

From:

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

To:

M/s. Biocon Limited
Konnar Industries, 29/A, 1st Floor,
Veerasandra Industrial Area, Electronics City
Bangalore-560100

New Delhi, dated
17 JUN 2011

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 19/05/2011 received by this office vide diary no. 24285 dt. 23/05/2011, I enclose licence(s) Nos. BD-795-19171 dated 17 JUN 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 (I) 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/31/08-DC

Copy together with copy of Licence No. BD-795-19171 dated 17 JUN 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-795-19171

Date 17 JUN 2011

1. M/S. Biocon Limited Konnar Industries, 29/A, 1st Floor, Veerasandra Industrial Area, Electronics City Bangalore-560100 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Zhejiang Hisun Pharmaceutical Co. Ltd. 1 Haizheng Avenue, jiaojiang District Taizhou City, Zhejiang Province, 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 14/06/2011 to 30/06/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Meropenem with Sodium Carbonate Mixture
2. Imipenem & Cilastatin USP (With Buffered Sodium Bicarbonate)

Item (Two) Only

Place: New Delhi

Date 17 JUN 2011


LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETY
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Services
E. D. A. Bhawan, Kolla Road
New Delhi - 110002

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

New Delhi, dated

To:

7 JUN 2011

M/s. Biocon Limited
20th K.M. Hosur Road Electronic City
Bangalore-560100

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 19/05/2011 received by this office vide diary no. 24286 dt. 23/05/2011, I enclose licence(s) Nos. BD-825-19172 dated 7 JUN 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 (I) 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/86/09-DC

Copy together with copy of Licence No. BD-825-19172 dated 7 JUN 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-825-19172

Date 17 JUN 2011

1. M/S. Biocon Limited 20th K.M. Hosur Road Electronic City Bangalore-560100
is hereby licensed into India during the period for which this licence is in force, the
drugs specified below, manufactured By M/s. Henan Topfond Pharmaceutical Co.
Ltd. No. 1219, Jiaotong Road, Zhumadian, Henan 463000 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 14/06/2011 to 31/01/2013 unless it is sooner
suspended or cancelled under the said rules.

Names of drugs to be imported

Simvastatin IP

Item (One) Only

Place: New Delhi

Date: 17 JUN 2011


LICENSING AUTHORITY

अरविन्द कुक्रेती
ARVIND KUKRETY
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Seal/Stamp

Conditions of Licence

Directorate General of Health Services
F.D.A. Bhawan, Kotla Road

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";

3

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
17 JUN 2011

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

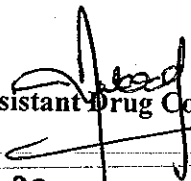
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 19/05/2011 received by this office vide diary no. 24364 dt. 23/05/2011, I enclose licence(s) Nos. BD-917-19173 dated 17 JUN 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 (I) 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/12/02-DC(Pt-1) (Re-Reg-2010)

Copy together with copy of Licence No. BD-917-19173 dated 17 JUN 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-917-19173

Date 17 JUN 2011

1. M/S. Harshada Corporation Bldg. No. 1, Gorund 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 Lihua Pharmaceutical Co. Ltd. Middle Of Huanghe Street, Anyang Hi-Tech Industry Development Zone Henan 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 14/06/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Prednisolone IP
2. Prednisolone Acetate IP
3. Hydrocortisone IP

Item (Three) Only

Place: New Delhi

Date : 17 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेली
ARVIND KUKRETI

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

स्वास्थ्य सेवा महानिदेशालय
General of Health Services

F.D.A. Bhawan,
New Delhi-110 002

Conditions of Licence

1. A photocopy of licence shall be displayed in a prominent place, ~~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";

4

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

17 JUN 2011

M/s. Mac-Chem Products (India) Pvt Ltd.
N-211/2/10, M.I.D.C Boisar,
Dist. Thane-401506

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. MCPIPL/DCGI/54/1011 Dated 20/05/2011 received by this office vide diary no. 24359 dt. 23/05/2011, I enclose licence(s) Nos. BD-917-19174 dated 17 JUN 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 (I) 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/12/02-DC(Pt-1) (Re-Reg-2010)

