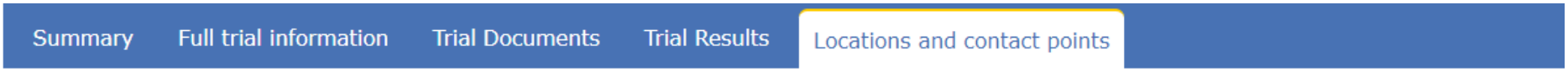
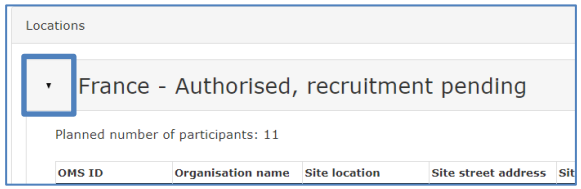
20 September 2024
EMA/441148/2024

CTIS public portal: Locations and contact points

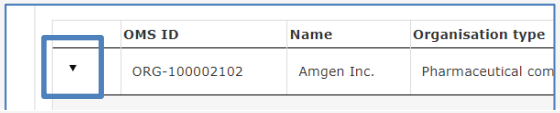


This section lists the countries and specific locations where the trial is conducted, the number of participants and sponsor details. You can see the relevant descriptions in the table below. To **view the detailed list of trial locations and the contact details of the main healthcare professional responsible for the trial**, you can click on the relevant country's name:



The participation criteria of a trial (i.e. criteria under which its participants are included or excluded from recruitment) can be seen in the section 'Full trial information'. Useful participants' documents such as the informed consent form or recruitment arrangements are available in the 'Trial documents' section.

You can also have a look at the patient/consumer organisations as well as healthcare professional organisations who regularly work with the European Medicines Agency (EMA) and may be able to provide information on clinical trials within various disease areas.

Term	Definition
Locations	
The following details are displayed per country, once you expand its box.	
Planned number of participants	The expected number of participants who will be enrolled in the clinical trial.
Site location	The exact location where the trial is foreseen to be conducted, at the time of its authorisation. The name of the organisation (generally a hospital), as well its address and *contact details of the main healthcare professional responsible for the trial* are provided here, and can be used by patients to ask for information.
Countries outside of the European Economic Area	
List of countries	A list of all the countries outside of the European Economic Area where the clinical trial is conducted. The European Economic Area (EEA) includes all the EU Member States and Iceland, Liechtenstein, and Norway. More information here.
Participants in the rest of the world	Planned number of trial's participants in countries outside of the European Economic Area.
Sponsor	
Sponsor	The person, company, institution or organisation which takes responsibility for the initiation, for the management and for setting up the financing of a clinical trial. There are two main types of sponsors: Commercial Sponsors (e.g., pharmaceutical companies) and Non-commercial Sponsors (e.g., universities or academic research centres). The sponsor's address, together with its main contact points are also displayed: the *Scientific contact point* for scientific or technical questions, while the *Public contact point* for general information about the clinical trial. In case the trial has more than one sponsor, the co-sponsor's details can be displayed through expanding the relevant table. 
Third parties	If applicable, organisations or individuals contracted by the sponsor to manage certain aspects of the clinical trial, e.g. Clinical Research Organisations (CROs).

Term	Definition
Legal representative	In case the sponsor the clinical trial is not established in the European Economic Area, a legal representative is nominated as the person or organisation designated to act on behalf of the sponsor in the European Economic Area, ensuring the trial complies with local regulations.