1. Recommended for approval (without any stipulations).

2. Recommended for approval with suggestions for improvement
   (please specify here)

3. Recommended for fulfillment of stipulated conditions before consideration for
   approval
   (please specify here)

4. Recommended for rejection with specific grounds
   (please specify here)

---

THE RECOMMENDATIONS OF THE SUB-COMMITTEE ON
REHABILITATION OF ANIMALS AFTER EXPERIMENTATION SET UP
BY CPCSEA

(i) The cost of aftercare and/or rehabilitation of animals post-experimentation
are to be part of research costs and should be scaled per animal in positive
correlation with the level of sentence of the animals.

(ii) **The average costs of rehabilitation and aftercare of different species of
animals were thereafter worked out based on the actual average expenses
incurred by various institutions. The Sub Committee was of the view that the
following minimum amounts may be necessary for maintenance of animals
after experimentation (for feed and husbandry but not including
infrastructure and other overheads). However, this may be reviewed once in
two years, if required. The rehabilitation of small animals is not necessary.

<table>
<thead>
<tr>
<th>Species</th>
<th>Minimum cost per day per animal (In Rs.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dogs/Cats</td>
<td>20</td>
</tr>
<tr>
<td>Sheep/Goats</td>
<td>15</td>
</tr>
<tr>
<td>Cattle</td>
<td>25</td>
</tr>
<tr>
<td>Horse</td>
<td>25</td>
</tr>
<tr>
<td>Monkey</td>
<td>25</td>
</tr>
</tbody>
</table>

In the context of amended Rule 9 (cc) which provides for provision of a lump sum
amount as costs for rehabilitation and care of such animals to cover its entire
statistical expected life span, the Sub Committee discussions indicated that provision
of a lump sum amount, payable to an AWO, would not cover the possibility that the
animal may not actually live for its statistically expected lifespan. Accordingly
payments at intervals, as also annual confirmation regarding the well being of the
animal may be required.

Rule 9(cc) of the Breeding of and Experiments on Animals (Control and
Supervision) Amendment Rules 2006 also provides for the AWOs to be under the
control of the Committee. However, given the administrative structure of CPCSEA
and the fact that proper scrutiny and registration of AWOs is already being done by
AWBI, prior to release of grants, the Sub Committee was of the view that AWOs
should be shortlisted from amongst those AWOs already registered with AWBI, on
the basis of a well defined selection criteria.

(iii) Since the basic responsibility for rehabilitation of the animals after
experimentation was of the concerned establishment, Government may
contribute financial support for rehabilitation of animals by NGOs or other
institutions only under available animal welfare schemes.

(iv) Breeding of animals, post-experimentation should not be allowed.
(v) For animals recommended for euthanasia post-experimentation, the relevant criteria could be that:

(a) the animal is not able to perform its natural functions and left in pain & suffering.
(b) the animal has been exposed to contagious/infectious diseases of zoonotic importance.
(c) the animal has been exposed to radioactive experiments.

In other cases, animals would have to be rehabilitated by the concerned institute unless they are fit for transfer to NGOs or other institutions.

Thus the Sub Committee was of the view that the possibility of the same animal being diverted for further use in scientific experimentation by another institution, should also be considered, as this would avoid use of additional animals. This could be operationalised by a scientific method of identifying each animal, and by putting in place a mechanism whereby details of experimental animals available with institutions, could be made available to other institutions for examining their suitability for further use. The IAEC of the concerned institution will consider the issue as per existing norms and guidelines and forwarded its recommendation to the Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) for consideration.

(vi) A format for maintaining health records of experimental animals has been devised, which will facilitate transfer of animals among institutions as well as to NGOs (Annexure-I).

(vii) ICMR agreed to develop a web solution for facilitating exchange of details of animals available post-experimentation.

Regarding the ethical aspect of reuse of an animal, this is permitted in many countries, though sometimes with conditions eg not subjecting the animal more than once, to experiments involving severe pain, distress or suffering.

