Swiss Biotech seeking several Clinical Research Scientists with different level of seniority (2 years to senior) for a 12 month contract in Switzerland. This is a long term assignment, with possibility for career progression on the long run, full-time and office-based in Switzerland (flexible to offer home working up to two days a week).

The programs will be in hemato-oncology.

REQUIREMENTS:

1. Review of clinical data within eCRF system, patient profiles, etc. to issue queries and clean data from clinical perspective.

2. Conduct review of ongoing summary data including: safety, primary efficacy variables, and laboratory data.

3. Assist the Clinical Research Physician to interface with project team members including: Clinical Operations, Data Management, Statistics, Drug Safety, Regulatory and Project Management.

4. Prepare or assist with preparation of key documents e.g. Investigator Brochures, regulatory submission documents, internal or external presentations, etc.

5. Protocol preparation (writing, reviewing, amending and cross-functional facilitation as appropriate).

6. Clinical study report preparation

7. Review literature and prepare summary documents for inclusion in IB, protocols regulatory submission documents, etc.

8. Participate and/or Lead team meetings as required.