

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  
06 MAY 2011

M/s. ABS Mercantiles Pvt. Ltd.  
Shop No.1, Saket Commercial Complex, 56/9  
New Railway Road  
Gurgaon

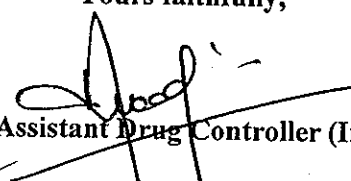
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 08/04/2011 received by this office vide diary no. 17138 dt. 11/04/2011, I enclose licence(s) Nos. BD-463-18978 dated 06 MAY 2011. These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

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

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/91/04-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-463-18978 dated

06 MAY 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-463-18978

Date 06 MAY 2011

1. M/S. ABS Mercantiles Pvt. Ltd. Shop No.1, Saket Commercial Complex, 56/9  
New Railway Road/ Gurgaon is hereby licensed into India during the period for  
which this licence is in force, the drugs specified below, manufactured By M/s.  
Hunan Dongting Pharmaceutical Co. Ltd., No. 16, Dongyan Road, Deshan,  
Changde City, Hunan Province , PC 415001, 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 04/05/2011 to 31/08/2013 unless it is sooner  
suspended or cancelled under the said rules.

Names of drugs to be imported

Tranexamic Acid BP

Item (One) Only

Place: New Delhi

Date: 06 MAY 2011

  
LICENSING AUTHORITY

अरविंद कुक्रेती  
ARVIND KUKRETY

अधीनस्थ अधिकारी, दिल्ली (भारत)

अधीनस्थ अधिकारी, दिल्ली (भारत)

अधीनस्थ अधिकारी, दिल्ली (भारत)

अधीनस्थ अधिकारी, दिल्ली (भारत)

अधीनस्थ अधिकारी, दिल्ली (भारत)

Seal/Stamp

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";

2

om:

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

To:

New Delhi, dated

06 MAY 2011

M/s. Gorang International  
Bldg No. 1, Gala No. 12, 2nd Floor  
Manish Compound, Anjurphata,  
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. Nil Dated 06/04/2011 received by this office vide diary no. 16870 dt. 11/04/2011, I enclose licence(s) Nos. BD-463-18979 dated 06 MAY 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/91/04-DC(Re-Reg-2010)

Copy together with copy of Licence No. BD-463-18979 dated 06 MAY 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-463-18979

Date 06 MAY 2011

1. M/S. Gorang International Bldg No. 1, Gala No. 12, 2nd Floor Manish Compound, Anjurphata, 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. Hunan Dongting Pharmaceutical Co. Ltd., No. 16, Dongyan Road, Deshan, Changde City, Hunan Province , PC 415001, 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 04/05/2011 to 31/08/2013 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Tranexamic Acid BP

Item (One) Only

Place: New Delhi

Date 06 MAY 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY

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

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

Conditions of Licence

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

From:

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

To:

New Delhi, dated  
06 MAY 2011

M/s. Twenty First Century Pharmaceuticals Pvt. Ltd.(Unit-II)  
Khasra No. 282, Nalheri Dehviran,  
Puhana-Iqbalpur Road, Roorkee,  
Distt. Haridwar, Uttarakhand

**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 25/04/2011 received by this office vide diary no. 19791 dt. 27/04/2011, I enclose licence(s) Nos. BD-198-18980 dated 06 MAY 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/52/02-DC(Re-Reg-2009)

Copy together with copy of Licence No. BD-198-18980 dated 06 MAY 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-198-18980

Date 06 MAY 2011

1. M/S. Twenty First Century Pharmaceuticals Pvt. Ltd.(Unit-II) Khasra No. 282, Nalheri Dehviran, Puhana-Iqbalpur Road, Roorkee, Distt. Haridwar, Uttarakhand is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. CKD Bio Corporation, 454, Moknae-dong, Danwon-GU, Ansan- SI, Kyunggi -DO, 425-100, South 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 04/05/2011 to 31/05/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

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

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

Item (Two) Only

Place: New Delhi

Date: 06 MAY 2011

LICENSING AUTHORITY

अरविन्द कुकरेली  
ARVIND KUKRETY  
सहायक औषधि  
Seal/Stamp

Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय

Directorate General of Health Services  
Department of Health, Government of India  
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

06 MAY 2011

M/s. Twenty First Century Pharmaceuticals Pvt. Ltd.  
No. 360, SIDCO Estate,  
Chennai-600098

