|  |  |  |
| --- | --- | --- |
|  |  | |
|  |  | |
| **G/SPS/N/EU/51/Add.1** | |
| 5 November 2014 | |
| (14-6442) | | Page: 1/2 |
| **Committee on Sanitary and Phytosanitary Measures** | | Original: English |

NOTIFICATION

Addendum

The following communication, received on 3 November 2014, is being circulated at the request of the Delegation of the European Union.

**\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

|  |
| --- |
| MRLs for benthiavalicarb, cyazofamid, cyhalofop-butyl, forchlorfenuron, pymetrozine and silthiofam |
| The proposal notified in G/SPS/N/EU/51 (30 August 2013) has been adopted as "Commission Regulation (EU) No. 398/2014 of 22 April 2014 amending Annexes II and III to Regulation (EC) No. 396/2005 of the European Parliament and of the Council as regards maximum residue levels for benthiavalicarb, cyazofamid, cyhalofop-butyl, forchlorfenuron, pymetrozine and silthiofam in or on certain products (Text with EEA relevance)" [OJ L 119, 23 April 2014, pp. 3-39]. The Regulation entered into force on 13 May 2014. All MRLs are applicable from 13 November 2014.  <http://members.wto.org/crnattachments/2014/sps/EEC/14_4933_00_e.pdf>  <http://members.wto.org/crnattachments/2014/sps/EEC/14_4933_00_f.pdf>  <http://members.wto.org/crnattachments/2014/sps/EEC/14_4933_00_s.pdf> |
| **This addendum concerns a:** |
| [] Modification of final date for comments |
| [**X**] Notification of adoption, publication or entry into force of regulation |
| [] Modification of content and/or scope of previously notified draft regulation |
| [] Withdrawal of proposed regulation |
| [] Change in proposed date of adoption, publication or date of entry into force |
| [] Other: |
| **Comment period: *(If the addendum extends the scope of the previously notified measure in terms of products and/or potentially affected Members, a new deadline for receipt of comments should be provided, normally of at least 60 calendar days. Under other circumstances, such as extension of originally announced final date for comments, the comment period provided in the addendum may vary.)*** |
| [] Sixty days from the date of circulation of the addendum to the notification and/or *(dd/mm/yy)*: Not applicable. |
| **Agency or authority designated to handle comments: [****X] National Notification Authority, [****X] National Enquiry Point. Address, fax number and e-mail address (if available) of other body:** |
| European Commission  DG Health and Consumers, Unit G6-Multilateral International Relations  Rue Froissart 101, B-1049 Brussels  Tel: +(32 2) 295 4263  Fax: +(32 2) 299 8090  E-mail: sps@ec.europa.eu |
| **Text(s) available from: [****X] National Notification Authority, [****X] National Enquiry Point. Address, fax number and e-mail address (if available) of other body:** |
| European Commission  DG Health and Consumers, Unit G6-Multilateral International Relations  Rue Froissart 101, B-1049 Brussels  Tel: +(32 2) 295 4263  Fax: +(32 2) 299 8090  E-mail: sps@ec.europa.eu |

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| |  |  | | --- | --- | | 世界贸易组织 | **G/SPS/N/EU/51/Add.1**  **分发日期：**2014-11-05  (14-6442) | | 卫生及植物卫生措施委员会 | 原文: 英文 |     通 报    补遗    **应欧盟代表团的要求，发送2014-11-03如下信息：**  苯噻菌胺(Benthiavalicarb)、氰霜唑（cyazofamid）、氰氟草酯（cyhalofop-butyl）、氯吡脲(Forchlorfenuron)、吡蚜酮(Pymetrozine)及硫硅菌胺(Silthiofam)最大残留限量  在G/SPS/N/EU/51(2013年8月30日)中通报的提案已批准为“2014年4月22日委员会第(EU)398/2014号法规——修改欧洲议会及理事会有关某些产品内/表苯噻菌胺(Benthiavalicarb)、氰霜唑（cyazofamid）、氰氟草酯（cyhalofop-butyl）、氯吡脲(Forchlorfenuron)、吡蚜酮(Pymetrozine)及硫硅菌胺(Silthiofam)最大残留限量的第(EC)396/2005号法规附件II和III(欧洲经济区相关文本)"[OJL119，2014年4月23日,3-39页]。本法规于2014年5月13日生效。所有最大残留限量将2014年11月13日起适用。http://members.wto.org/crnattachments/2014/sps/EEC/14\_4933\_00\_e.pdfhttp://members.wto.org/crnattachments/2014/sps/EEC/14\_4933\_00\_f.pdfhttp://members.wto.org/crnattachments/2014/sps/EEC/14\_4933\_00\_s.pdf  **该补遗通报涉及**:  [ ] 意见反馈截止日期的修订  [ X ] 法规批准、生效、公布的通报  [ ] 以前通报的法规草案的内容及/或范围的修改  [ ] 撤消拟定法规  [ ] 更改拟定批准日期, 公布或生效日期  [ ] 其它:  **评议期：***(如补遗通知增加了以前通报措施涉及的产品及/或可能受影响的成员范围，则应提供一个新的接收评议截止日期，通常至少为60天*。*其它情况，如延长原定的最终评议期，则可以更改补遗通报内的评议期。)*  [  **]补遗通报发布日后60天及或(*年/月/日)*：**不适用  **负责处理反馈意见的机构: [ X ]国家通报机构，[X ]国家咨询点，或其他机构的联系地址、传真及电子邮件地址(如能提供):** EuropeanCommissionDGHealthandConsumers,UnitG6-MultilateralInternationalRelationsRueFroissart101,B-1049BrusselsTel:+(322)2954263Fax:+(322)2998090E-mail:sps@ec.europa.eu  **文本可从以下机构得到: [X ]国家通报机构，[X ]国家咨询点，或其它机构的联系地址、传真及电子邮件地址(如能提供):**  EuropeanCommissionDGHealthandConsumers,UnitG6-MultilateralInternationalRelationsRueFroissart101,B-1049BrusselsTel:+(322)2954263Fax:+(322)2998090E-mail:sps@ec.europa.eu |