

## PROVIDER PROFILE

<b>Name of the organisation</b>	European Forum for Good Clinical Practice
<b>Type of the organisation</b>	Non-profit
<b>Contact email(s)</b>	<a href="mailto:info@efgcp.eu">info@efgcp.eu</a>
<b>Offering website</b>	<a href="http://www.efgcp.eu">www.efgcp.eu</a>
<b>Service language(s)</b>	English
<b>Target group</b>	Principal Investigators, Researchers, Students, Companies
<b>Type of the support</b>	Consulting, Training, Guidance (e.g. document templates, etc.), Research on status of conditions relevant for potential regulatory changes, Ethical and regulatory aspects of clinical trials, Patient engagement in medicines development, Quality/audit/inspection infrastructure in clinical research.
<b>Short support description</b>	EFGCP has different tools to enable consultation: Working Parties on e.g. paediatric clinical development, quality or ethical aspects in clinical trials, etc. consisting of multi-stakeholder experts in the field, willing to jointly elaborate answers/solutions/positions on important topics; outcome-focused multi-stakeholder workshops, standard setting multi-stakeholder conferences on a particular topic; webinars with selected experts to discuss and find agreement on a clearly defined special issue; organization of multi-stakeholder Roadmap Initiatives to work out solutions to a complex topic like proposed content for new legislation of guidances; performance of research projects with clearly defined objectives, timelines and budgets

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).