

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

New Delhi, dated

To:

15 MAR 2011

M/s. The Madras Pharmaceuticals  
No. 137-B, Old Mahabalipuram Road,  
Karapakkam, Chennai- 600 096

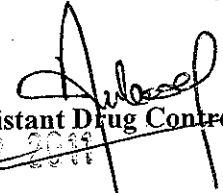
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 03/03/2011 received by this office vide diary no. 10022 dt. 03/03/2011, I enclose licence(s) Nos. BD-218-18716 dated \_\_\_\_\_. This/These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/12/03-DC(Re-Reg-2009)

Copy together with copy of Licence No. BD-218-18716 dated

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-218-18716

Date 5 MAR 2011

1. M/S. The Madras Pharmaceuticals No. 137-B, Old Mahabalipuram Road, Karapakkam, Chennai- 600 096 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Zhejiang East- Asia Pharmaceutical Co. Ltd. Economic Development Zone Of Sanmen County Zhejiang China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
2. This licence shall be in force from 08/03/2011 to 30/06/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Ofloxacin BP

2. Levofloxacin Hemihydrate

Item (Two) Only

Place: New Delhi

Date: 5 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती

ARVIND KUKRETI

रिजिस्ट्रार औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

आर्य समाज महाविद्यालय

Directorate General of Health Services,

23, Patna, Konda Road

New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in each part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi dated  
15 MAR 2011

M/s. Ruskin Chemi Pharm  
4/A Bhang wadi, Shopping Arcade, Kalbadevi Road,  
Mumbai-400002

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/02/2011 received by this office vide dt. no. 9359 dt. 28/02/2011, I enclose licence(s) Nos. BD-859-18717 dated 5 MAR 2011. This/These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

Assistant Drug Controller (India)

No. 6-3/BD/28/10-DC

Copy together with copy of Licence No. BD-859-18717 dated 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-859-18717

Date 5 MAR 2011

1. M/S. Ruskin Chemi Pharm 4/A Bhang wadi, Shopping Arcade, Kalbadevi Road, Mumbai-400002 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Henan Kangtai Pharmaceutical Group Corporation No. 338, Kangtai Road, Sishui Town, Xinyang 450141 China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
2. This licence shall be in force from 08/03/2011 to 30/06/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Norfloxacin IP

Item (One) Only

Place: New Delhi

Date: 5 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

स्वास्थ्य सेवा विभाग

Directorate General of Health Services

F.D.A. Bhawan, Kotla Road

New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, dated  
15 MAR 2011

M/s. Harshada Corporation  
Bldg, No.1, Ground Floor, No.36, Bomanji Master Road,  
Opp. Kalbadevi Post Office  
Mumbai-400002

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/02/2011 received by this office vide diary no. 9549 dt. 28/02/2011, I enclose licence(s) Nos. BD-859-18718 dated 5 MAR 2011. These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

- OFFICE COPY**
1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
  2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
  3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
  4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
  5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
  6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/28/10-DC

Copy together with copy of Licence No. BD-859-18718 dated 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-859-18718

Date 5 MAR 2011

1. M/S. Harshada Corporation Bldg, No.1, Ground Floor, No.36, Bomanji Master Road, Opp. Kalbadevi Post Office Mumbai-400002 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Henan Kangtai Pharmaceutical Group Corporation No. 338, Kangtai Road, Sishui Town, Xinyang 450141 China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.

2. This licence shall be in force from 08/03/2011 to 30/06/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Norfloxacin IP

Item (One) Only

Place: New Delhi

Date : \_\_\_\_\_

OFFICE COPY

LICENSING AUTHORITY

अरवि कुक्रेती  
ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)

स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
F.D.A. Bhawan, Kolla Road  
New Delhi - 110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, dated  
15 MAR 2011

M/s. Ruskin Chemi Pharm  
4/A Bhang wadi, Shopping Arcade, Kalbadevi Road,  
Mumbai-400002

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/02/2011 received by this office vide diary no. 9360 dt. 28/02/2011, I enclose licence(s) Nos. BD-709-18719 dated 5 MAR 2011. These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/85/07-DC(Re-Reg-2011)

Copy together with copy of Licence No. BD-709-18719 dated 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-709-18719

Date 5 MAR 2011

1. M/S. Ruskin Chemi Pharm 4/A Bhang wadi, Shopping Arcade, Kalbadevi Road, Mumbai-400002 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Zhejiang Guobang Pharmaceutical Co. Ltd. No. 6, Weiwu Road, Hangzhou Gulf Shangyu Industrial Zone Zhejiang, P.R.China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
2. This licence shall be in force from 08/03/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Annexure-1

