Serial Number **1947/13**

NEW ZEALAND

THE STOCK-REMEDIES REGULATIONS 1947

B. C. FREYBERG, Governor-General

ORDER IN COUNCIL

At the Government House at Wellington, this 17th day of September, 1947

Present:

HIS EXCELLENCY THE GOVERNOR-GENERAL IN COUNCIL

PURSUANT to the Stock-remedies Act, 1934, His Excellency the Governor-General, acting by and with the advice and consent of the Executive Council, doth hereby make the following regulations.

REGULATIONS

PRELIMINARY

1. These regulations may be cited as the Stock-remedies Regulations 1947.

2. These regulations shall come into force on the seventh day following notification in the *Gazette* of the making thereof.

3. In these regulations, unless inconsistent with the context,—

"The said Act" means the Stock-remedies Act, 1934:

"Board" means the Stock-remedies Registration Board constituted under the said Act:

"Label" includes any brand or writing on any stock-remedy or on any receptacle containing any stock-remedy or on any carton or cover for any such receptacle:

"Proprietor", in respect of a stock-remedy manufactured in New Zealand, means the manufacturer thereof, and in respect of a stock-remedy not manufactured in New Zealand, means the importer thereof:

"Stock-remedy" means any substance (including vaccines, sera, and other biological products) manufactured, advertised, or sold as a remedy for general use for the cure or prevention of disease in stock, or for the destruction or prevention of parasites of stock, or for the maintenance or improvement of the health or condition of stock, but does not include any substance which is used primarily as a food for stock:

"Vendor" means any person who, either on his own account or on behalf of any other person, sells in the ordinary course of his business any stock-remedy.

4. The Stock-remedies Registration Regulations 1935 and the Stock-remedies Registration Regulations 1935, Amendment No. 1, are revoked.

5. All applications, approvals, certificates, declarations, registers, registrations, and generally all acts of authority, and all other documents, matters, acts, and things, and all periods of time which orginated or had effect under the regulations hereby revoked and are of continuing effect at the time of coming into force of these regulations, shall enure for the purposes of these regulations as fully and effectually as if they had originated under these regulations and shall, where necessary, be deemed to have so originated.

Register of Stock-remedies

6. (1) For the purposes of the said Act a Register of Stock-remedies shall be kept by the Registrar of the Board at his office in Wellington.

(2) The register may be kept book-wise or in card form or in such other manner as may be deemed convenient.

(3) The entries in the register may be arranged in order of names of stock-remedies arranged alphabetically and in sub-order of names of proprietors arranged in like manner or in such other order as may be deemed convenient, and may from time to time be rearranged in different order.

(4) The particulars to be entered in the register shall be those set out in the form No. 1 in the Schedule hereto.

(5) No entry in the register shall be made by the Registrar without the direction in writing of the Board, and all entries shall be made in the order in which the directions to make the same are received by the Registrar.

(6) The direction in writing of the Board referred to in the last preceding subclause shall be sufficient if signed by the Chairman and given pursuant to a resolution of the Board.

Application for Registration

7. (1) Every application for the registration of a stock-remedy pursuant to section 6 (3) of the said Act (accompanied by the appropriate fee) shall be in the form No. 2 in the Schedule hereto.

(2) The specimen copy of every label or advertisement to be used or published in respect of the stock-remedy which by paragraph (*e*) of subsection (3) of section 6 of the said Act must accompany the application for registration shall be in duplicate.

(3) The statutory declaration of the applicant pursuant to subsection (4) of section 6 of the said Act in verification of the particulars in respect of the stock-remedy set out in his application for registration shall be in the form No. 3 in the Schedule hereto.

(4) Where the applicant is a company or other body corporate the declaration referred to in the last preceding clause of this regulation shall be made by the managing director, manager, secretary, or other principal executive officer thereof, and where the applicant is a firm or partnership such declaration shall be made by any member thereof acting for and on behalf of such firm or partnership.

Evidence of Registration

8. (1) On the registration of any stock-remedy there shall be issued to the proprietor of the stock-remedy a certificate of registration in the form No. 4 in the Schedule hereto.