(viii) A checklist for selection of NGOs for rehabilitation of experimental animals was also prepared (Annexure-II). The agencies/NGOs and their Animal House Facilities must be registered with Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) and should satisfy the conditions laid down by Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) in this regard. The permanent identification marks eg tattooing must be done before rehabilitation of animals. Further, once an institute handed over an animal to an NGO, it would be the responsibility of the NGO to ensure the well being of the animals, with annual inspection by the concerned institutions. In case, it is necessary to transfer an animal out of the charge of the NGO, the concerned institute will be responsible for ensuring the continued well being of the animal.

(ix) Since the concerned institute is otherwise required to look after or maintain animals post-experimentation, they would reach an agreement directly with the concerned NGO regarding any reimbursement for handling the animals, apart from any assistance that NGO may obtain directly from AWBI/any other source. Modalities for transfer of experimental animals would be handled directly between concerned institute/NGO and no guidelines would be required to be formulated except stipulating that all existing animal laws would be followed.

(x) As regards the generic guidelines in respect of small animals, the mandate for examining proposals relating to small animals is with the IAECs.

** the animals which are presently out of experimentation and need instant rehabilitation prior to the notification of the Rehabilitation guidelines, will be provided a lump-sum amount by the establishments on the basis of types of animals, their health status and remaining lifespan of such animals, to the agencies. For the animals which are in experimentation and will be rehabilitated after their use by the establishment, the payment for maintenance of such animals will be made annually till the survival of the animals by the concerned organization, on the basis of demand made by the rehabilitating agency.
Experimental Animal Health Records

Instructions and guidelines

It is expected that all registered animal houses (for breeding or for conducting experiments) should adopt an institutional policy for animal welfare and implement the code of practice at different stages of animals under their custody. One of the most important aspects of this is the compilation and maintenance of accurate records relating to animal admissions, treatment, health, experimentation, and rehabilitation. Records pertaining to these aspects provide valuable information and facilitate inspection at all times.

The following forms will help in achieving the objectives in line with the policy.

1. Items that are marked (*) in the forms are optional. However, if information is available, it should be presented.
2. These forms are to be filled up only by designated persons and signed by appropriate authority.
3. As an archival policy of the institute, these forms should be available in the animal facilities for at least 10 years for inspection and data preservation.
4. Copies of the data pertaining to large animals at different phases need to be submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source, submitted to CPCSEA office mainly information on acquiring source.
5. As per the current CPCSEA regulations, permissions for experimentation in large animals are obtained from the central subcommittee with recommendation routed through IAEC of the institute and it is important to mention the reference number of the approval in these forms.
6. Date of initiation and date of termination mentioned in the form is as per IAEC approval.

Following forms are compulsory for rehabilitation

1. Pre Experimentation details of Animal for Transfer to Rehabilitation
   Form is for maintaining records of animal’s origin, treatment and tests performed during quarantine by the acquiring institution prior to experimentation signed by a veterinarian.
2. Experimentation details
   Form is for maintaining records pertaining to animal experimentation to know the kind of material handled during experimentation and protocols followed signed by principal investigator.
3. Post Experimentation Details of Animal & Suitability Certification
   Form is for maintaining records of animals after performance of experimentation for its continuation either for next experiment or for rehabilitation signed by veterinarian.
4. Transfer Certificate for Rehabilitation
   Form is for certifying suitability for rehabilitation authorized by a veterinarian.

Following forms should be maintained for supportive information

1. Tuberculin Testing
2. X-Ray
3. Body Weight
4. Parasitology
5. Microbiology
6. Treatment
7. Surgery
8. Research and Experimentation
9. Blood drawn
10. Hematology
11. Menstrual cycle

Forms such as Postmortem details and Histopathology are essential to be maintained for all animals that have been sacrificed in experiments and died during rehabilitation. In addition, institutions are required to maintain records pertaining to regular transfers within the institution for any experimentation or for any rehabilitation program.

