



9th April 2013

Jeanette Drysdale
AR & JA Drysdale
P O Box 72 275
Papakura 2244

Dear Jeanette,

SQUEAK SUPER – V003416

C2 - The requested changes in manufacturer of the formulated product are:

- Removal of ChemSafe Group Ltd, 8 Alpito Place. Pukekohe; and
- Addition of ChemSafe Group Ltd, 12 Cape Hill Road, Pukekohe

Registration Renewal

The Agricultural Compounds and Veterinary Medicines Group has completed the technical appraisal and risk assessment of your smart track application requesting a variation to your registration of this product.

Authorisation has been granted under delegated authority from the Director General, Ministry for Primary Industries.

Please find enclosed the revised registration. It consists of the revised Certificate of Registration and Product Data Sheet. These along with the currently approved label content form the current registration of this product.

Note that the conditions of registration have changed.

Please read your certificate carefully. No changes to your product label are required.

The registration of this product will expire in three years time.

An invoice for payment is attached.

Yours Sincerely,

A handwritten signature in black ink, appearing to read 'N. Sanders'.

Natalie Sanders
Adviser (Approvals Operations)



Certificate of Registration

This Certificate of Registration is issued to:

Rentokil Initial New Zealand Ltd

of **Level 1, 89 Carbine Road
Mt Wellington, AUCKLAND**

The vetebrate toxic agent known by the trade name of:

SQUEAK SUPER

No. **V003416**

Is hereby registered under the Agricultural Compounds and Veterinary Medicines Act 1997

This registration expires on the 9th day of April 2016

The conditions placed on this approval are attached.



Dated on this 9th day of April 2013 (Date of first registration was 12 October 1984)

Marco Zinzley
Manager (Approvals Operations)

Signed:


Agricultural Compounds and Veterinary Medicines Group
Acting under delegated authority

This certificate is a record of the product approval that was current on the date the certificate was issued. It reflects the public register entry for the product at that time. From time to time the wording of registration conditions may change, resulting in a change on the public register for that product. As a certificate is not necessarily re-issued for such changes, consult the public register for the current approval of the product




Maree Zinley
Manager (Approvals Operations)



CONDITIONS

- 2 The product must be manufactured in accordance with the ACVM Standard for Good Manufacturing Practice and to the chemistry and manufacturing specifications provided by the registrant and approved as part of the registration.

- 4 The product must only be sold or imported according to the current registration.

- 31 This product must only be used as specified in the label content.

- 37 Ongoing obligations:
The registrant must provide an annual summary of adverse events to the ACVM Group. Adverse events which have serious implications for the continued use of the product must be notified immediately. The registrant must also advise the ACVM Group of any new studies or data that contradict information previously supplied.

- 51 Vertebrate Toxic Agents: In addition to any labelling, advertising or promotion requirements specified in the current registration, labelling, advertising or promotion of the product must comply with the current ACVM - New Zealand Labelling and Advertising Guide for Vertebrate Toxic Agents Requiring Registration

- 55 For pack sizes greater than 3kg, the product must be sold only by a person who has been approved by the ACVM Group

- 56 For pack sizes greater than 3kg, a register of sales must be kept (minimum of 3 years), recording who the product was sold to and the quantity sold
