

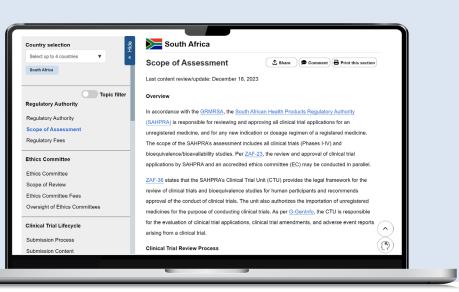
National Institute of Allergy and Infectious Diseases

Aggregating clinical research regulations from around the globe



What is ClinRegs and why you should use it

- Online database of countryspecific regulatory and ethics requirements
- 23 country profiles with links to official requirements and other sources
- · Easy to read and navigate



South Africa	Uganda	United States	Zimbabwe
Timeline of Review		1. Share	Comment
Last content review/update:	Last content review/update:	Last content review/update:	Last content review/update:
December 18, 2023	February 9, 2024	January 5, 2024	August 11, 2023
Overview	Overview	Overview	Overview
Based on ZAF-23 and the SA-	Per the NDPA-CTReg and UGA-	As delineated in 21CFR56 and	Per ZWE-GCP, parallel
GCPs, the review and approval	20, the National Drug Authority	21CFR312, institutional ethics	submissions are encouraged
of clinical trial applications by the	(NDA)'s review and approval of a	committee (EC) (institutional	among the regulatory authorities,
South African Health Products	clinical trial application are	review board (IRB) in the United	which comprise the Medicines
Regulatory Authority (SAHPRA)	dependent upon the applicant	States (US)) review of the clinical	Control Authority of Zimbabwe
and an accredited ethics	submitting proof in the	investigation may be conducted	(MCAZ), the National
committee (EC) may be	application of institutional ethics	in parallel with the Food & Drug	Biotechnology Authority of
conducted in parallel. The	committee (EC) (research ethics	Administration (FDA)'s review of	Zimbabwe (NBA), and the
applicant must notify each	committee (REC) in Uganda)	the investigational new drug	Research Council of Zimbabwe
regulatory body of the other's	approval and a research permit	application (IND). However, EC	(RCZ), and to the national ethics
approval once it has been	from the Uganda National	approval must be obtained prior	committee (EC)-the Medical

How you can make the most of your ClinRegs experience

- Compare up to 4 countries side-by-side
- View on-page notifications about recent changes to country requirements
- Sign up for email notifications about country profile updates
- Provide feedback on regulatory updates