**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 25/04/2011 received by this office vide no. 19790 dt. 27/04/2011, I enclose licence(s) Nos. BD-198-18981 dated 06 MAY 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/52/02-DC(Re-Reg-2009)

Copy together with copy of Licence No. BD-198-18981 dated 06 MAY 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-198-18981

Date 06 MAY 2011

1. M/S. Twenty First Century Pharmaceuticals Pvt. Ltd. No. 360, SIDCO Estate, Chennai-600098 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. CKD Bio Corporation , 454, Moknae-dong, Danwon-GU, Ansan- SI, Kyunggi -DO, 425-100, South 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 04/05/2011 to 31/05/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

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

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

Item (Two) Only

Place: New Delhi

Date: 06 MAY 2011

**LICENSING AUTHORITY**

अरविन्द कुक्रेती

ARVIND KUKRETI

सहायक औषधि नियंत्रक (भारत)

Assistant Drug 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

06 MAY 2011

M/s. VHB Medi Sciences Ltd.  
Plot No. 20-22, 49-51, I.I.E, Sector-5, Pant Nagar,  
Distt. Udham Singh Nagar

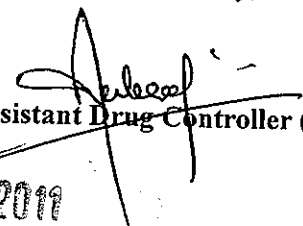
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 21/03/2011 received by this office with ~~reference~~ no. 13628 dt. 23/03/2011, I enclose licence(s) Nos. BD-882-18982 dated 06 MAY 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-882-18982 dated 06 MAY 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-882-18982

Date 06 MAY 2011

1. M/S. VHB Medi Sciences Ltd. Plot No. 20-22, 49-51, I.I.E, Sector-5, Pant Nagar, Dist. Udham Singh Nagar 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 04/05/2011 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: 06 MAY 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, Kotla Road  
New Delhi

Conditions of Licence

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

From:

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

To:

New Delhi, dated

06 MAY 2011

M/s. Dilipkumar & Co.  
606, 6th Floor, Arun Chambers, Tardeo Road,  
Mumbai-400034

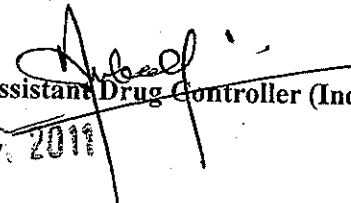
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 24/03/2011 received by this office vide diary no. 0 dt. 24/03/2011, I enclose licence(s) Nos. BD-574-18983 dated 06 MAY 2011. These licence(s) has/have been under the Drugs Act, 1940 and Rules thereunder.

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

Yours faithfully,

  
Assistant Drug Controller (India)

No. 6-3/BD/40/05-DC(Pt-1) (Re-registration 2009)

Copy together with copy of Licence No. BD-574-18983 dated 06 MAY 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-574-18983

Date 06 MAY, 2011

1. M/S. Dilipkumar & Co. 606, 6th Floor, Arun Chambers, Tardeo Road, Mumbai-400034 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Jiangsu Tohope Pharmaceutical Co. Ltd. 188, Wuyishan Road, Southeast Economic Development Zone, Changshu City, Jiangsu 215533, 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 04/05/2011 to 14/02/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Alpha Lipoic Acid

Item (One) Only

Place: New Delhi

Date : 06 MAY 2011

**LICENSING AUTHORITY**

अरविन्द कुकरेती

ARVIND KUKRETI

सहायक औषधि नियंत्रक (भारत)

Assistant Drug 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  
06 MAY 2011

M/s. Haresh Chemicals  
27-28, Anant Building , 2nd Floor, 217, S.G. Marg,  
Mumbai-400002

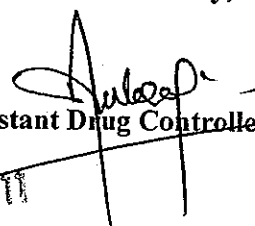
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 25/03/2011 received by this office vide diary no. 14582 dt 29/03/2011, I enclose licence(s) Nos. BD-574-18984 dated 06 MAY 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/40/05-DC(Pt-1) (Re-registration 2009)

Copy together with copy of Licence No. BD-574-18984 dated 06 MAY 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-574-18984

Date 06 MAY 2011

1. M/S. Haresh Chemicals 27-28, Anant Building , 2nd Floor, 217, S.G. Marg, 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. Jiangu Tohope Pharmaceutical Co. Ltd. 188, Wuyishan Road, Southeast Economic Development Zone, Changshu City, Jiangu 215533, 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 04/05/2011 to 14/02/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Alpha Lipoic Acid

