Medical Research and Development Division

Research Compliance Incident Form

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| Maintaining integrity in research is crucial to advancing knowledge and ensuring public trust. This section will be dedicated to addressing research misconduct, violations, and incident reporting. Our goal is to foster a culture of accountability and transparency while safeguarding the rights of researchers, participants, and stakeholders in the Emirate of Abu Dhabi. Your voice matters and will be heard. If you want to report/raise a complaint on research integrity and research misconduct, you can fill in the Research Compliance Incident Reporting Form and sending it to [medical.research@doh.gov.ae](mailto:medical.research@doh.gov.ae) | | |
| Submitter Information | | |
| Full Name |  | |
| Facility: |  | |
| Professional Title: |  | |
| Official E-Mail address: |  | |
| Mobile No.  (+[country code][number], ex 00971-XX-XXXXXXX) |  | |
| Incident details | | |
| Observation found: |  | |
| Are there any supporting documents for the observation found? | Yes  No  If yes, list of documents to be provided. | |
| Study details for the incident reported | | |
| Study Site(s) |  | |
| Study site(s) IRB/REC reference number |  | |
| DOH ADHRTC approval reference number |  | |
| Study Title: |  | |
| Principle Investigator Name: |  | |
| Clinical Research Organization (CRO) involved in the study | Yes  No  If yes, name to be provided: --------------------------------------- | |
| Declaration and Signature | | |
| I hereby attest that all the information is accurate to the best of my knowledge. | | |
| Name | Signature | Date |
|  |  |  |