GOVERNMENT OF THE PEOPLE'S REPUBLIC OF BANGLADESH
MINISTRY OF HEALTH & FAMILY WELFARE
DIRECTORATE GENERAL OF DRUG ADMINISTRATION
105-106, MOTIJHEEL COMMERCIAL AREA, DHAKA-1000

CERTIFICATE OF REGISTRATION

We hereby declare that **MENOMUNE- A/C/Y/W-135** manufactured by **M/S. Sanofi Pasteur Inc., Discovery Drive, Swiftwater, PA 18370 USA** and represented by **M/s. Sanofi ~~-aventis~~ Bangladesh Limited, Station Road, Tongi, Gazipur-1710, Bangladesh** is registered with Directorate General of Drug Administration and Licensing Authority (Drugs) under Reg. No. 102-4779-13. 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 : **MENOMUNE- A/C/Y/W-135**

Dosage Form : **Suspension for subcutaneous injection**

Pack Size : **1 dose vial with 0.78ml diluent**

Composition: Each vial (1.5ml) contains

A.	Active Ingredient	Specification	Quantity
	Meningococcal Polysaccharide of *Neisseria menningitidis* Group A	In- House	50 µg
	Meningococcal Polysaccharide of *Neisseria menningitidis* Group C	In- House	50 µg
	Meningococcal Polysaccharide of *Neisseria menningitidis* Group Y	In- House	50 µg
	Meningococcal Polysaccharide of *Neisseria menningitidis* Group W-135	In- House	50 µg
B.	**Excipients:**		
	Lactose Monohydrate	NF	2.5 to 5.0 mg
	Sodium Chloride	USP	4.25 to 4.75 mg

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

Major General Md Jahangir Hossain Mollik
Director General 31 JAN 2013
Directorate General of Drug Administration
&
Licensing Authority (Drugs)
Government of the People's Republic of Bangladesh

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/ 1608 Date 04/02/2013

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

04.02.13
For Director General,
Directorate General of Drug Administration