

				
	IRB-230	Posting of Clinical Trial Consent Forms	January 2019	1 of 1

1. PURPOSE

1.1. This policy establishes the posting of clinical trial consent forms to a federal website.

2. REVISIONS FROM PREVIOUS VERSION

2.1. None

3. POLICY

3.1. The revised Common Rule includes a requirement for the posting of one IRB-approved form to a federal website.