

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

03 FEB 2011

M/s. D.K. Enterprises
No. 41, Raghunayakulu street,
Ground floor,
Chennai- 600003

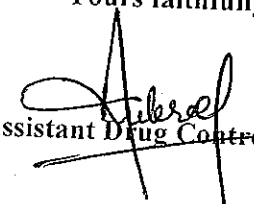
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 03/01/2011 received by this office vide dia. no. 924 dt. 07/01/2011, I enclose licence(s) Nos. BD-231-18445 dated 03 FEB 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)

03 FEB 2011
No. 6-3/BD/21/03-DC (Re-registration 2009)
Copy together with copy of Licence No. BD-231-18445 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-231-18445

Date 03 FEB 2011

1. M/S. D.K.Enterprises No. 41, Raghunayakulu street, Ground floor, Chennai-600003 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Hebei Shengxue Dacheng Pharmaceutical co. ltd., Shengxue road, Luancheng, Shijizuang, 051430, Hebei 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 28/01/2011 to 31/07/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Streptomycin sulphate BP

Item (One) Only

OFFICE COPY

Place: New Delhi

Date : _____

03 FEB 2011

LICENSING AUTHORITY

अरविन्द कुकरेती

ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

स्वास्थ्य सेवा महानिदेशालय

Directorate General of Health Services

F.D.A. Bhawan, Kolla Road

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

03 FEB 2011

M/s. D.K. Enterprises
No. 41, Raghunayakulu street,
Ground floor,
Chennai- 600003

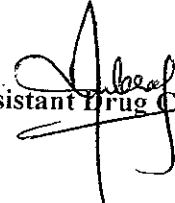
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 06/01/2011 received by this office vide No. 923 dt. 07/01/2011, I enclose licence(s) Nos. BD-890-18446 dated 03 FEB 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)

03 FEB 2011

No. 6-3/BD/09/10-DC

Copy together with copy of Licence No. BD-890-18446 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-890-18446

Date 03 FEB 2011

1. M/S. D.K.Enterprises No. 41, Raghunayakulu street, Ground floor, Chennai-600003 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Hangzhou Tiancheng Pharmaceutical co ltd Hongda road, Qiaonan zone, Hangzhou Qianjiang investment zone, hangzhou, 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 28/01/2011 to 30/11/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Methyldopa IP

Item (One) Only

OFFICE COPY

Place: New Delhi

Date 03 FEB 2011

LICENSING AUTHORITY

ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)
Assistant Drug Controller (India)
स्वास्थ्य सेवा महाविद्यालय

Conditions of Licence

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

No. 6-3/BD/43/04-DC (Re-registration 2011)

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

03 FEB 2011

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

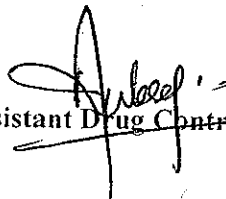
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. ACE/DCGI/13/2010-11 Dated 04/01/2011 received by this office vide diary no. 839 dt. 07/01/2011, I enclose licence(s) Nos. BD-427-18447 dated 03 FEB 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)

03 FEB 2011

No. 6-3/BD/43/04-DC (Re-registration 2011)

Copy together with copy of Licence No. BD-427-18447 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-427-18447

Date 03 FEB 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. Syn-Tech Chem & Pharma co. ltd. 168, kai yuan road, Hsin ying 73055, Taiwan, R.O.C. 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 28/01/2011 to 31/01/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Oxetacaine (Oxethazaine) BP

Item (One) Only

Place: New Delhi

Date: 03 FEB 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY

Assistant Drug (Seal/Stamp)

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

Conditions of Licence

9. 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.
10. 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.
11. The licensee shall be responsible for the business activities of the manufacturer in India along with the registration holder and his authorised agent.
12. 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

03 FEB 2011

M/s. Pioneer Bio-pharma pvt. Ltd.
Godown No. 5 & 6 , Bldg no. 3, Gr. Floor,
Arihant Compound, Purnagaon,
Dist. Thane, 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. PBPPL/C/274/2010-2011 Dated 04/01/2011 received by this office vide diary no. 838 dt. 07/01/2011, I enclose licence(s) Nos. BD-561-18448 dated 03 FEB 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,