A unique identification number is followed in all these forms for uniformity in every institution. The number to be placed in the appropriate box based on the following:

I  Species:
   01  Dog
   02  Monkey
   03  Cat
   04  Sheep
   05  Goat
   06  Cattle
   07  Horse
   08  Any other

II  Sex
   M  Male
   F  Female

III Year of Birth
   1997  97
   2001  01

IV Institutional Number Starts with CPCSEA
   Registration no.:
   0159
   00401

V  Individual number as per the institution
   002
   009
   019

Example: M 01 159 009
Pre Experimentation details of Animal for Transfer to Rehabilitation

1. Name of the organization currently holding the animal(s)

2. Details of the Animals
   a. Animal No:
   b. DOB
   c. Animals procured / Trapped from:
   d. Date of procurement / trapping
   e. Supplier / Source
   f. Colony breed / Wild caught
   g. Dam Number
   h. Sire Number
   i. Color
   j. Date of Induction
   k. Body Weight and Date

3. Physical Examination on arrival
   a. Fore Limbs
   b. Hind limbs
   c. Dental Formula: 1 C PM M
d. Teeth condition
   e. Eyes
   f. Nostrils
   g. Mouth lesions if any
   h. External Body Coat
   i. Abdominal palpation
   j. Lymph nodes
   k. Chest auscultation
   l. Body Temperature
   m. Any injuries physical deformities
   n. Clinical Symptoms if any
   o. Health condition

4. Quarantine period
   a. Introduced on
   b. Released from quarantine on

5. Microscopic examination for parasites
   a. Fecal Examination for Endo parasites
   b. Any Ectoparasites
   c. Parasites on Blood smear examination

6. Any Treatment given during Quarantine
   | Date | Clinical Symptoms | Diagnosis | Treatment | Recovered on |

7. Tuberculin Testing as per the animal:
   | Date | Source / Batch of Antigen | Result at each Observation |
   |      | First                     | Third                  | Second                  |

8. Chest X ray
   a. Date
   b. Report contents

9. Transferred to which Experiment

Signature
Name
Designation
### Experimentation Details

[Separate sheet for each experiment]

<table>
<thead>
<tr>
<th>1. Animal No</th>
<th>2. Body weight / Date</th>
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<table>
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<tr>
<th>3. Details of the Experiments:</th>
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<tbody>
<tr>
<td>a. Title/s of the project</td>
</tr>
<tr>
<td>b. Principal Investigator</td>
</tr>
<tr>
<td>c. Details of the protocol</td>
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<table>
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<tr>
<th>4. Age and weight range at the time of Initiation of the experiment</th>
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<tr>
<th>5. Experimental Endpoint Criteria</th>
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<tr>
<th>6. Date of Termination of experiment as per IACUC approval</th>
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<tr>
<th>7. Infectious agent used if any give details</th>
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<table>
<thead>
<tr>
<th>8. Radiotrace used if any give details</th>
</tr>
</thead>
</table>

**Signature of the Principal Investigator:**

Name  
Designation

---

### Post Experimentation Details of Animal & Suitability Certification

<table>
<thead>
<tr>
<th>1. Name of Organization currently possessing the animal</th>
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<tr>
<th>2. Animal No</th>
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<thead>
<tr>
<th>7. Body Weight &amp; Date</th>
<th>8. Details of Experiment / procedures:</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>9. Number of times anesthetized</th>
<th>10. Number of times blood withdrawn</th>
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<tr>
<th>11. Infectious organisms involved</th>
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</thead>
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<tr>
<th>12. Radioactive substance used</th>
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<tr>
<th>13. Is the animal suffering from any Zoonotic disease?</th>
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<tr>
<th>14. Physical abnormalities if any furnish details</th>
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<tr>
<th>15. Any clinical symptoms</th>
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<tr>
<th>16. Any attention of a veterinarian required during rehabilitation and how long</th>
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<th>17. Special nutritional requirements if any during rehabilitation</th>
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<tr>
<th>18. Present condition of the animal for rehabilitation. [Mark □]</th>
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</table>