(2) Every such certificate of registration shall be *prima facie* evidence of the registration of the stock-remedy to which the same relates.

APPROVAL OF LABELS OR ADVERTISEMENTS

9. (1) Every application by the proprietor of any stock-remedy for approval of any label or advertisement relating to the stock-remedy shall, when not combined with his application for the registration of the stock-remedy, be in the form No. 5 in the Schedule hereto, and every such application shall be accompanied by two copies of the label or advertisement.

(2) The approval by the Board of any label or advertisement relating to any stock-remedy shall be evidenced by endorsing on each of two copies of such label or advertisement a certificate in the following form, that is to say,—

" Approved for the period of registration ending the 30th day of September, 19..."

Signature :.....

Registrar, Stock-remedies Registration Board.

Date :.....

(3) One copy of such label or advertisement duly endorsed as provided by the last preceding clause of this regulation shall be forwarded to the proprietor of the stock-remedy, and the other copy shall be retained and filed in the office of the Registrar of the Board.

STANDARDS FOR STOCK-REMEDIES

Stock-licks

10. (1) Every stock-remedy in the form of a stock-lick shall contain only compounds of phosphorus, iodine, cobalt, or copper incorporated with common salt and such diluents and flavouring substances as may be necessary to render the stock-lick palatable and convenient for administration to stock.

(2) Every phosphatic stock-lick shall contain not less than 10 per centum by weight of phosphoric anhydride (P_2O_5).

(3) Every iodized stock-lick shall contain not less than four nor more than five ounces of potassium iodide per ton if incorporated in the stock-lick in a stabilized form, or not less than eight nor more than ten ounces of potassium iodide per ton if incorporated in the stock-lick otherwise than in a stabilized form.

(4) Every cobaltized stock-lick shall contain not less than 0·0023 per centum by weight of cobalt, or four ounces of cobalt sulphate ($CoSO_4$, 7 H_2O) per ton.

(5) Subject to the next succeeding clause of this regulation, every copperized stock-lick shall—

(*a*) If intended for sale for administration to sheep, contain not less than 0·064 nor more than 0·13 per centum by weight of copper, or 0·25 to 0·50 per centum of copper sulphate ($CuSO_4$, 5 H_2O) :

(*b*) If intended for sale for administration to cattle, contain not less than 0·38 nor more than 0·50 per centum by weight of copper, or 1·50 to 2·00 per centum of copper sulphate ($CuSO_4$, 5 H_2O).

(6) Where any copperized stock-lick contains substances other than common salt designed or intended to increase its palatability and consumption, the quantity of copper contained in such lick shall—

(*a*) If intended for sale for administration to sheep, be not less than 0·048 nor more than 0·064 per centum by weight of copper, or 0·19 to 0·25 per centum of copper sulphate ($Cu\ SO_4$, $5\ H_2O$) :

(*b*) If intended for sale for administration to cattle, be not less than 0·25 nor more than 0·38 per centum by weight of copper or 1·0 to 1·50 per centum of copper sulphate ($CuSO_4$, $5\ H_2O$).

Vitamin Preparations

11. (1) No stock-remedy shall contain any vitamins other than vitamin A and vitamin D, and such vitamins shall be prescribed only in the form of vitamin oils.

(2) Every vitamin oil intended for sale for administration to poultry shall contain not less than 1,000 international units of vitamin A and not less than 100 international chick units of vitamin D per gramme. For the purposes of this clause " chick unit " means the measure of the antirachitic activity in respect of poultry of 1 gramme of a solution of 0·000025 milligrams pure crystalline vitamin D_3 in a vitamin D free olive oil.

(3) Every vitamin oil intended for sale for administration to stock other than poultry shall contain not less than 1,000 international units of vitamin A and 100 international units of vitamin D per gramme.

Worm Drenches

12. No worm drench shall contain any drug other than phenothiazine, bluestone, nicotine sulphate, carbon tetrachloride, tetrachlorethylene, oil of chenopodium, carbon disulphide, arecoline hydrobromide or any derivative thereof, sodium arsenite, kamala, and hexylresorcinol. Every worm drench shall be so compounded as to contain adequate and safe amounts of the active ingredients of the drench.