03 FEB 2011

Assistant Drugs Controller (India)

No. 6-3/BD/10/05-DC (Re-registration 2008)

Copy together with copy of Licence No. BD-561-18448 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-561-18448

Date 03 FEB 2011

1. M/S. Pioneer Bio-pharma pvt. Ltd. Godown No. 5 & 6 , Bldg no. 3, Gr. Floor, Arihant Compound, Purnagaon, Dist. Thane, 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 Medicine Co. Ltd., Xinchang pharmaceutical factory, 98, east xinchang, dadao road, Xinchang, 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 28/01/2011 to 31/03/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported


D-Biotin USP

Item (One) Only

Place: New Delhi

Date: 03 FEB 2011

OFFICE COPY


LICENSING AUTHORITY
अरविन्द कुकर्णी
ARVIND KULKARNI
सहायक संचालक (आयुर्विज्ञान सेवाएँ)
Assistant Director (Health Services)
स्वास्थ्य सेवा महानिदेशक
Directorate General of Health Services
F.D.A. Bhawan, Kofla 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

03 FEB 2011

M/s. Kawarlal & Co.
No. 27, Raghunayakulu street,
Chennai- 600003

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

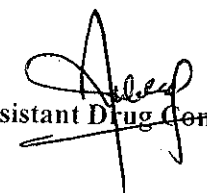
Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 06/01/2011 received by this office vide diary no. 925 dt. 07/01/2011, I enclose licence(s) Nos. BD-890-18449 dated 03 FEB 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,

03 FEB 2011


Assistant Drug Controller (India)

No. 6-3/BD/09/10-DC

Copy together with copy of Licence No. BD-890-18449 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-890-18449

Date 03 FEB 2011

1. M/S. Kawarlal & Co. No. 27, Raghunayakulu street, Chennai- 600003 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Hangzhou Tiancheng Pharmaceutical co ltd Hongda road, Qiaonan zone, Hangzhou Qianjiang investment zone, hangzhou, 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 28/01/2011 to 30/11/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Methyldopa IP

Item (One) Only

Place: New Delhi

Date : 03 FEB 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY
सहायक औषधि नियंत्रक (भारत)
Assistant Controller (India)
स्वास्थ्य सेवा महाविद्यालय
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
03 FEB 2011

M/s. Ranbaxy Laboratories Ltd.
Village Ganuwala, tehsil Paonta sahib,
Dist. Sirmour- 173 025 (H.P.)

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 06/01/2011 received by this office vide diary no. 942 dt. 07/01/2011, I enclose licence(s) Nos. BD-147-18450 dated 03 FEB 2011. The said 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)

03 FEB 2011

No. 6-3/BD/174/02-DC (Re-registration 2010)
Copy together with copy of Licence No. BD-147-18450 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.

No. 6-3/BD/194/03-DC (Re-registration 2010)

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
03 FEB 2011

M/s. Parabolic Drugs Ltd.
Village Sundhran, P.O. Mubarakpur,
Tehsil Dera Bassi,
District Mohali

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 23/12/2010 received by this office vide diary no. 915 dt. 07/01/2011, I enclose licence(s) Nos. BD-370-18451 dated **03 FEB 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 (h) 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,

03 FEB 2011


Assistant Drug Controller (India)

No. 6-3/BD/194/03-DC (Re-registration 2010)

Copy together with copy of Licence No. BD-370-18451 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-370-18451

Date 03 FEB 2011

1. M/S. Parabolic Drugs Ltd. Village Sundhran, P.O. Mubarakpur, Tehsil Dera Bassi, District Mohali is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Harbin Pharmaceutical Group Co. Ltd. General Pharma Factory, No. 109 Xuefu Road, Nangang Dist. Harbin, 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 28/01/2011 to 28/02/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported


Ceftriaxone Sodium (Bulk Non-Sterile) IP

Item (One) Only

Place: New Delhi

Date: 03 FEB 2011

OFFICE COPY


LICENSING AUTHORITY
Assistant Director (IP)
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

03 FEB 2011

M/s. Parabolic Drugs Ltd.
Village Sundhran, P.O. Mubarakpur,
Tehsil Dera Bassi,
District Mohali

