PHARMACOVIGILANCE AND CLINICAL TRIALS DIVISION

RECOMMENDED FORMAT FOR CVs OF INDIVIDUALS PARTICIPATING IN CLINICAL TRIALS

1. Study Title:

2. Protocol Number:

3. Designation:
   [e.g., National Principal Investigator, Investigator (Principal, Co- or sub-), Study Coordinator, Regional Monitor, Local Monitor, Clinical Research Associate]

4. Personal Details
   Name:
   Work Address:
   Telephone Number:
   Fax Number:
   Cell-phone:
   Number:
   E-mail address:

5. Academic and Professional Qualifications

6. Professional registration number

7. Current personal medical malpractice insurance details

8. Relevant related work experience (brief) and current position

9. Participation in clinical trials research in the last three years
   [Study title, protocol number, designation. If multiple trials, only list those with relevance to this application, or in the last year]

10. Peer-reviewed publications in the past 3 years

11. Date of last GCP training
    [As a participant or presenter]

12. Any additional relevant information supporting abilities to participate in conducting this trial *[Briefly]*

Signature: Date: