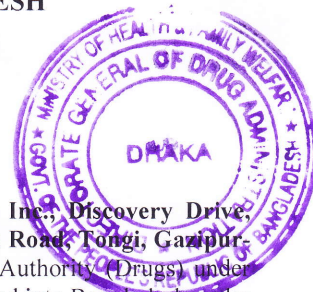


GOVERNMENT OF THE PEOPLE'S REPUBLIC OF BANGLADESH  
MINISTRY OF HEALTH & FAMILY WELFARE  
DIRECTORATE GENERAL OF DRUG ADMINISTRATION  
105-106, MOTIJHEEL COMMERCIAL AREA, DHAKA-1000



**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**

<b>A. <u>Active Ingredient</u></b>	<b><u>Specification</u></b>	<b><u>Quantity</u></b>
<b>Meningococcal Polysaccharide of <i>Neisseria meningitidis</i> Group A</b>	In- House	50 µg
<b>Meningococcal Polysaccharide of <i>Neisseria meningitidis</i> Group C</b>	In- House	50 µg
<b>Meningococcal Polysaccharide of <i>Neisseria meningitidis</i> Group Y</b>	In- House	50 µg
<b>Meningococcal Polysaccharide of <i>Neisseria meningitidis</i> Group W-135</b>	In- House	50 µg
<b>B. <u>Excipients:</u></b>		
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  
Directorate General of Drug Administration  
&  
Licensing Authority (Drugs)  
Government of the People's Republic of Bangladesh  
31 JAN 2013


**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...09.10.2013.....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