EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Veterinary Medicinal Products*
22 March 2019

CIRCABC Link: *https://circabc.europa.eu/w/browse/675c58a7-8429-498b-ba34-69043a7cc198*

AGENDA

Section A Information and/or discussion

A.01 Opening and adoption of the agenda. (CS)

A.02 HorStem – information about ongoing procedure for a marketing authorisation. (JP)

A.03 Implementation of the Regulation on veterinary medicinal products – update. (CS/AK)

A.04 Update from the International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH) meeting of 24/2 – 1/03/2019. (JP)

A.05 Update and information about the future inclusion of veterinary products in the scope of the Agreement on Mutual Recognition between the United States and the European Union for Good Manufacturing Practices (GMP). (LG)

A.06 Any Other Business. (CS)

Section B Draft(s) presented for an opinion

B.01 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision granting marketing authorisation under Regulation (EC) No 726/2004 of the European Parliament and of the Council for "Arti-Cell Forte - Chondrogenic induced equine allogeneic peripheral blood-derived mesenchymal stem cells", a veterinary medicinal product. (CS/JP)

(SANTE/3720784/2018)

Legal Basis: Regulation (EC) No 726/2004 - Art. 35

Procedure: Examination procedure