

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

**27 JUN 2011**

M/s. Wockhardt Limited  
Plot No. C-2, Wockhardt Towers  
Bandra Kurla Complex Bandra(E)  
Mumbai-400051

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

Dear Sir/Sirs,

~~With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/05/2011 received by this office vide diary no. 25998 dt. 01/06/2011, I enclose licence(s) Nos. BD-350-19241 dated 27 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/146/03-DC(Pt-1) (Re-Reg-2010)

Copy together with copy of Licence No. BD-350-19241 dated

**27 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-350-19241

Date 27 JUN 2011

1. M/S. Wockhardt Limited Plot No. C-2, Wockhardt Towers Bandra Kurla Complex Bandra(E) Mumbai-400051 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Kyowa Hakko Bio Co. Ltd. Hofu Plant, 1-1, Kyowa-Cho, Hofu, Yamaguchi, Japan 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 22/06/2011 to 31/01/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Annexure- 1

Item (Eighteen) Only

Place: New Delhi

Date 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुक्रेती  
ARVIND KUKRETY  
सहायक उपायुक्त (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय

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";

Seal/Stamp  
Directorate General of Health Services  
Shayan, Kolla Road  
New Delhi - 110 002

**ANNEXURE-1 (List of Products of Form 10 License No. BD-350-19241)**

1. L-Cysteine Hydrochloride
2. Glycine
3. L-Methionine
4. L-Tyrosine
5. L-Alanine
6. L-Arginine Hydrochloride
7. L-Aspartic Acid
8. L-Histidine Hydrochloride H<sub>2</sub>O
9. L-Isoleucine
10. L-Leucine
11. L-Lysine Hydrochloride
12. L-Phenyl Alanine
13. L- Threonine
14. L-Tryptophan
15. L-valine
16. L-Asparagine
17. L-Glutamic Acid
18. L- Alanyl-L-Glutamine

Item (Eighteen) Only

**OFFICE COPY**

*[Signature]*

अरविन्द कुकरेती  
ARVIND KUKRETY  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
F.D.A. Bhawan, Kotta Road  
New Delhi - 110 002

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

27 JUN 2011

M/s. Wockhardt Limited  
Plot No. C-2, Wockhardt Towers  
Bandra Kurla Complex Bandra(E)  
Mumbai-400051

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

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/05/2011 received by this office vide diary no. 25999 dt. 01/06/2011, I enclose licence(s) Nos. BD-351-19242 dated 27 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/147/03-DC(Pt-1) (Re-Reg-2010)

Copy together with copy of Licence No. BD-351-19242 dated

27 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-351-19242

Date 27 JUN 2011

1. M/S. Wockhardt Limited Plot No. C-2, Wockhardt Towers Bandra Kurla Complex Bandra(E) Mumbai-400051 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Kyowa Hakko Bio Co. Ltd. UBE Plant, 2548, Fujimagari, UBE, Yamaguchi, Japan 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 22/06/2011 to 31/1/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

1. L-Proline
2. L-Glutamine
3. L-Serine

Item (Three) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती  
ARVIND KUKRETI  
सहायक औषधि निरीक्षक (महाराष्ट्र)  
Assistant Drugs Controller (Maharashtra)

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

Conditions of Licence

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

From:

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

New Delhi, dated

**27 JUN 2011**

To:

M/s. Yeluri Formulations Pvt. Ltd.  
Sy. No. 296/7/6/, IDA, Bollaram,  
Medak 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. Nil Dated ~~02/05/2011~~ received by this office vide diary no. 25956 dt. 01/06/2011, I enclose licence(s) Nos. BD-582-19246 dated **27 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/07/09-DC(Re-Reg-2009

Copy together with copy of Licence No. BD-582-19246 dated **27 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-582-19246

Date 27 JUN 2011

1. M/S. Yeluri Formulations Pvt. Ltd. Sy. No. 296/7/6/, IDA, Bollaram, Medak Dist. is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Nexchem Pharmaceutical Co. Ltd. Bailongqiao Industrial District, Jinhua City, Zhejiang-321025 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 22/06/2011 to 31/03/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Meropenem Buffered (Sterile) USP

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरती  
ARVIND KUKRETY  
सहायक (संयोजक) (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:

27 JUN 2011

M/s. Allergan India Pvt. Ltd.  
KSEF Bldg. opp. Kims Hospital,  
Kondoji Basappa Road Kalasipalyam  
Bangalore-560002

**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 10/05/2011 received by this office vide diary no. 26350 dt. 02/06/2011, I enclose licence(s) Nos. FF-120-19243 dated 27 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. 6-4/FF/124/08-DC(RE-REG-2008)

Copy together with copy of Licence No. FF-120-19243 dated

27 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.