Item (Four) Only

Place: New Delhi

Date 5 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविंद कुक्रेती  
ARVIND KUKRETY  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
F.D.A. Bhawan, Kolla Road  
Kolkata - 700 017

Conditions of Licence

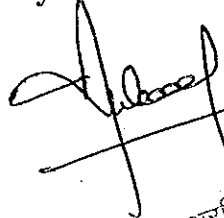
1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution?;

**ANNEXURE-1 (List of Products of Form 10 License No. BD-709-18719)**

1. Azithromycin IP
2. Ciprofloxacin Hydrochloride IP
3. Roxithromycin IP
4. Clarithromycin IP

**OFFICE COPY**

Item (Four) Only

  
15 MAR 2011  
अरविन्द कुकरती  
ARVIND KUKRETY  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Director General of Health Services  
L.O. A. Bhawan, Kola Road  
New Delhi-110 002

5

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

New Delhi, dated

To:

1 5 MAR 2011

M/s. Ace Pharma  
Basement No. 11-12, Star Trade Centre,  
Chamunda Circle, S.V. Road, Borivali (W)  
Mumbai-400092

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. ACE/DCGI/17/2010-11 Dated 23/02/2011 received by this office vide diary no. 9413 dt. 28/02/2011, I enclose licence(s) Nos. BD-709-18720 dated 1 5 MAR 2011. These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

OFFICE COPY

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/85/07-DC(Re-Reg-2011)

Copy together with copy of Licence No. BD-709-18720 dated 1 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-709-18720

Date 15 MAR 2011

1. M/S. Ace Pharma Basement No. 11-12, Star Trade Centre, Chamunda Circle, S.V. Road, Borivali (W) Mumbai-400092 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Zhejiang Guobang Pharmaceutical Co. Ltd. No. 6, Weiwu Road, Hangzhou Gulf Shangyu Industrial Zone Zhejiang, P.R.China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
2. This licence shall be in force from 08/03/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported


1. Azithromycin IP

2. Ciprofloxacin Hydrochloride IP

Item (Two), Only

OFFICE COPY

Place: New Delhi  
Date: 15 MAR 2011

  
LICENSING AUTHORITY  
ARVIND KULKARNI  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा मंत्रालय  
Directorate General of Health Services  
F.D.A. Bhawan, Kotla Road  
New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi dated 5 MAR 2011

M/s. Pradipkumar & Co.  
Gala No. 1, Gr. Floor, Munisuvrat Complex,  
Phase-II, Bldg. L-2, Rehnal Village, Tal- Bhiwandi  
Dist. Thane

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 21/02/2011 received by this office vide diary no. 9410 dt. 28/02/2011, I enclose licence(s) Nos. BD-901-18721 dated 5 MAR 2011. This/These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (f) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

Assistant Drug Controller (India)

No. 6-3/BD/51/10-DC

Copy together with copy of Licence No. BD-901-18721 dated 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-901-18721

Date 5 MAR 2011

1. M/S. Pradipkumar & Co. Gala No. 1, Gr. Floor, Munisuvrat Complex, Phase-II, Bldg. L-2, Rehnal Village, Tal- Bhiwandi Dist. Thane is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Tianjin Tianyao Pharmaceuticals Co. Ltd. No. 19, Xinye 9th Street, West Area of Tianjin Economic and Technological Development Area, Tianjin 300462 China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.

2. This licence shall be in force from 08/03/2011 to 31/12/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Prednisone IP
2. Prednisolone (Micronized) IP
3. Spironolactone (Micronized) IP

Item (Three) Only

Place: New Delhi

Date: 5 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY  
सहायक अधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा विभाग

Seal/Stamp  
Directorate General of Health Services  
F.D.A. Bhawan, Kolla Road  
New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, dated  
5 MAR 2011

M/s. Anuh Pharma Ltd.,  
E-17/3 & 17/4, MIDC, Opp. Brij Ice Factory, Tarapur, Boisar,  
Dist. Thane - 401 506

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

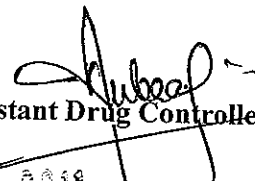
Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 22/02/2011 received by this office vide diary no. 9416 dt. 28/02/2011, I enclose licence(s) Nos. BD-901-18722 dated 1 5 MAR 2011. This/These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

OFFICE COPY

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/51/10-DC

Copy together with copy of Licence No. BD-901-18722 dated 1 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-901-18722

