20th September 2012

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

Dear Jeanette.

RENTOKIL RID RAT PREMIUM BLOCK – V009581
B2 – New Registration

The Agricultural Compounds and Veterinary Medicines (ACVM) Group are in receipt of the revised Registration and Product Datasheet and label content requested in our letter dated 13th August 2012

The Certificate of Registration, Registration and Product Datasheet and approved label content are enclosed. These constitute the current approval for the above named product.

Note that the registration of this product will expire in three years' time

Import Approval
This product may be redirected to the following address for repacking/relabeling:
Chemsafe Group Ltd, 8 Alpito Place, Pukekohe.
Please present this letter at the border to facilitate product clearance.

The invoice for payment is attached

Yours Sincerely,

Natalie Fagan
Adviser (Approvals, Operations)

BIOSECURITY NEW ZEALAND

NEW ZEALAND FOOD SAFETY AUTHORITY

Te Pou Oranga Kai o Aotearoa

Certificate of Registration

This Certificate of Registration is issued to:

Liphatech S.A.S

of **Bonnel**
BP3 - 47480, PONT DU CASSE, France

The vetebrate toxic agent known by the trade name of:

RENTOKIL RID RAT PREMIUM BLOCK No. **V009581**

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

This registration expires on the 20th day of September 2015

The conditions placed on this approval are attached, together with the registration and product datasheet and approved label content.

NEW ZEALAND FOOD SAFETY AUTHORITY
AGRICULTURAL COMPOUNDS & VETERINARY MEDICINES

Dated on this 20th day of September 2012

Glen Bradbury
Manager ACVM Appraisals and Programmes

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.

NEW ZEALAND FOOD SAFETY AUTHORITY
Te Pou Oranga Kai o Aotearoa

Glen Bradbury
Manager, ACVM Approvals and Programmes

21/2/12

NEW ZEALAND FOOD SAFETY AUTHORITY
AGRICULTURAL COMPOUNDS & VETERINARY MEDICINES

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.

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