DUPLICATE

GOVERNMENT OF THE PEOPLE'S REPUBLIC OF BANGLADESH
MINISTRY OF HEALTH & FAMILY WELFARE
DIRECTORATE GENERAL OF DRUG ADMINISTRATION
AUSHAD BHAVAN, MOHAKHALI, DHAKA-1212, BANGLADESH
CERTIFICATE OF REGISTRATION

We hereby declare that **VERORAB** manufactured by **Sanofi Pasteur, Parc Industriel d'Incarville 27100 Val de Reuil France,** Marketing Authorization Holder (MAH) M/S. **Sanofi Pasteur, 14 Espace Henry Vallee, 69007 Lyon, France** and represented by **M/s. Sanofi Bangladesh Limited, Station Road, Tongi, Gazipur-1710, Bangladesh** is registered with Directorate General of Drug Administration and Licensing Authority (Drugs) under Reg. No. **102-2402-91**. The drug as described below is allowed to be imported into Bangladesh under **The Drugs Act. 1940 (XXIII of 1940), The Drug (control) Ordinance, 1982** and The Drug (Control) (Amendment) Act, 2006 subjected to the provision of import policy published by Government from time to time.

Name of the Product : **VERORAB**
Dosage Form : Injectable (Subcutaneous or Intramuscular route)
Packing : 1Vial 1 Dose + 1 Syringe Diluent / 5vials 1 Dose + 5 Solvent Ampoule
Full composition with specification (including Active Substance and all excipients)

Composition : Each 0.5ml contains

Name of ingredients	Specification	Quantity
Active Ingredient:		
Rabies Virus (Wistar: rabies PM/WI 38-1503-3 M strain) obtained from culture on Vero continuous cell lines, inactivated with beta-propiolactone.	WHO requirement	One immunizing dose such that the protective activity is equal to or greater than 2.5 I.U.
Excipients:		
Maltose	B.P.	q.s.
Human Albumin	Ph. Eur.	q.s.
Diluent:		
Solution of 4% sodium chloride	Ph. Eur.	0.5 ml

Registration Date: 13-08-1991
Validity: 17-08-2021

Instructions
To be dispensed only by or on the prescription of a registered physician.

Major General Md. Mustafizur Rahman
Director General
Directorate General of Drug Administration
&
Licensing Authority (Drugs)
Government of the People's Republic of Bangladesh
06 MAY 2019

Conditions:
(1). Labeling should be done in accordance with the provision of the Drugs Act and Rules which require that the name and address of the manufacturer, batch number, manufacturing date, expire date, M.R.P. (Maximum Retail Price) DAR No. (Drug Administration Reg. Number) etc. should be displayed on the label or container and also on the outer cover containing the container
(2). The registration will be valid for 5 (five) years from its date of issue unless it is revoked, suspended or cancelled earlier.
(3). The certificate will be treated as cancelled if any violation of conditions and the name or the formula of this product is changed or modified qualitatively or quantitatively without approval of the Licensing Authority.

Memo. No. DA /15-5/102/97/

Date 18/08/1991

c.c. to: M/S. Sanofi Bangladesh Limited
Station Road, Tongi, Gazipur
Contact: 6/2/A., Segun Bagicha, Dhaka-1000, Bangladesh.

MINISTRY OF HEALTH & FAMILY WELFARE ★ DIRECTORATE GENERAL OF DRUG ADMINISTRATION ★ GOVT OF THE PEOPLE'S REPUBLIC OF BANGLADESH
DHAKA

09/8/2021

For Director General,
Directorate General

This Registration shall remain valid up to 17-08-2026 unless it is suspended Cancelled or revoked earlier.

30.03.22

Nipa Chowdhury
Assistant Director
For Director General
Directorate General of Drug Administrator
&
Licensing Authority (Drugs)
Government of the People's Republic of Bangladesh