Teat Salves

13. Every preparation described by the proprietor as a teat salve or intended for sale for the treatment of sore teats of cows shall contain a recognized antiseptic in effective concentration. Such preparation shall not have a paraffin or any other base which will remain on the teat for more than twelve hours after application, and shall not contain any substance capable of tainting milk.

Mastitis Remedies

14. No preparation in the form of a liniment intended for sale for the treatment of mastitis shall contain any substance capable of tainting milk.

15. Every drench intended for sale for the treatment of streptococcal mastitis shall contain a sulphonamide.

16. Every preparation intended for sale for the treatment of streptococcal mastitis by intra-mammary injection shall contain an acridine dye, iodine, silver oxide, a sulphonamide, tyrothricin, gramicidin, or penicillin. Every such preparation shall be so compounded that the drugs are present in an active form and in effective concentration.

Grass Staggers

17. Every preparation intended for sale for the treatment of grass staggers by injection shall be so compounded as to contain magnesium sulphate in effective concentration.

Milk Fever

18. Every preparation intended for sale for the treatment of milk fever by injection shall be so compounded as to contain calcium borogluconate in effective concentration.

Rotenone Sheep-dip

19. For the purpose of the preparation of a sheep-dip incorporating rotenone as the effective parasiticide, where the rotenone is derived from the roots of derris, cube, timbo, barbasco, or tuba, or other similar vegetable substances, these shall contain not less than 4·5 per centum by weight of rotenone. Such roots or other vegetable substances shall be so ground that not less than 50 per centum by weight is capable of being passed through a No. 300 British standard wire-mesh sieve and the remainder through a No. 200 British standard wire-mesh sieve.

REGISTRATION OF NEW PROPRIETOR

20. Where during the period of registration of any stock-remedy the proprietor thereof ceases to be such proprietor, the Board may, on the application of any person who satisfies the Board that he has become the proprietor of the stock-remedy, cause the name and address of the new proprietor to be entered in the register in the place of the name and address of the former proprietor. Every such application shall be accompanied by the appropriate fee as hereinafter prescribed.

SAMPLING

21. (1) Where for the purposes of the said Act a sample of any stock-remedy is taken by an Inspector, the size, weight, or volume of the sample so taken shall, unless otherwise directed by the Board, be not less than the minimum size, weight, or volume of the stock-remedy offered for sale.

(2) Each of the parts into which the sample is to be divided under section 12 of the said Act shall be placed into a clean, dry receptable on which shall be affixed a label in the form No. 6 in the Schedule hereto.

(3) The label shall contain the particulars in the said form No. 6 and shall be signed by the Inspector and by the proprietor or vendor or other witness present at the taking of the sample.

FORM OF CERTIFICATE OF ANALYST

22. (1) For the purposes of section 13 of the said Act the certificate of the Analyst in respect of any stock-remedy, not being a vaccine, serum, or other biological product, shall be in the form No. 7 in the Schedule hereto.

(2) The certificate of the Analyst in respect of any stock-remedy being a vaccine, serum, or other biological product shall be in the form No. 8 in the Schedule hereto.

FEES

23. (1) The fees payable by a proprietor on application for the registration of stock-remedies shall be as follows :—

	£	s.	d.
(*a*) On application for the registration of one stock-remedy	3	3	0
(*b*) On application by the same proprietor for the registration of each additional stock-remedy in respect of the same period of registration	0	5	0
(*c*) On application by the same proprietor for re-registration from time to time of one stock-remedy	1	10	0
(*d*) On application by the same proprietor for the re-registration of each additional stock-remedy in respect of the same period of registration ..	0	2	6

(2) The fee payable by a proprietor on an application for approval of any label or advertisement relating to any stock-remedy made at any time after registration of the stock-remedy and during the same period of registration shall be 10s.

(3) The fee payable on an application for the registration of a new proprietor shall be 2s. 6d. in respect of the first stock-remedy and 1s. in respect of each additional stock-remedy to which the application relates.