Date: 5 MAR 2011

1. M/S. Anuh Pharma Ltd., E-17/3 & 17/4, MIDC, Opp. Brij Ice Factory, Tarapur, Boisar, Dist. Thane - 401 506 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Tianjin Tianyao Pharmaceuticals Co. Ltd. No. 19, Xinye 9th Street, West Area of Tianjin Economic and Technological Development Area, Tianjin 300462 China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
2. This licence shall be in force from 08/03/2011 to 31/12/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Prednisolone (Micronized) IP

Item (One) Only

Place: New Delhi

Date: 9 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरती  
ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)

स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
F. D. A. Bhawan, Kolla Road  
New Delhi

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, dated  
15 MAR 2011

M/s. Black Rose Industries Ltd.  
145-A, Mittal Tower, Nariman Point  
Mumbai-400021

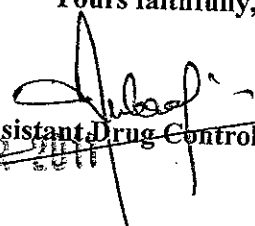
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 22/02/2011 received by this office vide diary no. 9411 dt. 28/02/2011, I enclose licence(s) Nos. BD-901-18723 dated 15 MAR 2011 these licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/51/10-DC

Copy together with copy of Licence No. BD-901-18723 dated

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-901-18723

Date 15 MAR 2011

1. M/S. Black Rose Industries Ltd. 145-A, Mittal Tower, Nariman Point Mumbai-400021 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Tianjin Tianyao Pharmaceuticals Co. Ltd. No. 19, Xinye 9th Street, West Area of Tianjin Economic and Technological Development Area, Tianjin 300462 China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.

2. This licence shall be in force from 08/03/2011 to 31/12/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Prednisone IP
2. Prednisolone (Micronized) IP
3. Spironolactone (Micronized) IP

Item (Three) Only

Place: New Delhi  
Date: 15 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती  
ABVIND KUKRETI  
राष्ट्रीय औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, dated

**15 MAR 2011**

M/s. Black Rose Industries Ltd.  
145-A, Mittal Tower, Nariman Point  
Mumbai-400021

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 22/02/2011 received by this office vide diary no. 9412 dt. 28/02/2011, I enclose licence(s) Nos. BD-59-18724 dated 15 MAR 2011. These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/78/02-DC(Pt-1)(Re-Registration 2008)

Copy together with copy of Licence No. BD-59-18724 dated 15 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-59-18724

Date 15 MAR 2011

1. M/S. Black Rose Industries Ltd. 145-A, Mittal Tower, Nariman Point Mumbai-400021 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Fermic S.A. de C.V. Reforma No. 873, Colonia San Nicolas Tolentino 09850 Mexico, D.F. Mexico any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.

2. This licence shall be in force from 08/03/2011 to 31/12/2011 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Clavulanate Potassium+Avicel

2. Clavulanate Potassium+Syloid

3. Clavulanate Potassium - Sterile+Amoxicillin Sodium Sterile

Item (Three) Only

Place: New Delhi 15 MAR 2011

Date : \_\_\_\_\_

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती  
ARVIND KUKRETI

सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)

स्वास्थ्य सेवा महानिदेशालय

Directorate General of Health Services  
New Delhi

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, dated  
**15 MAR 2011**

M/s. G.C. Chemie Pharmie Ltd.  
5-C, Shri Laxmi Ind. Estate New Link Road, Andheri (West),  
Mumbai-400053

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 26/02/2011 received by this office vide diary no. 9414 dt. 28/02/2011, I enclose licence(s) Nos. BD-404-18725 dated 15 MAR 2011 this licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/07/04-DC(Re-Registration 2010)

Copy together with copy of Licence No. BD-404-18725 dated 15 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-404-18725

Date 15 MAR 2011

1. M/S. G.C. Chemie Pharmie Ltd. 5-C, Shri Laxmi Ind. Estate New Link Road, Andheri (West), Mumbai-400053 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Hubei Huazhong Pharmaceutical Co. Ltd., No. 71, West Chunyuan Road, Xiangfan City, Hubei Province China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
2. This licence shall be in force from 08/03/2011 to 31/08/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Thiamine Hydrochloride IP (Vitamin B1 Hydro Chloride)
2. Thiamine Mononitrate IP (Vitamin B1 Mononitrate)

Item (Two) Only

Place: New Delhi

Date: 15 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती  
ARVIND KUKRETI

सहायक औषधि नियंत्रक  
Assistant Drugs Controller (India)

स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi, 28/02/2011  
9.5 MAR 2011