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  
**27 JUN 2011**

M/s. Getwell Pharmaceutical  
474 Udyog Vihar, Phase-V Gurgaon  
Haryana

**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 ~~09/06/2011~~ received by this office vide diary no. ~~29234~~ dt. ~~17/06/2011~~, I enclose licence(s) Nos. BD-865-19244 dated **27 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/87/09-DC

Copy together with copy of Licence No. BD-865-19244 dated

**27 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-865-19244

Date 27 JUN 2011

1. M/S. Getwell Pharmaceutical 474 Udyog Vihar, Phase-v Gurgaon Haryana is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Jiangsu Hengrui Medicine Co. Ltd. Jinqiao Road, Dapu Industrial Park Economic & Technological Development Zone, Lianyungang, Jiangsu 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 22/06/2011 to 31/07/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

**Oxaliplatin BP**

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

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

Directorate General of Health Services  
F.O.A. Bhambhata  
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  
**27 JUN 2011**

M/s. Astellas Pharma India Pvt. Ltd.  
505 Medows, Sahar Plaza Complex,  
J.B. Nagar Andheri Kurla Road,  
Mumbai 400059

**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 09/06/2011 received by this office vide diary no. 27674 dt. 09/06/2011, I enclose licence(s) Nos. FF-584-19245 dated 27 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. 6-4/FF/57/2010-DC

Copy together with copy of Licence No. FF-584-19245 dated **27 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 FF-584-19245

Date 27 JUN 2011

1. M/S. Astellas Pharma India Pvt. Ltd. 505 Medows, Sahar Plaza Complex, J.B. Nagar Andheri Kurla Road, Mumbai 400059 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Astellas Pharma Europe B.V. Hogemaat 2,7942 JG Meppel The Netherlands 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 22/06/2011 to 31/03/2014 unless it is sooner suspended or cancelled under the said rules.

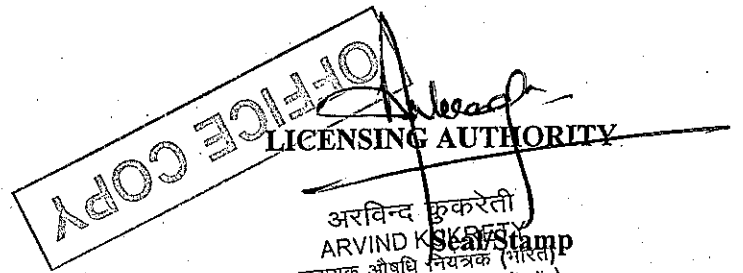
Names of drugs to be imported

Solifenacin Succinate 5 mg & 10 mg Film coated tablets

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011



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

**27 JUN 2011**

M/s. United Biotech Pvt. Ltd.  
Vill. Bagbania, Baddi- Nalagarh Road,  
Dist. Solan, 174101(H.P)

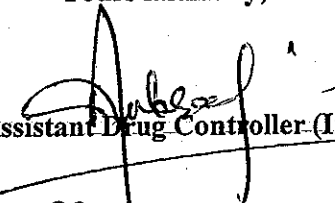
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. UBPL/2011-12/CDSO/SK/005 Dated 10/06/2011 received by this office vide diary no. 27842 dt. 10/06/2011, I enclose licence(s) Nos. BD-644-19247 dated **27 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/87/06-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-644-19247 dated

**27 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-644-19247

Date 27 JUN 2011

1. M/S. United Biotech Pvt. Ltd. Vill. Bagbania, Baddi- Nalagarh Road, Dist. Solan, 174101(H.P) is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Biogene Co. Ltd. 307-11, Songsan-RI, Yangkam-Myun Hwaseong-SI, Gyeonggi-Do, Korea 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 22/06/2011 to 28/02/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Teicoplanin-JP(Bulk Non Sterile)

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

LICENSING AUTHORITY

अरविन्द कुकरेती

ARVIND KUKRETI

सहायक औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

ए. डी. ए. भवान, कोला रोड

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 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