Copy together with copy of Licence No. BD-917-19174 dated

17 JUN 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-917-19174

Date 7 JUN 2011

1. M/S. Mac-Chem Products (India) Pvt Ltd. N-211/2/10, M.I.D.C Boisar, Dist. Thane-401506 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Henan Lihua Pharmaceutical Co. Ltd. Middle Of Huanghe Street, Anyang Hi-Tech Industry Development Zone Henan 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 14/06/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

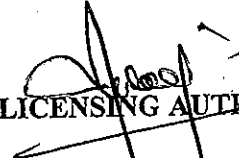
Names of drugs to be imported

Hydrocortisone IP

Item (One) Only

Place: New Delhi

Date: 17 JUN 2011


LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND Kulkarni

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

स्वास्थ्य सेवा महानिदेशालय
Directorate 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

New Delhi, dated

To:

17 JUN 2011

M/s. Add-Biotec,
Unit 110-B, 1st Floor, Andheri Indst Prms,
Co-Op. Soc. Plt N 22, Vira Desai Rd Andheri W
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 19/05/2011 received by this office vide diary no. 24360 dt. 23/05/2011, I enclose licence(s) Nos. BD-917-19175 dated 17 JUN 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 (I) 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/12/02-DC(Pt-1) (Re-Reg-2010)

Copy together with copy of Licence No. BD-917-19175 dated 17 JUN 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-917-19175

Date 17 JUN 2011

1. M/S. Add-Biotec, Unit 110-B, 1st Floor, Andheri Indst Prms, Co-Op. Soc. Plt N 22, Vira Desai Rd Andheri W 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. Henan Lihua Pharmaceutical Co. Ltd. Middle Of Huanghe Street, Anyang Hi-Tech Industry Development Zone Henan 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 14/06/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Prednisolone IP
2. Prednisolone Acetate IP
3. Hydrocortisone IP

Item (One) Only

Place: New Delhi

Date : 17 JUN 2011

LICENSING AUTHORITY

अरविंद कुक्रेजी
ARVIND KUKRETY

Conditions of Licence

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

संयुक्त स्वास्थ्य सेवा विभाग
Directorate General of Health Services

F.D.A. Bhawan, Kotta Road

New Delhi - 110 002

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

17 JUN 2011

M/s. Lupin Limited
Shree Arihant Compound, Gala No. 1 to 9,
Bldg No. 21, 1st Fl Kalher Pipeline, kalher,
Bhiwandi -421302

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 19/05/2011, received by this office vide diary no. 24142 dt. 23/05/2011, I enclose licence(s) Nos. BD-901-19176 dated 17 JUN 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 (I) 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-901-19176 dated 17 JUN 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-901-19176

Date 7 JUN 2011

1. M/S. Lupin Limited Shree Arihant Compound, Gala No. 1 to 9, Bldg No. 21, 1st
Fl Kalher Pipeline, kalher, Bhiwandi -421302 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 14/06/2011 to 31/12/2013 unless it is sooner
suspended or cancelled under the said rules.

Names of drugs to be imported

Spironolactone(Micronized)IP

Item (One) Only

Place: New Delhi

Date : 7 JUN 2011

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

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

New Delhi, dated

To:

17 JUN 2011

M/s. Dr. Reddy's Laboratories Ltd.
Sy. No. 41, Bachupally (V) Qutubullapur(M)
Ranga Reddy Dist.

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. DRL/RA/1007-11 Dated 19/05/2011 received by this office vide diary no. 24250 dt. 23/05/2011, I enclose licence(s) Nos. BD-556-19177 dated 17 JUN 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/23/05-DC(Pt-1)(Re-Rreg-2008)

Copy together with copy of Licence No. BD-556-19177 dated

17 JUN 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-556-19177

Date 17 JUN 2011

1. M/S. Dr. Reddy's Laboratories Ltd. Sy. No. 41, Bachupally (V) Outubullapur(M) Ranga Reddy Dist. is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. DSM Nutritional Products Pvt. Ltd. Dairy, Ayrshire Ka 245 JJ, Scotland United Kingdom 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 14/06/2011 to 31/07/2011 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Ascorbic Acid IP

Item (One) Only

Place: New Delhi

Date : 17 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETI
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
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;

⑧

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:

7 JUN 2011

M/s. Sun Pharmaceutical Industries
6-9 EPIP, Kartholi, Bari-Brahmana, Jammu,
J & K