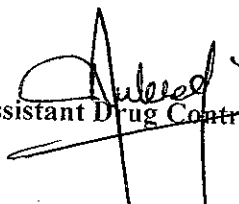
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 05/01/2011 received by this office vide diary no. 1292 dt. 10/01/2011, I enclose licence(s) Nos. BD-756-18452 dated **03 FEB 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/05/06-DC(Pt-1)

03 FEB 2011

Copy together with copy of Licence No. BD-756-18452 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-756-18452

Date 03 FEB 2011

1. M/S. Parabolic Drugs Ltd. Village Sundhran, P.O. Mubarakpur, Tehsil Dera Bassi, District Mohali is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. CSPC Ouyi Pharmaceutical Co. Ltd. 276, Zhongshan West Road, Shijiazhuang 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 28/01/2011 to 30/09/2011 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Azithromycin IP

Item (One) Only

Place: New Delhi

Date: 03 FEB 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY

सहायक निदेशक (आर.डी.)

Assistant Director (India)

स्वास्थ्य सेवा विभाग

Conditions of Licence Directorate General of Health Services
F.D.A. Bhawan, Kofa 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

03 FEB 2011

M/s. Essix Biosciences Ltd.
B4 & 5 Focal Point Dera Bassi District
S.A.S. Nagar
Mohali

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 29/12/2010 received by this office vide entry no. 1052 dt. 10/01/2011, I enclose licence(s) Nos. BD-658-18453 and BD-658-18454 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,

03 FEB 2011

Assistant Drug Controller (India)

No. 6-3/BD/78/06-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-658-18453 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-658-18453

Date 03 FEB 2011

- M/S. Essix Biosciences Ltd. B4 & 5 Focal Point Dera Bassi District S.A.S. Nagar Mohali is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Hec Biochem Co. Ltd., 62, Binjiang Road, Yidu City, Hebei Province 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 28/01/2011 to 30/04/2013 unless it is sooner suspended or cancelled under the said rules.


Names of drugs to be imported

1. Erythromycin IP
2. Erythromycin Thioamylate
Item (Two) Only

OFFICE COPY

Place: New Delhi

Date : 03 FEB 2011


LICENSING AUTHORITY
ARVIND KULKRETY
सहायक औषधि नियंत्रक (भारत)
Assistant Drug Controller (India)
एवमरुध्य सेवक (भारत)
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
03 FEB 2011

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

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

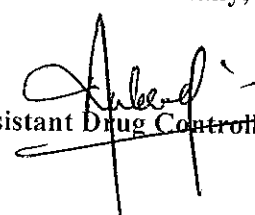
Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 06/01/2011 received by this office vide diary no. 1250 dt. 10/01/2011, I enclose licence(s) Nos. BD-639-18454 dated **03 FEB 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,

03 FEB 2011


Assistant Drug Controller (India)

No. 6-3/BD/81/06-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-639-18454 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-639-18454

Date 03 FEB 2011

1. M/S. Pradipkumar Pharma Pvt. Ltd. Gala No. 2 Gr Floor, Munisuvrat Complex, Phase-II, Bldg. No. 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. Guangzhao Baiyunshan Pharmaceutical Co. Ltd. No. 78, Tongbao Road, Tonghe Street, Baiyun District, Guangzhou, 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 28/01/2011 to 31/01/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

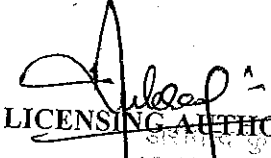
Annexure-1

Item (Six) Only

OFFICE COPY

Place: New Delhi

Date: 03 FEB 2011


LICENSING AUTHORITY
ARVIND HUKRETY
सहायक डी.डी. नियंत्रक (भारत)
Assistant Drug Controller (India)
स्वास्थ्य सेवा विभाग, दिल्ली
Seal/Stamp
Directorate General of Health Services
F.D.A. Bhawan, Kotla Road
New Delhi - 110 002

Conditions of Licence

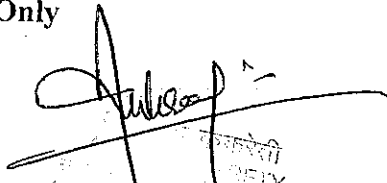
1. A photocopy of licence shall be displayed in a prominent place in a part of the premises, and the original licence shall be produced, whenever required.
2. Each batch of drug imported into India shall be accompanied with a detailed batch test report and a batch release certificate, duly signed and 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";

