

A report to the Parliament and Council by the European Commission (**COM(97)362**).  
It is a critical assessment of the legal regime brought about by Directives 92/73/EEC and 92/74/EEC.

Commission proposed to take the following amendments into consideration:

- The adoption of a mutual recognition procedure under Article 6(1).
- The broadening of the scope of Article 7(1) to include routes of administration other than oral and external.
- The use of fantasy names for combination products and a more neutral wording on the labelling with regard to the assessment of effectiveness under Article 7(2).
- To make the provisions in Article 9(2) enabling Member States to adopt special rules on preclinical tests and clinical trials in accordance with the homeopathic traditions and principles in the Member States compulsory.

To expand the scope of Directive 92/74/EEC with respect to food producing animals.