Referral to the Coordination Group under Article 35 of Regulation (EU) No 528/2012

Executive summary

Type of referral: Referral to the Coordination Group of a disagreement on conclusions of the assessment report in accordance with Delegated Regulation (EU) No 492/2014 on the renewal of authorisations subject to mutual recognition in accordance with Regulation (EU) No 528/2012

Case type: Application of renewal of national authorisation (NA-RNL)

Reference Member State (rMS): Italy

Initiating concerned Member State (iCMS): France

Other Concerned Member States (CMSs): Austria, Cyprus, Germany, Estonia, Spain, Greece, Hungary, Ireland, Lithuania, Latvia, Malta, Czech republic, Poland, Portugal, Romania, Sweden, Slovenia, Slovakia,

Product type(s): 14

Active substance(s): Brodifacoum

Brief summary of the point of disagreement:

1) A composition change is indicated in the PAR. Clarification is needed on the evaluation of this change of composition.
2) A risk assessment must be performed for the water compartment for the sewer uses, and for groundwater for uses in and around buildings.

Outcome of the discussion within the Coordination Group (CG):

The CG members agreed by consensus on 7 March 2018 by written procedure:

1) The refMS confirmed that no changes in composition occurred since the first authorisation.
2) A groundwater risk assessment has been performed and is included in the PAR. Sewer use will be removed.

The product meets the condition for granting an authorisation in Article 19(1)(b)(iv) and Article 19(1)(c) of the BPR. This formal referral is therefore closed.