ANNEXURE-1 (List of Products of Form 10 License No. BD-639-18454)

1. Ceftriaxone Sodium IP (Bulk Sterile)
2. Ceftazidime IP (Bulk Sterile)
3. Cefuroxime Sodium IP (Bulk Sterile)
4. Cefprozil USP
5. Cefixime USP
6. Cefotaxime Sodium IP (Bulk Sterile)

03 FEB 2011

Item (Six) Only


DIRECTOR
GENERAL
INDIA)
Directorate of Pharmaceutical Health Services
F-20, Connaught Place, Kirti Road
New Delhi-110 002

OFFICE COPY

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

03 FEB 2011

M/s. Harish Enterprises
Shop No. 1, 1st Floor, Survey No. 145/3,
Ausadh Compound, Inside Dalmill Comp. Purna,
Bhiwandi

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 05/01/2011 received by this office vide diary no. 1251 dt. 10/01/2011, I enclose licence(s) Nos. BD-404-18455 dated **03 FEB 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,

03 FEB 2011


Assistant Drug Controller (India)

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

Copy together with copy of Licence No. BD-404-18455 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-404-18455

Date 03 FEB 2011

1. M/S. Harish Enterprises Shop No. 1, 1st Floor, Survey No. 145/3, Ausadh Compound, Inside Dalmill Comp. Purna, Bhiwandi 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, Chunyuan West 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 28/01/2011 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: 03 FEB 2011

LICENSING AUTHORITY

ARVIND KUKRETY

सहायक दवा वि नियंत्रक (भारत)

Assistant Drugs Controller (India)

सहायक दवा वि नियंत्रक

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

03 FEB 2011

M/s. Ipca Laboratories Ltd.
Regd. Off. 48 Kandivli Industrial Estate,
Kandivli (W)
Mumbai-400067

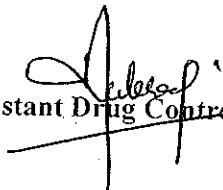
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 28/12/2010 received by this office vide diary no. 1219 dt. 10/01/2011, I enclose licence(s) Nos. BD-78-18456 dated 03 FEB 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/105/02-DC

03 FEB 2011

Copy together with copy of Licence No. BD-78-18456 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-78-18456

Date 03 FEB 2011

1. M/S. Ipca Laboratories Ltd. Regd. Off. 48 Kandivli Industrial Estate, Kandivli (W) Mumbai-400067 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Cilag AG, Hochstrasse 201, CH 8205 Schaffhausen, Switzerland 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 28/01/2011 to 31/12/2011 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Mefloquine Hydrochloride Ph

Item (One) Only

OFFICE COPY

Place: New Delhi

Date: 03 FEB 2011

LICENSING AUTHORITY

ARVIND KUKRETY

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

रक्षाध्यक्ष सेना, दिल्ली

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
03 FEB 2011

M/s. Indoco Remedies Ltd.
L-14, Verna Industrial Estate, Verna Salcete
Goa.

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

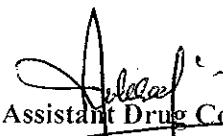
Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. RMK/IND-IMP/DCGI/100 Dated 04/01/2011 received by this office vide diary no. 1126 dt. 10/01/2011, I enclose licence(s) Nos. BD-13-18457 dated **03 FEB 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,

03 FEB 2011


Assistant Drug Controller (India)

No. 6-3/BD/20/02-DC(Pt-1) (Re-Reg-2008)

Copy together with copy of Licence No. BD-13-18457 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-13-18457

Date 03 FEB 2011

1. M/S. Indoco Remedies Ltd. L-14, Verna Industrial Estate, Verna Salcete Goa.
is hereby licensed into India during the period for which this licence is in force, the
drugs specified below, manufactured By M/s. Basf Se, Carl- Bosch-Strasse 38,
67056 Ludwigshafen, Germany having Premises at M/s. Basf
Pharmachemikalien GMBH & Co.KG. Plant Minden, Karlstrabe 15-39, 42-44,
32423 Minden Federal Republic of Germany 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 28/01/2011 to 31/12/2011 unless it is sooner
suspended or cancelled under the said rules.