(4) The fee payable under section 13 (4) of the said Act by the proprietor or the vendor for a copy of the Analyst's certificate where no discrepancy materially to the prejudice of a purchaser is found in any sample of a stock-remedy taken and analysed under the said Act shall be £1 1s.

(5) The fee payable under section 14 (2) of the said Act on an application for the taking and analysing of any sample of any stock-remedy in the possession of the applicant shall be £2 2s.

(6) Where pursuant to the provisions of this regulation a fee is made payable on application, the application shall not be entertained until such fee is paid.

SCHEDULE

[Form No. 1, Reg. 6 (4)

The Stock-remedies Regulations 1947

REGISTER OF STOCK-REMEDIES

Application File No.	Name of Proprietor.	Address.	Name of Stock-remedy.	Date of Registration.	Period of Registration.	Remarks.

[Form No. 2, Reg. 7 (1)

The Stock-remedies Regulations 1947

APPLICATION FOR REGISTRATION OF A STOCK-REMEDY

The Registrar,
Stock-remedies Registration Board,
Care of Department of Agriculture,
Wellington C. 1.

PURSUANT to section 6 of the above Act application is hereby made for the registration of the undermentioned stock-remedy, the particulars of which are set out below :—

1. Name of stock-remedy :.....
2. Description or representation of every trade-mark to be used in respect of the stock-remedy, and registered number if registered :.....
3. Composition of stock-remedy :—
   Names of ingredients :.....
   Form of ingredients :.....
   Proportions present :.....*
4. Method of preparation of stock-remedy (if a vaccine, serum, or other biological product) :.....
5. Preventive or remedial properties claimed :.....
6. Date beyond which stock-remedy will be ineffective (where applicable) :.....
7. Enclosed with this application are the following documents :—
   (i) Specimen copy (in duplicate) of the label to be used in respect of the stock-remedy :
   (ii) Specimen copy (in duplicate) of the advertisement (if any) to be published in respect of the stock-remedy.
8. The prescribed fee for registration is also enclosed, viz. :.....

Dated at ....., this ..... day of ....., 19..

Signature of Applicant :.....

* Minimum proportion to be stated (to be expressed as a percentage by weight in the case of a solid stock-remedy or as a number of grammes per 100 c.c. in the case of a liquid stock-remedy) except where the ingredient is a poison within the meaning of the Poisons Act, 1934, when the maximum as well as the mimimum proportion must be stated. A semi-fluid stock-remedy is deemed to be a solid only if recommended for use by weight. It is unnecessary to state the proportions of ingredients which are shown under the heading "adjuvants," "emulsifiers," or "inert ingredients" and for which no specific efficacy is claimed.

N.B.—The prescribed fee may be lodged to the credit of the Public Account at any branch of the Bank of New Zealand and the first portion of the lodgment-slip should accompany the application for registration.

If payment of the prescribed fee is made by cheque exchange must be added where necessary.

[Form No. 3, Reg. 7 (3)

The Stock-remedies Regulations 1947

STATUTORY DECLARATION

I, A. B., of [*Address*], being the proprietor (*or* the managing director *or* manager *or* secretary *or* other responsible official [*Describing his office*], of the [*Name of firm or company*], the proprietors) of the stock-remedy (stock-remedies) set out in the application(s) for registration dated the ..... day of ....., 19.. attached or appended hereto, do solemnly and sincerely declare as follows :—

1. That I have been duly authorized to make the said application(s) and this declaration in support thereof on behalf of the said firm (*or* company). (Where declaration is made by the proprietor this paragraph should be deleted.)
2. That the signature to the said application(s) is in the proper handwriting of me this declarant.
3. That to the best of my knowledge, information, and belief the particulars relating to the stock-remedy (stock-remedies) set out in the said application(s) are true and correct in all respects.

And I make this solemn declaration conscientiously believing the same to be true and by virtue of the Justices of the Peace Act, 1927.

A. B.

Declared at ..... this ..... day of ....., 19.., before me—

Affix 3s. Stamp.

