

Roles and Responsibilities of IEC Members
INSTITUTIONAL ETHICS COMMITTEE
HM PATEL CENTER FOR MEDICAL CARE AND EDUCATION, KARAMSAD

Role of member [s] of IEC

1. Chairperson:

- Conduct of meeting
- Ensure appropriate quorum
- Ensure active participation by all the members
- Appoint another member as Chairperson, in his/ her absence
- Approval of minutes of meeting
- Sign all documents on behalf of IEC
- Ensure that any conflict of interest is well taken care of

2. Deputy Chairperson:

- Manages role of Chairperson in his absence

3. Member Secretary:

- Manage administrative work of IEC
- Call for proposals
- Review and check submitted proposals for completeness
- Propose and circulate the agenda for meetings
- Ensure adequate number of clinical trials in a single meeting
- Review and decide on qualification of a proposal for exempt/ expedite review
- Review and reporting of Serious Adverse Event [s] [SAE]
- Review protocol deviations
- Communication with various stake holders [researchers, regulatory body etc.]
- Archiving of all IEC documents
- Monitor conduct of trials and their progress
- Prepare for audits
- Meetings and agenda for subcommittees
- Preparing minutes of meeting
- Review of Standard Operating Procedures [SOP]
- Preparation of annual reports
- Updating of new rules and regulations
- Arrange for capacity building among IEC members

4. Lay Person:

- Review of all the protocols from point of view of study participant
- Review of informed consent process
- Review of compensation processes
- Analysis of risks and benefits
- Review of post trial benefits
- Review of non trial related injuries
- Issues with vulnerability
- Monitoring of ongoing research projects

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5. Social Scientist:

- Review of all the protocols from point of view of social implications
- Review of informed consent process
- Review of compensation processes
- Analysis of risks and benefits
- Review of post trial benefits
- Review of non trial related injuries
- Issues with vulnerability
- Monitoring of ongoing research projects

6. Lawyer:

- Review of informed consent process
- Review of compensation processes
- Review of Clinical Trial Agreement [s]
- Compliance with current National Laws
- Analysis of risks and benefits
- Review of post trial benefits
- Issues with vulnerability
- Monitoring of ongoing research projects

7. Basic Scientist:

- Review of all the protocols for scientific merit and validity
- Review of informed consent process
- Review of compensation processes
- Analysis of risks and benefits
- Review of post trial benefits
- Review of non trial related injuries
- Issues with vulnerability
- Monitoring of ongoing research projects

8. Clinician:

- Review of all the protocols for scientific merit and validity
- Review of informed consent process
- Review of compensation processes
- Analysis of risks and benefits
- Review of post trial benefits
- Review of non trial related injuries
- Issues with vulnerability
- Monitoring of ongoing research projects

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9. Office Assistant/ Co-ordinator

- Maintaining an effective and efficient tracking procedure for each proposal received
- Preparation, maintenance and distribution of study files
- Allocation of project reviews to specific members to facilitate efficient dispensation of the projects
- Organizing IEC meetings
- Preparation and maintenance of meeting agenda and minutes
- Receive and check for the completeness of the documents for review by the EC
- Co-ordinate with the investigators
- Maintaining the IEC's documentation and archival
- Communicating with the IEC members and investigator applicants
- Arrangement of training for personnel and IEC members
- Organizing the preparation, review, revision and distribution of SOPs
- Work in unison with the EC members and the investigators to reduce the turn-around time of the study proposals sent to the EC for review
- Providing updates on relevant and contemporary issues related to ethics in health research, as well as relevant contemporary literature to the Committee members