**27 JUN 2011**

M/s. United Biotech Pvt. Ltd.  
Vill. Bagbania, Baddi- Nalagarh Road,  
Dist. Solan, 174101(H.P)

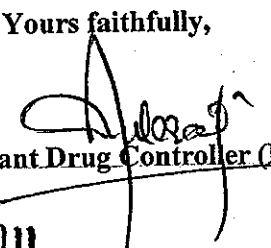
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. UBPL/2011-12/CDSKO/SK/007 Dated 10/06/2011 received by this office vide diary no. 27840 dt. 10/06/2011, I enclose licence(s) Nos. BD-35-19248 dated 27 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/48/02-DC(Pt-1) (Re-Reg-2008)

Copy together with copy of Licence No. BD-35-19248 dated **27 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-35-19248

Date 27 JUN 2011

1. M/S. United Biotech Pvt. Ltd. Vill. Bagbania, Baddi- Nalagarh Road, Dist. Solan, 174101(H.P) is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Crystal Pharma S.A. Parque Tecnologico- Parcela, 105, 47151, Boecillo (Valladolid) Spain 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 22/06/2011 to 31/12/2011 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Hydrocortisone Sodium Succinate Buffered 5%, Sterile USP

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

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";

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  
**27 JUN 2011**

M/s. United Biotech Pvt. Ltd.  
Vill. Bagbania, Baddi- Nalagarh Road,  
Dist. Solan, 174101(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. UBPL/2011-12/CDSCO/SK/006 Dated 10/06/2011 received by this office vide diary no. 27841 dt. 10/06/2011, I enclose licence(s) Nos. BD-392-19249 dated **27 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/30/04-DC (Re-Reg-2011)

Copy together with copy of Licence No. BD-392-19249 dated

**27 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-392-19249

Date 27 JUN 2011

1. M/S. United Biotech Pvt. Ltd. Vill. Bagbania, Baddi- Nalagarh Road, Dist. Solan, 174101(H.P) is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Nantong Jinghua Pharmaceutical Co. Ltd., 43, Yaogang Road , Nantong , Jiangsu, China-226006 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 22/06/2011 to 30/09/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

**5-Fluorouracil IP**

Item (One) Only

Place: New Delhi

Date : 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

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

Conditions of Licence

Directorate General of Health Services  
P.O. Bawana, Indraprastha  
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

New Delhi, dated

To:

27 JUN 2011

M/s. United Biotech Pvt. Ltd.  
Vill. Bagbania, Baddi- Nalagarh Road,  
Dist. Solan, 174101(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. UBPL/2011-12/CDSCO/SK/008 Dated 10/06/2011 received by this office vide diary no. 27839 dt. 10/06/2011, I enclose licence(s) Nos. BD-882-19250 dated 27 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/73/09-DC

Copy together with copy of Licence No. BD-882-19250 dated

27 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-882-19250

Date 27 JUN 2011

1. M/S. United Biotech Pvt. Ltd. Vill. Bagbania, Baddi- Nalagarh Road, Dist. Solan, 174101(H.P) is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Shanghai Techwell Biopharmaceutical Co. Ltd. No. 4258, jindu Road, Minhang District Shanghai-201108 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 22/06/2011 to 31/01/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

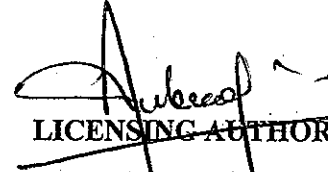
1. Chorionic Gonadotrophin (HCG)IP
2. Menotropins-Human Menopausal Gonadotrophin (HMG)USP
3. Urofollitropin (FSH)

Item (Three) Only

Place: New Delhi

Date : 27 JUN 2011

OFFICE COPY

  
LICENSING AUTHORITY

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

Conditions of Licence

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

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

27 JUN 2011

M/s. Covidien Healthcare India Pvt. Ltd.  
No. 156, Doshi Towers 6th Floor,  
Poonamallee High Road, Kilpauk  
Chennai-600010

**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. CHIL/IRA/RB/F-10/FF-581/2011 Dated 06/05/2011 received by this office vide diary no. 21628 dt. 06/05/2011, I enclose licence(s) Nos. FF-581-19251 dated 27 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. 6-4/FF/13/09-DC (Re-Reg-2009)

