Annex 1

Notification Report
AUTHORISATION IN THE SCOPE OF ARTICLE 53

Please note that, due to a danger to plant health that cannot be contained by any other reasonable means, an authorisation in accordance with Article 53 of Council Regulation 1107/2009, has been granted for the preparation:

1	Member State, and MS notification number	*(The number shall be a sequence number per year and by MS, e.g. ES- 2011- 05.)*
2	Names of active substances	*e.g. thiametoxam*
3	Trade name of Plant Protection Product	
4	Pest	*(pest name EPPO, English and scientific)*
5	Crop	*(crop name EPPO, English, no group names)*
6	Time period for authorisation	
7	Technical contact point	*(Name, e-mail and telephone of the contact person responsible for the technical contents of the reasoning report.)*

8. Type of formulation and contents
(e.g 80% dry granule)

9. Applicant

10. GAP
(Attach GAP: dose rate, frequency, application type etc)

11. Compliance with existing MRLs

12. Value of tMRL if needed, *including information on the measures taken in order to confine the commodities resulting from the treated crop to the territory of the notifying MS. (PRIMO EFSA model to be attached)*

13. Function of the product
(e.g. systemic long acting insecticide; foliar fungicide, used for regular control, elimination scenario etc)

14. Type of danger to plant health.
(Provide reasoning for what category the 120 day authorisation is given: quarantine pest; emergent pest, either invading non-native, or native; emerging resistance in a pest, etc. Whereas reference to the EU quarantine legislation may suffice for quarantine pests elaborate reasoning should be provided for the category ' any harmful pest')

15. Size and effect of danger
(Describe shortly the area affected and the agronomic and economic effects it has)

16. Absence of any other reasonable means
(Describe the alternative control measures (chemical, non-chemical and cultural) and indicate why they do not (in combination) suffice.

17. Mitigation measures
(Describe what mitigation measures are taken to minimise the risk to man, animals, and the environment, attach summary risk attachment)

18. Applications in progress
(The use notified may have been applied for already, or suitable alternative PPP may be in the process of authorisation. Describe such applications, including a possible date of authorisation)

19. Research activities
(Describe the research efforts undertaken and/or in progress, their aims, their funding, and their expected date of results. This is needed for all categories of dangers, except quarantine pests that can still be eliminated, or infrequent pests.
In case of a repeated notification: indicate the state of works of the research projects.)

to be uploaded on CIRCA[1]

[1] IG "Plant Protection Products and their residues, section of the next SCFCAH-Meeting, subsection "Notifications under Art 53