

## COMPREHENSIVE INVENTORY

### PROVIDER PROFILE

<b>Name of the organisation</b>	University of Oxford – Clinical Trials and Research Governance (CTRG), Research Services
<b>Type of the organisation</b>	Non-profit
<b>Contact email(s)</b>	<a href="https://researchsupport.admin.ox.ac.uk/contacts/ctrg">https://researchsupport.admin.ox.ac.uk/contacts/ctrg</a>
<b>Offering website</b>	<a href="https://researchsupport.admin.ox.ac.uk/ctrg">https://researchsupport.admin.ox.ac.uk/ctrg</a>
<b>Service language(s)</b>	English
<b>Target group</b>	Principal Investigators, Researchers, Students
<b>Type of the support</b>	Consulting, Training, Guidance (e.g. document templates, etc.)
<b>Short support description</b>	CTRG's role: <ul style="list-style-type: none"><li>• review and authorise sponsorship of clinical research when it involves: NHS patients, resources, staff, data or facilities, tissue samples, administration of a drug, herbal remedy or food product</li><li>• record, facilitate and oversee clinical research carried out within the University and ensure that the University is compliant with relevant legislation and regulations</li><li>• reduce and manage exposure to risk by advising on insurance</li><li>• work with University researchers and associated NHS trusts to enable high quality design and conduct of clinical research</li><li>• guide researchers in writing protocols, participant information sheets and informed consent forms prior to research ethics committee, MHRA and NHS approval submissions</li><li>• provide training and guidance in Good Clinical Practice and research governance</li><li>• monitor and/or audit clinical research sponsored/hosted by the University</li><li>• co-ordinate safety reporting and facilitate the University/OUH NHS Trust Trial Safety Group</li><li>• work with partner organisations to create and maintain a Joint Research Office with the aim of streamlining research processes</li></ul>

This profile is part of the [Comprehensive Inventory \(CI\)](#) provided by the [STARS project](#).

Purpose of the CI is to assist European academic drug developers in finding support on regulatory affairs. The inventory lists various support services provided by national competent authorities, public actors and private entities.

For information about the STARS project please visit [www.csa-stars.eu](http://www.csa-stars.eu) or contact [CoordinationSTARS@bfarm.de](mailto:CoordinationSTARS@bfarm.de).