DRAFT AGENDA

22nd meeting of the Veterinary Pharmaceutical Committee
4 July 2016 (10:00-17:00)

Conference Centre Albert Borschette (CCAB)
room AB-1B
Rue Froissart 36
BE-1040 Brussels, Belgium

1. Opening and adoption of the agenda.

2. Maximum residue limits – discussion on draft implementing measures to the Regulation (EC) No 470/2009:
   a) draft Commission Implementing Regulation on the form and content of the applications and requests referred to in Articles 3 and 9 of the Regulation (EC) No 470/2009;
   b) draft Commission Regulation implementing Regulation (EC) No 470/2009 with regard to the rules on the use of a maximum residue limit established for a pharmacologically active substance in a particular foodstuff for another foodstuff derived from the same species, or a maximum residue limit established for a pharmacologically active substance in one or more species for other species;
   c) update on the planned Commission implementing measure regarding the methodological principles for the risk assessment and risk management recommendations referred to in Articles 6 and 7 of Regulation (EC) No 470/2009, including technical requirements in accordance with internationally agreed standards.

   a) draft Commission Regulation on the maximum residue limits to be considered for control purposes for foodstuffs derived from animals which have been treated under Article 11 of Directive 2001/82/EC of the European Parliament and of the Council;
b) draft Commission Regulation on reference points for action for non-allowed pharmacologically active substances present in food of animal origin.


5. Follow up to the discussion on diclofenac:

   a) Member State's update on the measures taken by the Member States at national level, including in particular information on reported vulture deaths in the Member States that have diclofenac authorised;

   b) discussion on the state of play and possible further steps.

6. Follow up to the discussion on gentamicin:

   a) outcome of the consultation with the Member States on the implementation into the national legislation of Article 6(3) of Directive 2001/82/EC;

   b) veterinary medicinal products for animals of the equidae family – situation in the Member States.


8. AOB