



LICENCE TO MANUFACTURE VETERINARY CHEMICAL PRODUCTS

Licence Holder: Bioproton Pty. Ltd.
ACN 059 093 417

Licence No: 4096

The APVMA hereby issues a licence under section 123 of the Agricultural and Veterinary Chemicals Code (Agvet Code) to the above named person (the Licence Holder) to carry out the following step(s) of manufacture:

Quality assurance (QA) of raw materials, formulation including blending, filling, granulation, fermentation (bacterial and fungal), packaging, labelling, analysis and testing (physical and enzyme assay), storage, and release for supply.

This licence authorises the manufacture of the following type(s) of veterinary chemical products only:

Category 4 (Premixes and supplements) - Enzyme supplements.


—at the following premises:

**55 Dulacca Street
ACACIA RIDGE QLD 4110**

This licence is subject to the conditions set out in subsection 126(4) of the Agvet Code, regulations 60, 61 and 62 of the *Agricultural and Veterinary Chemicals Code Regulations 1995* (Agvet Code Regulations) and the **additional conditions in the attached Schedule.**

This licence comes into force on the date of issue and replaces the previous licence issued on 31 May 2016. This licence remains in force unless otherwise suspended or cancelled by the APVMA.

Dated this ^{29th} day of March 2017


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Susan Hanns
Acting Director, Manufacturing Quality and
Licensing
Delegate of the Australian Pesticides and
Veterinary Medicines Authority

This licence remains the property of the APVMA and must be returned on request

SCHEDULE OF ADDITIONAL LICENCE CONDITIONS


The following additional conditions apply to and form part of Licence No. **4096** issued to the Licence Holder:

Bioproton Pty. Ltd.

ACN 059 093 417

- S1.1 This Licence authorises only those steps of manufacture, product type(s) and premises listed.
- S1.2 The Licence Holder must provide the original signed copy of the audit report together with details of all corrective actions they propose to make with respect to the identified non-conformances and the timeframe for their implementation. This documentation must be received by the APVMA within 25 working days of the audit in accordance with Regulation 61(8C)(a)(i).

Dated this ^{29th} day of March 2017


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Susan Hanns
Acting Director, Manufacturing Quality and
Licensing
Delegate of the Australian Pesticides and
Veterinary Medicines Authority

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