Commission Directive 2005/34/EC of 17 May 2005 amending Council Directive 91/414/EEC to include etoxazole and tepraloxydim as active substances (Text with EEA relevance)

Article 3

1 Member States shall review the authorisation for each plant protection product containing etoxazole or tepraloxydim to ensure that the conditions relating to those active substances set out in Annex I to Directive 91/414/EEC are complied with. Where necessary, they shall amend or withdraw authorisations in accordance with Directive 91/414/EEC by 30 November 2005 at the latest.

2 For each authorised plant protection product containing etoxazole or tepraloxydim as either the only active substance or as one of several active substances all of which were listed in Annex I to Directive 91/414/EEC by 31 May 2005 at the latest, Member States shall re-evaluate the product in accordance with the uniform principles provided for in Annex VI to Directive 91/414/EEC, on the basis of a dossier satisfying the requirements of Annex III to that Directive. On the basis of that evaluation, they shall determine whether the product satisfies the conditions set out in Article 4(1)(b), (c), (d) and (e) of Directive 91/414/EEC.

Following that determination Member States shall:

a in the case of a product containing etoxazole or tepraloxydim as the only active substance, where necessary, amend or withdraw the authorisation by 30 November 2006 at the latest; or

b in the case of a product containing etoxazole or tepraloxydim as one of several active substances, where necessary, amend or withdraw the authorisation by 30 November 2006 or by the date fixed for such an amendment or withdrawal in the respective Directive or Directives which added the relevant substance or substances to Annex I to Directive 91/414/EEC, whichever is the latest.