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**

Signature valid
Digitally signed by
Date: 2023.05.31
15:45:52 +05:30
Reason: Sign
Location: AHD

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	Prod ID : **67249** Permission Date : **10-Dec-1993** Type : **Normal** Permission Purpose: **G-General**

Generic Name : **Sodium Starch Glycolate IP** Brand Name : **N.A.**
Brand No :1
Composition Title :

Composition	Ingredients	Standards	Strength	UOM	Equivalent to
Excipients	Sodium Starch Glycolate	IP	0	-----	

Product Package Size Details:

Product Size	UOM	Container	Dose	Remark
25	Kg.	Fibre Drums	Multi Dose	

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 Commisioner)
For Commissioner
Food & Drugs Control Administration
Gujarat State, Gandhinagar

Signature valid
Digitally signed by
Date: 2023.06.01 13:22:15 +05:30
Reason: Sign
Location: AHD

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	Prod ID : **67254** Permission Date : **10-Dec-1993** Type : **Normal** Permission Purpose: **E-Export only**

Generic Name : **Sodium Starch Glycolate B.P.** Brand Name : **N.A.**
Brand No :1
Composition Title :

Composition	Ingredients	Standards	Strength	UOM	Equivalent to
API	SODIUM STARCH GLYCOLATE	BP	0	-----	

Product Package Size Details:

Product Size	UOM	Container	Dose	Remark
25	Kg.	Fibre Drums	Multi Dose	

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).
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(Deputy Commisioner)
For Commissioner
Food & Drugs Control Administration
Gujarat State, Gandhinagar

Signature valid
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Reason: Sign
Location: AHD

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	Prod ID : **67259** Permission Date : **28-Sep-1999** Type : **Normal** Permission Purpose: **G-General**

Generic Name : **CROSCARMELLOSE SODIUM U.S.P./B.P.** Brand Name : **N.A.**
Brand No :1
Composition Title :

Composition	Ingredients	Standards	Strength	UOM	Equivalent to
API	CROSCARMELLSE SODIUM U.S.P.B.P.	BP	0	-----	

Product Package Size Details:

Product Size	UOM	Container	Dose	Remark
25	Kg.	Fibre Drums	Multi Dose	

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.
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RAVAT H. L.
(Deputy Commisioner)
For Commissioner
Food & Drugs Control Administration
Gujarat State, Gandhinagar

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