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. SPIJ/F/05/14 Dated 21/05/2011 received by this office vide diary no. 24418 dt. 23/05/2011, I enclose licence(s) Nos. BD-845-19178 dated 7 JUN 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/123/09-DC

Copy together with copy of Licence No. BD-845-19178 dated

7 JUN 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-845-19178

Date 17 JUN 2011

1. M/S. Sun Pharmaceutical Industries 6-9 EPIP, Kartholi, Bari-Brahmana, Jammu, J & K is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Pharmaxyn Laboratories Ltd. No. 09 Minpu Road, Qingyang, Jiangyin, Wuxi, Jiagsu 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 14/06/2011 to 30/04/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Voglibose

Item (One) Only

Place: New Delhi

Date: 17 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKURETI

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

स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Services
F-15, Pawan, 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;

9

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
7 JUN 2011

M/s. Sun Pharmaceutical Industries
6-9 EPIP, Kartholi, Bari-Brahmana, Jammu,
J & K

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. SPIJ/F/05/13 Dated 21/05/2011 received by this office vide diary no. 24417 dt. 23/05/2011, I enclose licence(s) Nos. BD-918-19179 dated 7 JUN 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/29/11-DC

Copy together with copy of Licence No. BD-918-19179 dated

7 JUN 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-918-19179

Date 7 JUN 2011

1. M/S. Sun Pharmaceutical Industries 6-9 EPIP, Kartholi, Bari-Brahmana, Jammu, J & K is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Symbiotica Speciality Ingredients Sdn. Bhd. #518, Jalan Waja 4, Taman Industri Waja, 09000 Kulim, Kedah, Malaysia 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 06/06/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Tibolone

Item (One) Only

Place: New Delhi

Date: 7 JUN 2011


LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KULKARNI
सहायक औषधि नियन्त्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय

Conditions of Licence

Directorate, General of Health Services
P.O.A. Bihawali, Kolia Road
New Delhi-110 002

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";

(10)

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:

17 JUN 2011

M/s. S.A. Pharmachem Pvt. Ltd.
220, Udyog Bhavan, Sonawala Road, Goregaon(East),
Mumbai-400063

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 09/05/2011 received by this office vide diary no. 24275 dt. 23/05/2011, I enclose licence(s) Nos. BD-674-19180 dated 17 JUN 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/76/06-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-674-19180 dated

17 JUN 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-674-19180

Date 7 JUN 2011

1. M/S. S.A. Pharmachem Pvt. Ltd. 220, Udyog Bhavan, Sonawala Road, Goregaon(East), Mumbai-400063 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Sichuan Neijiang Huixin Pharmaceutical Co. Ltd. No. 188 Linchang Road, Baima Town Nejiang City Sichuan 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 14/06/2011 to 31/07/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Vitamin D2(Ergocalciferol) IP

Item (One) Only

Place: New Delhi

Date: 7 JUN 2011


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

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";

(11)

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
17 JUN 2011

M/s. Kantilal Manilal & Co. Pvt. Ltd.
S.No. 120, Hissa No.2, Pajki, L-2 Gala No. 3&4,
Munisuvrat Complex Phase-II, Rehnal Village, 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 19/05/2011 received by this office vide diary no. 24104 dt. 23/05/2011, I enclose licence(s) Nos. BD-917-19181 dated 17 JUN 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 (I) 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/12/02-DC(Pt-1)(Re-Reg-2010)

Copy together with copy of Licence No. BD-917-19181 dated 17 JUN 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-917-19181

Date 17 JUN 2011

1. M/S. Kantilal Manilal & Co. Pvt. Ltd. S.No. 120, Hissa No.2, Pajki, L-2 Gala No. 3&4, Munisuvrat Complex Phase-II, Rehnal Village, 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. Henan Lihua Pharmaceutical Co. Ltd. Middle Of Huanghe Street, Anyang Hi-Tech Industry Development Zone Henan 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 14/06/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Prednisolone IP

Item (One) Only

Place: New Delhi

Date: 17 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETI

सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Director 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;

(12)

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:

17 JUN 2011

M/s. Kantilal Manilal & Co. Pvt. Ltd.
S.No. 120, Hissa No.2, Pajki, L-2 Gala No. 3&4,
Munisuvrat Complex Phase-II, Rehnal Village, 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 17/05/2011 received by this office vide diary no. 24103 dt. 23/05/2011, I enclose licence(s) Nos. BD-160-19182 dated 7 JUN 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/07/03-DC(Re-Reg-2009)