Copy together with copy of Licence No. FF-581-19251 dated

27 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 FF-581-19251

Date 27 JUN 2011

1. M/S. Covidien Healthcare India Pvt. Ltd. No. 156, Doshi Towers 6th Floor, Poonamallee High Road, Kilpauk Chennai-600010 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Tyco Healthcare 7500 Trans Canada Highway, Pointclaire, Quebec, Canada H9R 5H8 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 22/06/2011 to 31/03/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

**Optiray (Ioversol Injections 300, 320, 350)**

**IN Glass Vial, Glass Bottle and Syringe Preparations**

**Item (One) Only**

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

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

**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:

**27 JUN 2011**

M/s. Yeluri Formulations Pvt. Ltd.  
Sy. No. 296/7/6/, IDA, Bollaram,  
Medak 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. Nil Dated ~~31/05/2011~~ received by this office vide diary no. 25955 dt. 01/06/2011, I enclose licence(s) Nos. BD-593-19252 dated 27 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/06/06-DC (Re-Reg-2009)

Copy together with copy of Licence No. BD-593-19252 dated **27 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-593-19252

Date 27 JUN 2011

1. M/S. Yeluri Formulations Pvt. Ltd. Sy. No. 296/7/6/, IDA, Bollaram, Medak Dist. is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. NCPG Shanxi Guardian Pharmaceuticals Co. Ltd. No. 1, Huagong Road, Jinyuan Distict, Taiyuan, Shanxi 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 22/06/2011 to 30/04/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Piperacillin Sodium/Tazobactam Sodium (8:1)

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

LICENSING AUTHORITY

अरविन्द कुकरेती  
ARVIND KUKRETI

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

स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services

P.O. Arvind Kukreti  
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:

27 JUN 2011

M/s. M.J. Biopharm Pvt. Ltd.  
Plot No. L-7, M.I.D.C. Industrial Area, Taloja, Navi Mumbai  
Dist. Raigad

**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 27/04/2011 received by this office vide diary no. 20895 dt. 04/05/2011, I enclose licence(s) Nos. BD-710-19253 dated 27 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/06/07-DC (Re-Reg-2011)

Copy together with copy of Licence No. BD-710-19253 dated 27 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-710-19253

Date 27 JUN 2011

1. M/S. M.J. Biopharm Pvt. Ltd. Plot No. L-7, M.I.D.C. Industrial Area, Taloja, Navi Mumbai Dist. Raigad 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 22/06/2011 to 31/03/2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

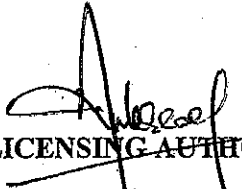
**Prednisolone Sodium Phosphate. IP**

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

  
LICENSING AUTHORITY  
अरविन्द कुक्रेती  
ARVIND KUKRETI  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Health Services  
F-10, Bhawan, Kailash 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:

27 JUN 2011

M/s. Novartis Healthcare Pvt. Ltd.  
Gala No. 1-A & 2-A, Bldg No. 28,  
Arihant Comp. Kopar, Purna 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 01.06.2011 received by this office vide diary no. 27727 dt. 03.06.2011, I enclose licence(s) Nos. FF-463-12536 dated 27 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. 6-4/FF/53/07-DC (Re-Registration-2011)

Copy together with copy of Licence No. FF-463-12536 dated 27 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 FF-463-12536

Date 27 JUN 2011

1. M/S. Novartis Healthcare Pvt. Ltd. Gala No. 1-A & 2-A, Bldg No. 28, Arihant Comp. Kopar, Purna 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. Novartis Pharma Produktions GMBH, Oeflingerstrasse 44 D-79664 Wehr, 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 22-06-2011 to 31-05-2014 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

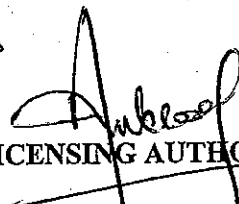
Elidel (Pimecrolimus) Cream 1%

Item (One) Only

Place: New Delhi

Date: 27 JUN 2011

OFFICE COPY

  
LICENSING AUTHORITY

अरविन्द कुकरेती  
ARVIND KUKARETI  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय  
Directorate General of Medicines and  
P.D.A. Bhawan, Kola Road  
New Delhi-110 002

Conditions of Licence

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