

# **RETENTION OF LICENCE**

**Retention of Licence to manufacture for sale (or for distribution) of drugs other than those specified in Schedules C, C (1) and X**

Certified that Licence in Form 25 No: G/25/1014 Issue Date: 18-Jan-1991

To M/s. ASCOT PHARMACHEM PVT. LTD

to manufacture the following categories of drugs being drugs other than those specified in Schedules C and C (1) and Schedule "X" to the Drugs and Cosmetics Rules 1945, on the premises situated at 98/7, G.I.D.C. ESTATE, NANDESARI - 391 340, VADODARA

has been retained from: 01-Jan-2023 To 31-Dec-2027

Name(s) of drugs

: As per list Approved

Names of approved competent technical staff

: As per list Approved

Signature :

(This Document is Digitally Signed.)  
**Dr. H. G. KOSHIA**

Designation :

**Commissioner**  
Food & Drugs Control Administration  
Gujarat State

Date: 31-May-2023

Note: You are requested to apply for the Retention of the above licence in 3 months before its VALIDITY EXPIRES.



Reg ID : 636450

Doc ID: **RC704300001014**

**ASCOT PHARMACHEM PVT. LTD**

Print Date : **31/05/2023 03:42 PM**

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# Food & Drugs Control Administration

Office of the Commissioner

Block No. 8, 1st Floor, Dr. Jivraj Mehta Bhavan, Gandhinagar, Gujarat State

## RETENTION PRODUCT PERMISSION

To,

**ASCOT PHARMACHEM PVT. LTD**

**98/7, G.I.D.C. ESTATE, NANDESARI - 391 340, VADODARA**

Reference : Your Application inward ID: **660786** Dated : **02-Jan-2023** (Reg ID : **636450**)

With reference to your Inward application, this is to inform you that your said application is considered & following **RETENTION PRODUCT PERMISSION** has been granted, under the **LICENSE NO. G/25/1014 IN FORM NO. 25**.

Product Section : **Bulk Drugs**, Product Sub Section : **Excipients**

Sr. No.	Name of Drugs																						
1	<p>Prod ID : <b>67249</b> Permission Date : <b>10-Dec-1993</b> Type : <b>Normal</b> Permission Purpose: <b>G-General</b></p> <p>Generic Name : <b>Sodium Starch Glycolate IP</b> Brand Name : <b>N.A.</b> <b>Brand No :1</b> Composition Title :</p> <table><thead><tr><th>Composition</th><th>Ingredients</th><th>Standards</th><th>Strength</th><th>UOM</th><th>Equivalent to</th></tr></thead><tbody><tr><td>Excipients</td><td>Sodium Starch Glycolate</td><td>IP</td><td>0</td><td>----</td><td></td></tr></tbody></table> <p>Product Package Size Details:</p> <table><thead><tr><th>Product Size</th><th>UOM</th><th>Container</th><th>Dose</th><th>Remark</th></tr></thead><tbody><tr><td>25</td><td>Kg.</td><td>Fibre Drums</td><td>Multi Dose</td><td></td></tr></tbody></table>	Composition	Ingredients	Standards	Strength	UOM	Equivalent to	Excipients	Sodium Starch Glycolate	IP	0	----		Product Size	UOM	Container	Dose	Remark	25	Kg.	Fibre Drums	Multi Dose	
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### Terms and Conditions

- 1) Licensee should comply with all the provisions of Drugs & Cosmetics Act, 1940 & Rules 1945 as amended time to time.
- 2) Licensee should comply with all the provisions of Drugs (Price Control) Order 2013 as amended time to time (wherever applicable).
- 3) Licensee should abide by all the provision of Drugs & Magic Remedies (Objectionable Advertisement) Act, 1954 & Rules 1955 as amended up to date.
- 4) Licensee should not manufacture any drug by a name belonging to another manufacturer.
- 5) Licensee should not manufacture or sell drugs even if it is included in the approved list of product, if it is or as and when banned by Licensing Authority or Drugs Controller General of India or Government of India.
- 6) The permission is granted subject to the condition that, the product is safe for use in context of pharmaceutical Aids additives and excipients used in the formulation.
- 7) Any addition thereto or any deletion therefore will not be carried out without permission of Licensing Authority.
- 8) Above Retention Product Permission is granted based on undertaking with respect to BCS classification and declaration under Form - 51.