M/s. Sheetal Pharma  
Shantivilla, Shantivan Tower Compound,  
Plot No. 3, 3rd floor, Devidas Lane, Borivali (West)  
Mumbai-400103

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/02/2011 received by this office vide diary no. 9388 dt. 28/02/2011, I enclose licence(s) Nos. BD-679-18726 dated 9.5 MAR 2011. This/these licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

OFFICE COPY

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/08/2007-DC

Copy together with copy of Licence No. BD-679-18726 dated 9.5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-679-18726

Date 1.5 MAR 2011

- M/S. Sheetal Pharma Shantivilla, Shantivan Tower Compound, Plot No. 3, 3rd floor, Devidas Lane, Borivali (West) Mumbai-400103 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Chongqing Daxin Pharmaceutical Co. Ltd. No. 22, Chuangzao Road, Beibei District, Chongqing 400700 P. R.China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.
- This licence shall be in force from 08/03/2011 to 31/07/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Amikacin Sulfate IP

Item (One) Only

Place: New Delhi

Date : 1.5 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती

ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

स्वास्थ्य सेवा महानिदेशालय

Directorate General of Health Services

New Delhi - 110 002

Conditions of Licence

- A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
- Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
- The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
- The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";

From:

The Drugs Controller (India)  
Directorate General of Health Services  
FDA Bhawan, Kotla Road  
Phone No.-011-23236965, Fax No.-23236973

To:

New Delhi dated  
15 MAR 2011

M/s. Man Mill Chemicals Pvt. Ltd.  
Gala No. 20, Bldg, No. 3, 'B' Wing,  
1st Fl, Manish Comp. Anjurphata, Rehnal Village  
Bhivandi

**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 26/02/2011 received by this office vide diary no. 9415 dt. 28/02/2011, I enclose licence(s) Nos. BD-218-18727 dated 15 MAR 2011. This/These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

1. I am to point out the provisions of Drugs Act, 1940 are in addition to and not derogation of any other law for the time being in force and as such the licences issued under Drugs Act will be in addition to and distinct from any licences which may be necessary under the Import Trade Control Regulations made of the Government of India, Ministry of Commerce.
2. The import licence(s) mentioned in para (1) above will not accordingly to itself/themselves be sufficient authority for import of Drugs covered by that/those licence if under the Import Trade Control Regulations of the Commerce Ministry separate licences are required for import of such drug(s).
3. I am therefore, to advise you to obtain, where necessary licences for import of drugs in question under the Import Trade Control Regulations.
4. Any literature or packing accompanying the drug or any matter stated on the label should not contravene the provisions of the Drugs and Magic Remedies (Objectionable Advertisement) Act.
5. The Assistant Drugs Controller (India) and Technical Officer of the Central Organization at the ports will be Officers authorized to inspect the premises of importers establishments for the purpose of Rule 26 of the Drugs Rules.
6. Please acknowledge receipt of this letter and its enclosures.

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/12/03-DC(Re-Reg-2009)

Copy together with copy of Licence No. BD-218-18727 dated 5 MAR 2011

1. Asstt. Drugs Controller (India), New Customs, Fort, Mumbai.
2. Asstt. Drugs Controller (India), Customs House, Kolkata.
3. Asstt. Drugs Controller (India), Customs House, Chennai.
4. Asstt. Drugs Controller (India), IGIA, New Delhi.

FORM 10  
(See rules 23 and 27)

Licence to import drugs (excluding those specified in Schedule X) to the Drugs and Cosmetic  
Rules, 1945

Licence Number BD-218-18727

Date 1.5 MAR 2011

1. M/S. Man Mill Chemicals Pvt. Ltd. Gala No. 20, Bldg. No. 3, 'B' Wing, 1st Fl, Manish Comp. Anjurphata, Rehnal Village Bhivandi is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Zhejiang East- Asia Pharmaceutical Co. Ltd. Economic Development Zone Of Sanmen County Zhejiang China any other drugs manufactured by the said manufacturer as may from time to time be endorsed on this licence.

2. This licence shall be in force from 08/03/2011 to 30/06/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

**Ofloxacin BP**

Item (One) Only

Place: New Delhi  
Date: 1.5 MAR 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरती  
ARVIND KUKRETI  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
New Delhi - 110 002

- Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and authenticated by the manufacturer with date of testing, date of release and date of forwarding such reports. The imported batch of each drug shall be subjected to examination and testing as the licensing authority deems fit prior to its marketing.
3. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
4. The licensee shall inform the licensing authority forthwith in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the licensing authority in the name of the firm with the changed constitution";