C. D.,
Justice of the Peace (*or* Solicitor *or* Notary Public).

[Form No. 4, Reg. 8 (1)

The Stock-remedies Regulations 1947

CERTIFICATE OF REGISTRATION OF A STOCK-REMEDY

THIS to certify that, pursuant to the application of [*Name of proprietor*] of [*Address of proprietor*], dated the ..... day of ....., 19.., the undermentioned stock-remedy (stock-remedies) is (are) hereby registered under the Stock-remedies Act, 1934, for the registration period ending the 30th day of September, 19..

Name of stock-remedy : .....

Dated at Wellington, this ..... day of ....., 19..

.....

Registrar, Stock-remedies Registration Board.

[Form No. 5, Reg. 9 (1)

The Stock-remedies Regulations 1947

APPLICATION FOR APPROVAL OF LABEL OR ADVERTISEMENT RELATING TO A REGISTERED STOCK-REMEDY

The Registrar,
Stock-remedies Registration Board,
Care of Department of Agriculture,
Wellington C. 1.

PURSUANT to section 8 (3) of the above Act, application is hereby made for approval of the label ((and) advertisement) of which two copies are attached or appended hereto relating to the undermentioned registered stock-remedy (stock remedies) :—

Dated at ....., this ..... day of ....., 19..

Signature of Applicant :.....

[Form No. 6, Reg. 21 (2)

The Stock-remedies Regulations 1947

FORM OF LABEL TO BE AFFIXED TO SAMPLE OF STOCK-REMEDY

SAMPLE of [*Name of stock remedy*] duly registered under the Stock-remedies Act, 1934, on the application of [*Name and address of proprietor*] taken on the ..... day of ....., 19.., for analysis under the said Act in the presence of [*Name of proprietor or vendor or other witness present when sample is taken.*]

Signature of Inspector :.....

Signature of proprietor, vendor, or other witness present : .....

[Form No. 7, Reg. 22 (1)

The Stock-remedies Regulations 1947

CERTIFCATE OF ANALYSIS OF STOCK-REMEDY, NOT BEING A VACCINE, SERUM, OR OTHER BIOLOGICAL PRODUCT

I, THE undersigned, an Analyst under the Stock-remedies Act, 1934, do hereby certify that on the .....day of ....., 19.., there was delivered to me personally by (or I received by registered post from) [*Insert name and address of the Inspector from whom the sample was received*] a sample of [*State name of stock-remedy*] for analysis in a [*State the nature of the package in which the sample was enclosed, the particulars set out in the label on the package, and a description of the impress of the seal*], and that I have analysed the same and that the result of my analysis is as follows : [*Analysis and statement of opinion as to whether any discrepancy found in the sample on comparison pursuant to section 13 (3) of the said Act would be materially to the prejudice of a purchaser*].

As witness my hand at ....., this .....day of ....., 19..

Signature of Analyst :.....

[Form No. 8, Reg. 22 (2)

The Stock-remedies Regulations 1947

CERTIFCATE OF ANALYSIS OF STOCK-REMEDY, BEING A VACCINE, SERUM, OR OTHER BIOLOGICAL PRODUCT

I, THE undersigned, an Analyst under the Stock-remedies Act, 1934, do hereby certify that on the ..... day of ....., 19.., there was delivered to me personally by (or I received by registered post from) [*Insert name and address of the Inspector from whom the sample was received*] a sample of [*State name of stock-remedy*] for analysis in a [*State the nature of the package in which the sample was enclosed, the particulars set out in the label on the package, and a description of the impress of the seal*], and that I have conducted a bacteriological (serological, biological, or as the case may be) examination of the same and that the result of my examination is as follows: [*Analysis and statement of opinion as to whether any discrepancy found in the sample on comparison pursuant to section 13 (3) of the said Act would be materially to the prejudice of a purchaser*].

As witness my hand at ....., this .....day of ....., 19..

Signature of Analyst:.....

W. O. HARVEY,
Clerk of the Executive Council.

Issued under the authority of the Regulations Act, 1936.
Date of notification in *Gazette:* 25th day of September, 1947.
These regulations are administered in the Department of Agriculture.

(Notice No. Ag. 4448.)