(This Document is Digitally Signed)

**RAVAT H. L.**

**( Deputy Commissioner )**

For Commissioner

Food & Drugs Control Administration  
Gujarat State, Gandhinagar



Reg ID : **636450**

Doc ID: **PP882200067249**

**ASCOT PHARMACHEM PVT. LTD**

License No - **G/25/1014** From Date: **01-Jan-2023** To Date: **31-Dec-2027**

Print Date : **31/05/2023 04:26 PM**

Page 1 of 1

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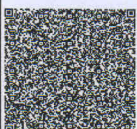
Sr. No.	Name of Drugs																						
1	<p>Prod ID : <b>67254</b> Permission Date : <b>10-Dec-1993</b> Type : <b>Normal</b> Permission Purpose: <b>E-Export only</b></p> <p>Generic Name : <b>Sodium Starch Glycolate B.P.</b> Brand Name : <b>N.A.</b> Brand No : <b>1</b> Composition Title :</p> <table><thead><tr><th>Composition</th><th>Ingredients</th><th>Standards</th><th>Strength</th><th>UOM</th><th>Equivalent to</th></tr></thead><tbody><tr><td>API</td><td>SODIUM STARCH GLYCOLATE</td><td>BP</td><td>0</td><td>----</td><td></td></tr></tbody></table> <p>Product Package Size Details:</p> <table><thead><tr><th>Product Size</th><th>UOM</th><th>Container</th><th>Dose</th><th>Remark</th></tr></thead><tbody><tr><td>25</td><td>Kg.</td><td>Fibre Drums</td><td>Multi Dose</td><td></td></tr></tbody></table>	Composition	Ingredients	Standards	Strength	UOM	Equivalent to	API	SODIUM STARCH GLYCOLATE	BP	0	----		Product Size	UOM	Container	Dose	Remark	25	Kg.	Fibre Drums	Multi Dose	
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**( Deputy Commissioner )**  
For Commissioner  
Food & Drugs Control Administration  
Gujarat State, Gandhinagar



Reg ID : **636450**

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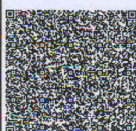
Sr. No.	Name of Drugs																						
1	<p>Prod ID : <b>67259</b> Permission Date : <b>28-Sep-1999</b> Type : <b>Normal</b> Permission Purpose: <b>G-General</b></p> <p>Generic Name : <b>CROSCARMELLOSE SODIUM U.S.P./B.P.</b> Brand Name : <b>N.A.</b> Brand No : <b>1</b> Composition Title :</p> <table><thead><tr><th>Composition</th><th>Ingredients</th><th>Standards</th><th>Strength</th><th>UOM</th><th>Equivalent to</th></tr></thead><tbody><tr><td>API</td><td>CROSCARMELLSE SODIUM U.S.P.B.P.</td><td>BP</td><td>0</td><td>----</td><td></td></tr></tbody></table> <p>Product Package Size Details:</p> <table><thead><tr><th>Product Size</th><th>UOM</th><th>Container</th><th>Dose</th><th>Remark</th></tr></thead><tbody><tr><td>25</td><td>Kg.</td><td>Fibre Drums</td><td>Multi Dose</td><td></td></tr></tbody></table>	Composition	Ingredients	Standards	Strength	UOM	Equivalent to	API	CROSCARMELLSE SODIUM U.S.P.B.P.	BP	0	----		Product Size	UOM	Container	Dose	Remark	25	Kg.	Fibre Drums	Multi Dose	
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