<table>
<thead>
<tr>
<th>Excellent [Answers to 11-17 are No]</th>
<th>It can be Retained</th>
</tr>
</thead>
<tbody>
<tr>
<td>Good [Answer to 11-16 are No]</td>
<td>It can be given for Experimentation</td>
</tr>
<tr>
<td>Satisfactory [Answers to 11-15 are No]</td>
<td>Fit for transfer to NGO for Rehab</td>
</tr>
<tr>
<td>Poor [Answers 11-14 are Yes]</td>
<td>Can be sent for Euthanasia</td>
</tr>
</tbody>
</table>

**Signature of Veterinarian:**

Name  
Designation
**Transfer Certificate for Rehabilitation**

[To be filled in for every animal separately]

1. Animal No

2. Body weight / Date

3. Name & CPCSEA No. of Organization handing over the animal/s

4. Name & CPCSEA No. (if any) of the Organization taking over the animal/s

5. CPCSEA Registration No of the Rehabilitation unit:

6. In case of Emergencies Contact person/s details:
   - Name
   - Telephone No(s)
   - Fax:

7. Person handing over the animal
   - Signature
   - Name
   - Designation
   - Date

8. Person under taking over the animal
   - Signature
   - Name
   - Designation
   - Date

---

**Tuberculin Testing**

<table>
<thead>
<tr>
<th>Animal No</th>
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<table>
<thead>
<tr>
<th>Date</th>
<th>Source &amp; Batch number</th>
<th>Result of readings</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>First</td>
<td>Second</td>
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Signature of Veterinarian
- Name
- Designation

X-Ray

<table>
<thead>
<tr>
<th>Animal No</th>
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<table>
<thead>
<tr>
<th>Date</th>
<th>Organ / Site</th>
<th>Radiological Findings</th>
</tr>
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<tbody>
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</table>

Signature of Radiologist
- Name
- Designation
### Blood Drawn

<table>
<thead>
<tr>
<th>Animal No</th>
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<tbody>
<tr>
<td></td>
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</tbody>
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<table>
<thead>
<tr>
<th>Date</th>
<th>Quantity</th>
<th>If any anesthetic used &amp; Quantity</th>
<th>Purpose</th>
<th>Name of the Person drawn the blood</th>
</tr>
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Signature of Animal Technician
Name
Designation

### Microbiology

<table>
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<tr>
<th>Animal No</th>
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<table>
<thead>
<tr>
<th>Date</th>
<th>Test</th>
<th>Material</th>
<th>Organisms found</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
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</table>

Signature of Veterinarian/ Microbiologist
Name
Designation

### Body Weight

<table>
<thead>
<tr>
<th>Animal No</th>
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<tr>
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<table>
<thead>
<tr>
<th>Date</th>
<th>Weight</th>
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Signature of Animal Technician
Name
Designation

### Parasitology

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<th>Animal No</th>
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<table>
<thead>
<tr>
<th>Date</th>
<th>Ectoparasites</th>
<th>Endoparasites</th>
<th>Remarks</th>
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<tbody>
<tr>
<td></td>
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Signature of Veterinarian/ Parasitologist
Name
Designation
### Treatment

<table>
<thead>
<tr>
<th>Animal No</th>
<th>Date</th>
<th>Clinical Symptoms</th>
<th>Diagnosis</th>
<th>Treatment</th>
<th>Remarks</th>
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Signature of Veterinarian
Name
Designation

### Hematology

<table>
<thead>
<tr>
<th>Animal No</th>
<th>Date</th>
<th>RBC Millions/cc</th>
<th>WBC Cells/cc</th>
<th>PCV</th>
<th>HB g%</th>
<th>Differential Leukocyte Count</th>
<th>ESR</th>
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</table>

