

## **Section 12 Notice**

## Additional Information Required to Fulfill a Condition of Registration

**Product Name:** Mosquiron 0.12P

Registration Number: 31078
Application Number: 2011-1681
PMRA #: 2970207

Date of Issuance: March 11, 2019

The information specified below is required to be submitted to the Pest Management Regulatory Agency in accordance with section 12 of the *Pest Control Products Act* by **October 15, 2019**.

PART 0 INDEX

**DACO:** 0 Index

Required Data: Please submit an electronic index of the data package submitted in

response to this letter. Please refer to Regulatory Directive 2006-05, Requirements for Submitting Data Index, Documents and Forms, for

additional information.

PART 3 CHEMISTRY REQUIREMENTS FOR THE REGISTRATION OF

MANUFACTURING CONCENTRATES AND END-USE

PRODUCTS FORMULATED FROM REGISTERED TECHNICAL

GRADE OF ACTIVE INGREDIENTS OR INTEGRATED

SYSTEM PRODUCTS

**DACO:** 3.5.10/3.5.14

**Title:** Storage stability data/ Corrosion characteristics

**Deficiency:** A storage stability study, where corrosion characteristics are confirmed,

was not submitted

Required Data: A storage stability study, where corrosion characteristics are

confirmed, must be conducted under at least one of the following

regimes:

at least one year's duration at a constant ambient temperature of 20 or 25°C and, if the package is permeable, at a relative humidity of



50%, with quantitative analysis for the active ingredient(s) at study commencement and following storage periods of 3, 6 and 12 months; or

at least one year's duration under warehouse conditions that reflect the expected storage conditions of the commercial product (this may include the need for freeze-thaw studies). Where possible, the storage environment should approximate any extremes of temperature or climate expected to occur under actual storage conditions. Quantitative analysis for the active ingredient(s) is required at study commencement and following storage periods of 3, 6 and 12 months; or

of 14 days' duration under accelerated conditions at a constant temperature of 54°C, with quantitative analysis for the active ingredient(s) at study commencement and after 14 days.