CONLUSIONS AND RECOMMENDATION
X SEMINAR ON HARMONIZATION OF REGISTER AND CONROL OF VETERINARY MEDICINES
BRASILIA, BRAZIL, 9-12 AUGUST 2004

1. HARMONIZATION

1.1. Technical Guidelines

Technical Guidelines were put into consideration of the attendees and due to the many observations made to, it was agreed to harmonize them during next Seminar. In addition, the need to define the contents of the technical guidelines was set forth.

1.2. Register Forms

1.2.1. Harmonization of Pharmacological Products Register Form was achieved

1.2.2. Harmonization of Biological Products Register Form was achieved.

Both forms are in the CD handed over to the attendees and also will be published in the Regional Representation Web page.

1.3. Labeling

SINDAN presented a proposal of labeling which was approved in general. Some comments on specific topics were left for further review. SINDAN document will be forwarded to all CAMEVET members for comments. Due date for presentation of comments is 30th November 2004. After end of analysis by CAMEVET members will be published in the Regional representation web page.

1.4. Norms identification

The document on identification and follow up of CAMEVET working documents, presented by Dr. Adela Encinosa Liñero, was approved.

2. CAMEVET

2.1. Executive Board

2.1.1. According to CAMEVET bylaws the Executive Board was constituted as follows:

President: Dr. Ricardo PAMPLONA (Brazil)
Vice-President: Dr. Elia Muñoz (Uruguay)
Official Member: Dr. Adela ENCINOSA LIÑERO (Cuba)
Adherent Members:
Dr. Milson Da Silva Pereyra (SINDAN)
Dr. Enrique Argento (CAPROVE)

The term for the elected members are valid from 12th August 2004 to date of next Seminar in 2005.

2.2. Secretary ex-office Report

2.2.1. The Secretary ex-office informed on the activities encouraged by CAMEVET

Dr. Silvia Pizarro from SAG (Chile) made a presentation regarding mission to the International Cooperation on the Harmonization of technical Requirements for the Registration of Veterinary Medicinal Products (VICH). This presentation was enhanced by members of VICH Steering Committee Members.

Dr. Elia Muñoz gave a detailed report concerning Good Manufacture Practice, on the Course hold in November 2004 in Uruguay. It was agreed on the need to hold similar courses in each OIE Country Members for the Americas in accordance to the countries needs on GMP.

Bearing in mind Paraguay proposal it was approved the organization of a GMP Course to be hold in Asuncion, Paraguay in November 2004. In addition, it was also agreed to hold a similar course in Brazil in year 2005.

3. VENUE OF NEXT CAMEVET'S MEETING

3.1. The motion to consider Uruguay as host country for next meeting was approved. In addition, the motion presented by Central-American Countries to organize CAMEVET's meeting in 2006 was approved.

4. WORKING PLAN AND FUTURE ACTIONS.

CAMEVET's working lines to follow in the future were approved. On this purpose CAMEVET'S Working Plan was written.

A proposal by Bolivia to include determination of MLRs for biological products in the agenda of next meetings, was approved. The proposal by Cuba to include medicinal food as topic for the Agenda of next meetings was also approved unanimously.

It was agreed on the need that CAMEVET's Executive Board shall inform to all the American countries responsible for the Veterinary Services putting stress on the important activities hold by CAMEVET as well as their achievements, thus they could request OIE to continue supporting CAMEVET in all aspects.