6th June 2014

Jeanette Drysdale
Rentokil Initial Limited
PO Box 72 275
Papkura 2244

Dear Jeanette

V2964 – Rid Rat Super
Registration Renewal

The Agricultural Compounds and Veterinary Medicines Group is in receipt of your letter dated 27th March 20014 confirming no changes in the product and manufacturing specification of the above named product.

The Certificate of Registration, approved Registration and Product Datasheet (PDS) and Label Content are enclosed. These documents constitute the registration for this product.

Please read your certificate carefully.

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

Please find invoice attached.

Condition 108 has been applied to the registration, which means that the shelf life statement on the label can be removed or amended if you are satisfied that the shelf-life can be managed in an alternative way.

Yours sincerely

Jo-ellen Powell
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:

Rid Rat Super No. **V002964**

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

This registration expires on the 26th day of May 2017

The conditions placed on this approval are attached.

MINISTRY FOR PRIMARY INDUSTRIES
NEW ZEALAND

Dated on this 6th day of June 2014 (Date of first registration was 16 July 1981)

Signed: Maree Zinzley
Manager Approvals Operations
Regulation and Assurance

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 Zinzley
Manager Approvals Operations
Regulation and Assurance

MINISTRY FOR PRIMARY INDUSTRIES
NEW ZEALAND

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