EU CTIMPS

If you are planning a clinical trial which involves administering medicinal products to participants, your study may fall under the EU Clinical Trials Directive. These trials are referred to as Clinical Trials of Investigational Medicinal Products (CTIMPs) and will need to meet more regulatory requirements than other kinds of clinical research projects.

Please contact the Cambridge Clinical Trials Unit (CCTU) in the first instance at cctu@addenbrookes.nhs.uk