Names of drugs to be imported

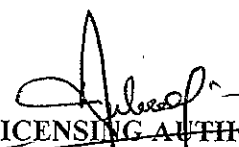
Caffeine Anhydrous

Item (One) Only

Place: New Delhi

Date : 03 FEB 2011

OFFICE COPY


LICENSING AUTHORITY
अरविन्द कुकरेती
ARVIND KUKRETY
सहायक औषधि नियंत्रक (भारत)
Assistant Director (India)
स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Serv.
F.D.A. Bhawan, Kolla Road
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:

03 FEB 2011

M/s. Kem Organics
Arihant Complex, Near Koper Bus Stop,
Bldg. No. 2, Godown No. 6, Purna Village
Bhivandi

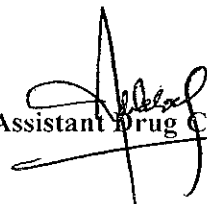
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 05/01/2011 received by this office vide diary no. 1014 dt. 10/01/2011, I enclose licence(s) Nos. BD-160-18458 dated 03 FEB 2011 as/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)

03 FEB 2011

No. 6-3/BD/07/03-DC (Re-Reg-2009)

Copy together with copy of Licence No. BD-160-18458 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-160-18458

Date 03 FEB 2011

1. M/S. Kem Organics Arihant Complex, Near Koper Bus Stop, Bldg. No. 2, Godown No. 6, Purna 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 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 28/01/2011 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: 03 FEB 2011

OFFICE COPY

LICENSING AUTHORITY
ARVIND KUKRETY

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

Seal/Stamp
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

New Delhi, dated

To:

03 FEB 2011

M/s. Param Pharma
2/12, Morarji Velji Building, 7-A,
Dr. M.B. Velkar Street, Kalbadevi Road
Mumbai-400002

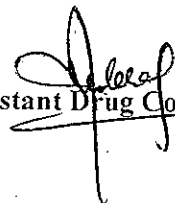
Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 05/01/2011 received by this office vide diary no. 1253 dt. 10/01/2011, I enclose licence(s) Nos. BD-694-18459 dated 03 FEB 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)

03 FEB 2011

No. 6-3/BD/17/07-DC (Re-Reg-2010)

Copy together with copy of Licence No. BD-694-18459 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-694-18459

Date 03 FEB 2011

1. M/S. Param Pharma 2/12, Morarji Velji Building, 7-A, Dr. M.B. Velkar Street, 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. Minsheng Group Shaoxing Pharmaceutical Co. Ltd. 315, Tanggong Road, Paojiang Industrial Zone Shaoxing Zhejiang 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 28/01/2011 to 30/11/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Furazolidone IP

Item (One) Only

Place: New Delhi

Date : 03 FEB 2011

OFFICE COPY
LICENSING AUTHORITY
रजिस्ट्रार जीवधि नियंत्रक (भारत)
Registrar of Drugs Controller (India)
रजिस्ट्रार सेवा महाविद्यालय
Directorate of Health Services
Seal/Stamp
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

03 FEB 2011

M/s. Medi Pharma Drug House
Bulakhidas Bldg, Gr. Floor, 13,
Vithaladas Road, Princess Street,
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. FM/49/10 Dated 05/01/2011 received by this office vide diary no. 1200 dt. 10/01/2011, I enclose licence(s) Nos. BD-825-18460 dated 03 FEB 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 to the effect 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)

03 FEB 2011

No. 6-3/BD/86/09-DC

Copy together with copy of Licence No. BD-825-18460 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-825-18460

Date 03 FEB 2011

1. M/S. Medi Pharma Drug House Bulakhidas Bldg, Gr. Floor, 13, Vithaladas Road, Princess Street, 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 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 28/01/2011 to 31/01/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. Ofloxacin IP

2. Simvastatin USP

Item (Two) Only

Place: New Delhi

Date: 03 FEB 2011

LICENSING AUTHORITY

अरविन्द कुकरेती
ARVIND KUKRETY
सहायक औषधि नियंत्रक (भारत)
Assistant Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Directorate General of Health Services
5-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";