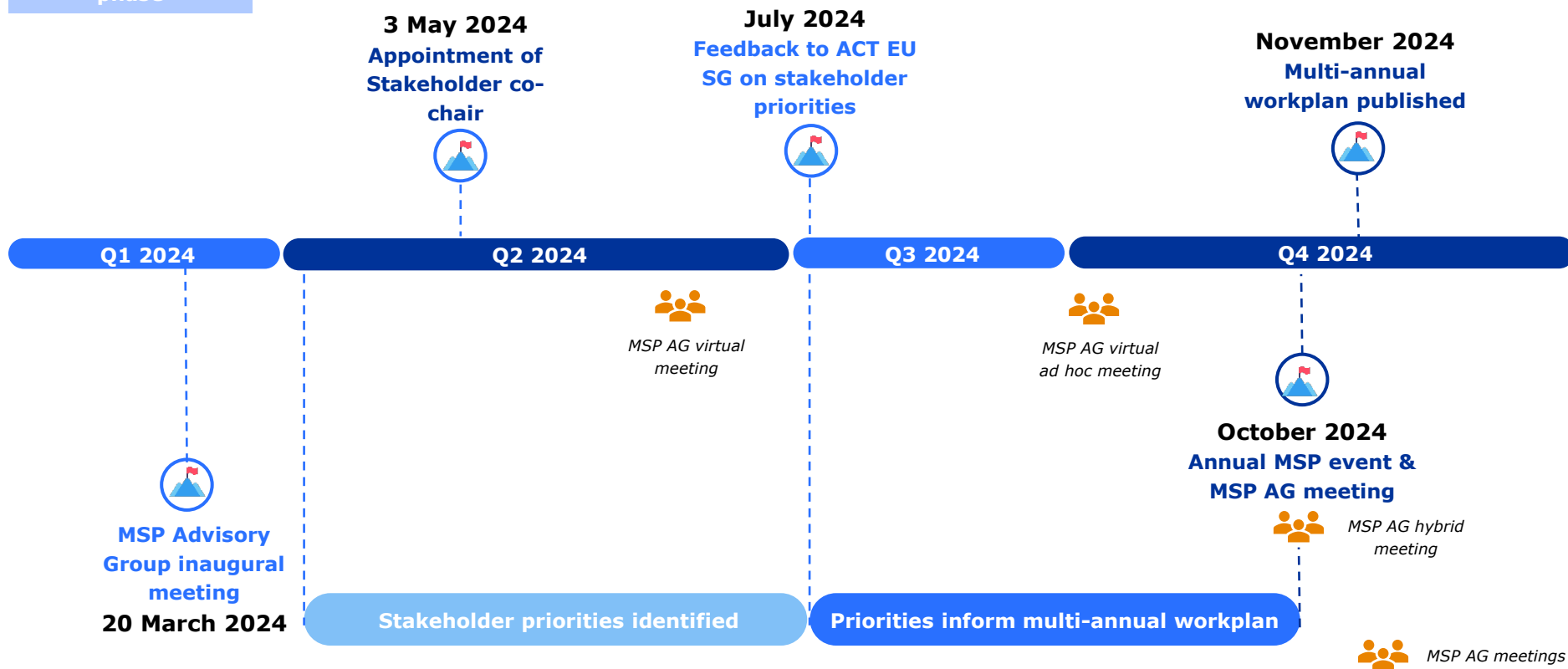
How stakeholders are involved

- Providing the ACT EU Steering Group with their views and strategic advice on the [ACT EU multi-annual workplan](#)
- Identifying stakeholder needs, concerns, challenges and priorities, and communicating these to the ACT EU regulatory partners
- Advising ACT EU regulatory partners on stakeholder engagement and communication
- Keeping their respective stakeholder group informed of the output from ACT EU MSP initiatives and overall ACT EU activities
- Reviewing and agreeing on the mandate, workplan and any governance-related documents of the advisory group

Strategic input into ACT EU workplan

2024
Establishment
phase

Identifying stakeholder needs, concerns, challenges and priorities



Operational input into ACT EU events



2024
Establishment
phase

Advising ACT EU regulatory partners on stakeholder engagement and communication

Apr 2024
SNSA webinar for
sponsors

3 May 2024
Appointment of
Stakeholder co-
chair

July 2024
Feedback to ACT EU
SG on stakeholder
priorities

**Workshop on CT
methodologies**

November 2024
Multi-annual
workplan published

Q1 2024

Q2 2024

Q3 2024

Q4 2024

Q1 2025



MSP Advisory
Group inaugural
meeting
20 March 2024



**Launch of
consolidated advice
pilots
April/May 2024**



MSP AG virtual
meeting

**Training on
consolidated advice
pilots
May 2024**



MSP AG virtual
ad hoc meeting



October 2024
Annual MSP event &
MSP AG meeting



MSP AG hybrid
meeting



**Workshop on
ICH E6 R3**

Written consultations and ad hoc meetings as needed



MSP AG meetings

Let's discuss

How can we ensure comprehensive identification of stakeholder priorities and needs, and integration into the work of ACT EU?

How can we ensure widespread sharing of information?

What are the essential ingredients for ensuring the success of the MSP AG?

Any questions?

Further information

misp-agsecretariat@ema.europa.eu

[ACT EU website](#)

[Clinical trials in human medicines | European Medicines Agency \(europa.eu\)](#)

Official address Domenico Scarlattilaan 6 • 1083 HS Amsterdam • The Netherlands

Telephone +31 (0)88 781 6000

Send us a question Go to www.ema.europa.eu/contact

Follow us on  **@EMA_News**