Item (One) Only

Place: New Delhi

Date: 06 MAY 2011

LICENSING AUTHORITY

ARVIND KUKRETY

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

स्वास्थ्य विभाग, नया दिल्ली  
Seal/Stamp

Conditions of Licence

Directorate General of Health Services  
F.D.A. Bhawan, Kolla 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/40/05-DC(Pt-1) (Re-registration 2009)

From:

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

③

To:

New Delhi, dated

06 MAY 2011

M/s. Medi Pharma Drug House  
Bulakhidas Bldg, Gr. Floor, 13,  
Vithaldas Road, Princess Street,  
Mumbai-400002

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

Dear Sir/Sirs,

OFFICE COPY

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 30/03/2011 received by this office vide diary no. 15708 dt. 04/04/2011, I enclose licence(s) Nos. BD-574-18985 dated 06 MAY 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,

*[Signature]*  
Assistant Drug Controller (India)

06 MAY 2011

No. 6-3/BD/40/05-DC(Pt-1) (Re-registration 2009)

Copy together with copy of Licence No. BD-574-18985 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), ...

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-574-18985

Date 06 MAY 2011

1. M/S. Medi Pharma Drug House Bulakhidas Bldg, Gr. Floor, 13, Vithaldas 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. Jiangsu Tohope Pharmaceutical Co. Ltd. 188, Wuyishan Road, Southeast Economic Development Zone, Changshu City, Jiangsu 215533, 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 04/05/2011 to 14/02/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

**Alpha Lipoic Acid**

**Item (One) Only**

Place: New Delhi

Date: 06 MAY 2011

**LICENSING AUTHORITY**

ARVIND KUMRETY

सहायक औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

स्वास्थ्य सेवा विभाग

Seal/Stamp

**Conditions of Licence**

Directorate General of Health Services  
F.D.A. Bhawan, Koka 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

06 MAY. 2011

M/s. Gorang Interational  
Bldg. No. 1, Gala NO. 12, 2nd Floor,  
Manish Compound Anjurphata,  
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. Nil Dated 23/03/2011 received by this office vide diary no. 14365 dt. 28/03/2011, I enclose licence(s) Nos. BD-574-18986 dated 06 MAY 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/40/05-DC(Pt-1) (Re-registration 2009)

Copy together with copy of Licence No. BD-574-18986 dated 06 MAY 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-574-18986

Date 06 MAY 2011

1. M/S. Gorang Interational Bldg. No. 1, Gala NO. 12, 2nd Floor, Manish Compound Anjurphata, 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. Jianguo Tohope Pharmaceutical Co. Ltd. 188, Wuyishan Road, Southeast Economic Development Zone, Changshu City, Jianguo 215533, 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 04/05/2011 to 14/02/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Alpha Lipoic Acid

Item (One) Only

Place: New Delhi

Date: 06 MAY 2011

OFFICE COPY

LICENSING AUTHORITY

ARVIND KUKRETY

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

सहायक औषधि नियंत्रक (भारत)

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

6 MAY 2011

M/s. Aquatic Remedies Pvt. Ltd.  
F 108, 109, 110, Sambhav Complex, Village Rahnal  
Dist. Thane Bhiwandi-421302

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

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 15/03/2011 received by this office vide diary no. 13722 dt. 23/03/2011, I enclose licence(s) Nos. BD-574-18987 dated 06 MAY 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/40/05-DC(Pt-1) (Re-registration 2009)

Copy together with copy of Licence No. BD-574-18987 dated 06 MAY 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-574-18987

Date 06 MAY 2011

1. M/S. Aquatic Remedies Pvt. Ltd. F 108, 109, 110, Sambhav Complex, Village  
Rahnal Dist. Thane Bhiwandi-421302 is hereby licensed into India during the  
period for which this licence is in force, the drugs specified below, manufactured By  
M/s. Jiangsu Tohope Pharmaceutical Co. Ltd. 188, Wuyishan Road, Southeast  
Economic Development Zone, Changshu City, Jiangsu 215533, 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 04/05/2011 to 14/02/2012 unless it is sooner  
suspended or cancelled under the said rules.

