15 YEARS OF EXCELLENCE IN RESEARCH

cliantha research

Regulatory Affairs Services

Service	Details
Clinical Trial Applications (CTAs), Amendments (CTA-As), Notifications (CTA-Ns)	Preparation and compilation of CTAs and related submissions for Phase I, II, III and BA/BE clinical trials conducted in Canada
Investigational Testing Authorization (ITA)	For unlicensed Class II, III, and IV Medical Devices to be used in a clinical trial in Canada
Medical Device Licence Applications	License to sell Class II, III, and IV Medical Devices in Canada
Cannabis Research License	Consultation on how to obtain a Cannabis Research license in Canada including compilation of submission on CTLS
Controlled Substance License	Consultation on how to prepare and compile required information in obtaining a Controlled Substance License from Health Canada
Investigational New Drug Applications (INDs)	For clinical trials that require USFDA approval
Pre-IND Meeting Request/Briefing Package	Preparation and compilation of Pre-IND meeting request and briefing package for FDA review

For business inquiry and more information ✉ info@cliantha.com