Signature of Pathologist / Veterinarian
Name
Designation

### Surgery

<table>
<thead>
<tr>
<th>Animal No</th>
<th>Date</th>
<th>Type of Surgery</th>
<th>Purpose</th>
<th>Treatment</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
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</tbody>
</table>

Signature of Veterinarian
Name
Designation
Post Mortem Report

1. Animal No

2. Source

3. Date of Induction

4. Was the animal Sacrificed / Died naturally

5. Condition of Carcass

6. Date of Death

7. History of Animal

8. External Examination
   a. Head
   b. Body
   c. Limbs

9. Discharges if any
   a. Mouth
   b. Eyes
   c. Ears
   d. Nose
   e. Anus
   f. Vagina/Penis
   g. Lesions in the Buccal Cavity
   h. Teeth
   i. Tongue

10. Internal Examination
    a. Lesions in Brain:
    b. Lesions in Sinuses:
    c. Heart:
    d. Liver:
    e. Spleen:
    f. Pancreas:
    g. Kidney/ Ureter:
    h. Stomach
    i. Intestine
    j. Cecum:
    k. Colon:
    l. Rectum
    m. Uterus
    n. Ovaries / Testes

11. Material Collected for Laboratory as deemed necessary by pathologist conducting PM
    Samples for Microbiology
    Samples for Parasitology
    Samples for Histopathology

12. Important Findings:

13. Cause of Death

Signature of Pathologist / Veterinarian
Name
Designation

Histopathology

1. Animal No

Date | Material | Findings
--- | --- | ---

Signature: Pathologist / Veterinarian
Name:
Designation:
Rehabilitation

Animal should be examined and assessed accurately by a person who is knowledgeable in particular requirements of that species by a veterinarian or by an experienced in rehabilitator process with primary objective of animal welfare.

The criteria may vary however following are a few conditions that preclude animal from rehabilitation and animals may require prompt euthanasia

1. Where the animal is found to be suffering from significant pain, distress, trauma or disease that cannot be relieved.

2. Where the animal would not survive without extended treatment or surgery, and is unlikely to recover sufficiently to return to the normal life.

Following conditions also preclude successful rehabilitation:

1. Loss of limbs or function of limbs
2. Permanent vital sensory loss (hearing, sight, smell, feeding)
3. Untreatable infectious disease
4. Permanent damage to the nervous system
5. Inability to adjust to temporary captivity
6. Chronic ill health
7. Abnormal behavioral patterns

---

Annexure –II

Check list for examining response of NGOs to be considered for rehabilitation of retired laboratory animals

1. Name of the Organisation:
2. Address:
3. Tel: Fax: Email:
4. Registration No. Status: Trust/ Society/ Any other:
5. Registration No. with Animal Welfare Board of India / CPCSEA (Enclose copy of registration Certificate)
6. Specify choice of animals for rehabilitation:
   a) Dogs   b) Goats/ sheep   c) Cattle   d) Monkeys   e) Others
7. Number of animals proposed to accommodated?
8. Details regarding location of shelter
   Where is the shelter located? Specify place relating climatic condition.
9. Details of availability of Veterinary care including surgical facilities available.
10. Details of amenities available at Shelter:

   (i) Food
   (ii) Water
   (iii) Bedding
   (iv) Safety/security
   (v) Hygiene / sanitation
11. Describe facilities for euthanasia, if required.