Copy together with copy of Licence No. BD-160-19182 dated

17 JUN 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-160-19182

Date 17 JUN 2011

1. M/S. Kantilal Manilal & Co. Pvt. Ltd. S.No. 120, Hissa No.2, Pajki, L-2 Gala No. 3&4, Munisuvrat Complex Phase-II, Rehnal Village, 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. Zhejiang Neo-Dankong Pharmaceutical Co. Ltd. 259 Binhai Road, Yantou Ind. Zone, Jiaojiang Dist. Taizhou. 318000 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 14/06/2011 to 31/03/2012 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 : 17 JUN 2011


LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETI
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Services

Conditions of Licence

Department of Health, Government of India
New Delhi-110 002

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";

13

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:

17 JUN 2011

M/s. Kantilal Manilal & Co. Pvt. Ltd.
S.No. 120, Hissa No.2, Pajki, L-2 Gala No. 3&4,
Munisuvrat Complex Phase-II, Rehnal Village, 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 16/05/2011 received by this office vide diary no. 24102 dt. 23/05/2011, I enclose licence(s) Nos. BD-706-19183 dated 17 JUN 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/28/07-DC(Re-Reg-2011)

Copy together with copy of Licence No. BD-706-19183 dated

17 JUN 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-706-19183

Date 07 JUN 2011

1. M/S. Kantilal Manilal & Co. Pvt. Ltd. S.No. 120, Hissa No.2, Paiki, L-2 Gala No. 3&4, Munisuvrat Complex Phase-II, Rehnal Village, 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. Hebei Dongfeng Pharmaceutical Co. Ltd. Address-West Of Yongnian County, Handan City, Hebei, 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 14/06/2011 to 28/02/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Doxycycline Hyclate IP

Item (One) Only

Place: New Delhi

Date: 07 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETI
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Services
F.D.A. Plot No. 10, Connaught Place,
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";

14

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:

17 JUN 2011

M/s. Samarth Lifesciences Pvt.Ltd.
Village-Nangal Uperia-Swarghat Road, Nalagarh,
Disst. Solan-174101

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. SLS/02/05/11 Dated 18/05/2011 received by this office vide diary no. 24361 dt. 23/05/2011, I enclose licence(s) Nos. BD-861-19184 dated 17 JUN 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 (I) 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/92/09-DC

Copy together with copy of Licence No. BD-861-19184 dated

17 JUN 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-861-19184

Date 17 JUN 2011

1. M/S. Samarth Lifesciences Pvt.Ltd. Village-Nangal Uperia-Swarghat Road, Nalagarh, Disst. Solan-174101 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Nanjing Xinbai Pharmaceutical Co. Ltd. No. 68, Xingang Road, Nanjing Economic And Technological Development Zone, 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 14/06/2011 to 31/07/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Heparin Sodium(Non Sterile Bulk)IP

Item (One) Only

Place: New Delhi

Date: 17 JUN 2011


LICENSING AUTHORITY

अरविन्द कुक्रेती
ARVIND KUKRETY

सहायक अवरुधि 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 JUN 2011

M/s. Brooks Laboratories Ltd.
Vill, Kishanpura, Nalagarh Road, Baddi,
Dist. Solan

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 17/05/2011 received by this office vide diary no. 25864 dt. 31/05/2011, I enclose licence(s) Nos. BD-687-19185 dated 15 JUN 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/46/07-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-687-19185 dated

15 JUN 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-687-19185

Date 15 JUN 2011

1. M/S. Brooks Laboratories Ltd. Vill, Kishanpura, Nalagarh Road, Baddi, Dist. Solan is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Shandong New Time Pharmaceutical Co. Ltd. No. 1, North Outer Ring Road, Feixian County, Shandong 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 14/06/2011 to 31/08/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Clavulanate Potassium With Silicon Dioxide USP(1:1)

2. Clavulanate Potassium With Microcrystalline Cellulose USP (1:1)

3. Amoxicillin Sodium and Clavulanate Potassium(5:1) (Bulk Sterile)

Item (Three) Only

Place: New Delhi

Date : 15 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETI
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Services
P. O. Bhawan, Roza 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";