Names of drugs to be imported

Alpha Lipoic Acid

Item (One) Only

Place: New Delhi

Date 06 MAY 2011

LICENSING AUTHORITY

ARVIND KUKRETY

सहायक औषधि नियंत्रक (भारत)

Assistant Drugs Controller (India)

प्राथम्य सेवा निदेशिका

Directorate General of Health Services

F.D.A. Shawan, 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

06 MAY. 2011

M/s. Pravin Pharma  
Gala No. 102, 1st Flr, Bldg-B, Sri Sambhav Complex,  
Rabala Purna 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. PP/III-A/2011 Dated 29/03/2011 received by this office vide diary no. 15739 dt. 04/04/2011, I enclose licence(s) Nos. BD-178-18988 dated 06 MAY. 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/194/02-DC (Re-Registration 2009)

Copy together with copy of Licence No. BD-178-18988 dated

06 MAY 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-178-18988

Date 06 MAY 2011

**1. M/S. Pravin Pharma Gala No. 102, 1st Flr, Bldg-B, Sri Sambhav Complex, Rabala Purna 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. Yancheng Suhai Pharmaceutical Co. Ltd., No. 92, E, Jiankang Road, Dafeng City, Jiangsu, 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 04/05/2011 to 31/05/2012 unless it is sooner suspended or cancelled under the said rules.**

Names of drugs to be imported

**Doxycycline Hydrochloride EP**

Item (One) Only

Place: New Delhi

Date: 06 MAY 2011

**LICENSING AUTHORITY**

ARVIND KUKRETY

अरविंद कुक्रेती (भारत)

Ministry of Health & Family Welfare, Government of India

General of Health Services

L.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

06 MAY 2011

M/s. Aquatic Remedies Pvt. Ltd.  
F 108, 109, 110, Sambhav Complex, Village Rahnal  
Dist. Thane Bhiwandi-421302

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

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 22/03/2011 received by this office vide diary no. 14012 dt. 25/03/2011, I enclose licence(s) Nos. BD-895-18989 dated 06 MAY 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/75/10-DC

Copy together with copy of Licence No. BD-895-18989 dated 06 MAY 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-895-18989

Date 06 MAY 2011

1. M/S. Aquatic Remedies Pvt. Ltd. F 108, 109, 110, Sambhav Complex, Village  
Rahnal Dist. Thane Bhiwandi-421302 is hereby licensed into India during the  
period for which this licence is in force, the drugs specified below, manufactured By  
M/s. Guangzhou Hanpu Pharmaceutical Co. Ltd. Bai yunling, Aotou Town,  
Conghua city, Guangzhou, Guangdong 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 04/05/2011 to 30/11/2013 unless it is sooner  
suspended or cancelled under the said rules.

Names of drugs to be imported

Clotrimazole IP

Item (One) Only

Place: New Delhi

Date: 06 MAY 2011

  
LICENSING AUTHORITY

अरविन्द कुकरती  
ARVIND KUKRETY  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drug 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  
06 MAY 2011

M/s. Supreem Pharmaceuticals Mysore Pvt. Ltd.,  
Plot No. 73,74 & 48-P1, KIADB Indl Area,  
Nanjangud-571 302

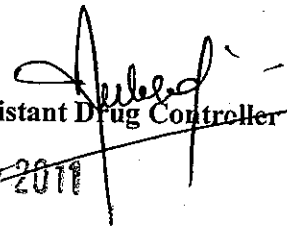
**Sub: Import Licence under the Drugs Act, 1940 and Drugs Rules thereunder.**

Dear Sir/Sirs,

With reference to your application for import licence forwarded to this office with your letter No. Nil Dated 20/04/2011 received by this office vide diary no. 19696 dt. 27/04/2011, I enclose licence(s) Nos. BD-574-18990 dated 06 MAY 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/40/05-DC(Pt-1) (Re-registration 2009)

Copy together with copy of Licence No. BD-574-18990 dated 06 MAY 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-574-18990

Date 06 MAY 2011

1. M/S. Supreem Pharmaceuticals Mysore Pvt. Ltd., Plot No. 73,74 & 48-P1, KIADB Indl Area, Nanjangud-571 302 is hereby licensed into India during the period for which this licence is in force, the drugs specified below, manufactured By M/s. Jianguo Tohope Pharmaceutical Co. Ltd. 188, Wuyishan Road, Southeast Economic Development Zone, Changshu City, Jianguo 215533, 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 04/05/2011 to 14/02/2012 unless it is sooner suspended or cancelled under the said rules.

Names of drugs to be imported

Alpha Lipoic Acid

Item (One) Only

Place: New Delhi

Date: 06 MAY 2011

  
LICENSING AUTHORITY

अरवि कुक्रेती  
ARVIND KUKRETY  
सहायक औषधि नियंत्रक (भारत)  
Assistant Drugs Controller (India)  
स्वास्थ्य सेवा महानिदेशालय

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 licence 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";