For office use

Name of observer(s):

Date of visit (DAY/MONTH/YEAR):

Start time: End time: Duration

Region: State:

Shelter name and address:

1. Whether all physical facilities match the particulars/ specifications given in the application.
2. Whether adequate veterinary / care available.
3. Remarks / comments.
4. Recommendation

Date: Signature:

Minimum space requirements for specific animals:

<table>
<thead>
<tr>
<th>Dogs/ Cats</th>
<th>Goat or Sheep</th>
<th>Cattle</th>
<th>Monkey</th>
<th>Horse</th>
</tr>
</thead>
<tbody>
<tr>
<td>8-14</td>
<td>10 (Calf)</td>
<td>24 (Full Grown)</td>
<td>8 (Minimum height 10 feet)</td>
<td>60</td>
</tr>
</tbody>
</table>

Minimum ratio of animals to attendants for specific animals:

<table>
<thead>
<tr>
<th>Dogs / Cats</th>
<th>Goat or Sheep</th>
<th>Cattle</th>
<th>Monkey</th>
<th>Horse</th>
</tr>
</thead>
<tbody>
<tr>
<td>10:2</td>
<td>15:1</td>
<td>10:1</td>
<td>6:1</td>
<td>4:1</td>
</tr>
</tbody>
</table>
ROLE OF NOMINEE

Nominee should be familiar with the CPCSEA guidelines, the concept of 3 Rs and other animal welfare issues. He/She must keep liaison with CPCSEA and also browse the MoEF website for any update of information or draft guidelines or rules and regulation, placed for comment. He/She should be familiar with animal experimentation procedures, husbandry conditions, supplies, humane way of dealing with animals, handling procedures etc and other requirements as published in the CPCSEA/INSA guidelines for GLP methods. He/she must learn/be familiar with alternatives, reduction and refinement procedure available in biomedical research studies/programmes.

Nominee attached to educational institution should be through with the syllabus prescribed by the University/college for a particular course. Nominee attached with research institutions should learn about the over all goal/mundate of the institute and keep himself/herself abreast with the current and past work done by the institute in animal studies and thus be able to link the new studies proposed. Nominees attached with institutions undertaking regulatory toxicology work should read and be familiar with various guidelines for registration of products such a pesticide, drugs and pharma, biotechnology and etc. For any clarifications on the protocols, methods, either they should see more details with the investigator when the agenda is circulated or seek comments from CPCSEA.

Nominee must read the proposal attached with Agenda thoroughly and come prepared to attend the IAEC meeting. He/She should aim for humane/ethical issues and compliance to GLP principles, avoid repetitions of animal studies, expected out come and it's impact, number and sex of animals used, source, health status, alternatives, reduction without loosing scientific conclusions etc.

Nominee should not indulge in arguments and heated debate but understand the merit of a study based on above principles.

Nominee is expected to sign the necessary forms of each protocol and maintain a copy/list in his/her records.

In case of proposals dealing with large animal, nominees should note that IAEC is only a recommending authority for such studies. Nominee should ascertain the capability of institution to perform studies on large animals and make sure his/her recommendations find place in the minutes attached to CPCSEA.

Nominees should forward a copy of the minutes to CPCSEA office within 15 days from the date of meeting/inspection of animal house.

If no meetings are held for 6 months in a row because of lack of projects or due to other reasons, nominee should notify to CPCSEA about "no meetings" after confirmation/verification with the organization.

Nominee should visit animal house at least once in a calendar year and submit the report in the prescribed form to CPCSEA office within a month from the date of inspection. Follow up may be taken up and reports also submitted to CPCSEA with ATR.

If nominee requires assistance of any other expert to review a protocol he/she can do so by consulting CPCSEA.

The nominee would be paid sitting fee and reimbursement of travel expenditure by the establishment / institute as decided by CPCSEA, from time to time.

Nominee not supposed to do

1. Nominee is not allowed to print visiting cards, letter pad with his/her name with Government of India official seal as IAEC nominee.

2. Nominee is not permitted to make any campaign/publicity about his/her role and solicit any sponsorship from any organization falling under his/her jurisdiction.

3. Nominee should not sign the minutes of the meeting/Forms/Register without attending the meeting in person.

4. Further, nominees should not allow any tele/audio conference with a non-participating member during the IAEC meeting.

5. Nominee should not allow any outside member to attend the IAEC meeting.

6. The nominees must keep themselves away from the media and press and will not disclose the confidential information related to the CPCSEA.