J. K. INDUSTRIES

FDA Approved and ISO 9001:2015

CORPORATE OFFICE AND FACTORY

Plot No. D-3, Gane Khadpoli MIDC, Tal-Chiplun, Dist- Ratnagiri. Maharashtra-415605.

Contact: +91-8866669682, +91-8975990934

Email: info@jkinduschem.com | sales@jkinduschem.com | jagdishtandle99@gmail.com

www.jkinduschem.com

About Company:

- **J. K. Industries** is FDA Approved and an ISO 9001: 2015 Certified Bulk drug manufacturing company.
- **Engaged** in manufacturing and marketing of Bulk Drugs and Fine Chemicals with a state-of–art manufacturing facility at Gane Khadpoli MIDC, Chiplun Taluka, Ratnagiri District, and Maharashtra.
- **Specialized** in manufacture of products used in the production of Neutraceuticals, ORS Powder, Cosmetics, Food & Beverages, Veterinary feeds and in compliance to Indian and International Pharmacopoeia standards.
- **Backed** by a qualified team of professional with requisite expertise in technical, quality control, R & D and management.

Quality:

- ❖ Quality is the forte of J K Industries and is one of the biggest factors for the growth.
- ❖ Core of our Quality Philosophy is the commitment to achieve perfection to match the highest international standards.
- ❖ A reputed team of experts and in-house sophisticated lab with the latest equipments for Quality Assurance and Quality Control throughout the production process.
- ❖ Consistent up gradation the technology through precise innovations.
- ❖ Maintain clean, safe and hygiene work environment.
- ❖ Ensure personnel validation through training and evaluation.

Our Products:

Product	Grade
Zinc Sulphate Heptahydrate	IP,BP
Zinc Sulphate Monohydrate	BP, USP
Magnesium Sulphate Heptahydrate	IP, BP
Magnesium Sulphate Monohydrate	USP
Magnesium Sulphate Anhydrous	USP
Potassium Chloride	IP, BP, USP
Ammonium Chloride	IP, BP, USP

Note: Packing to suit customer requirements.

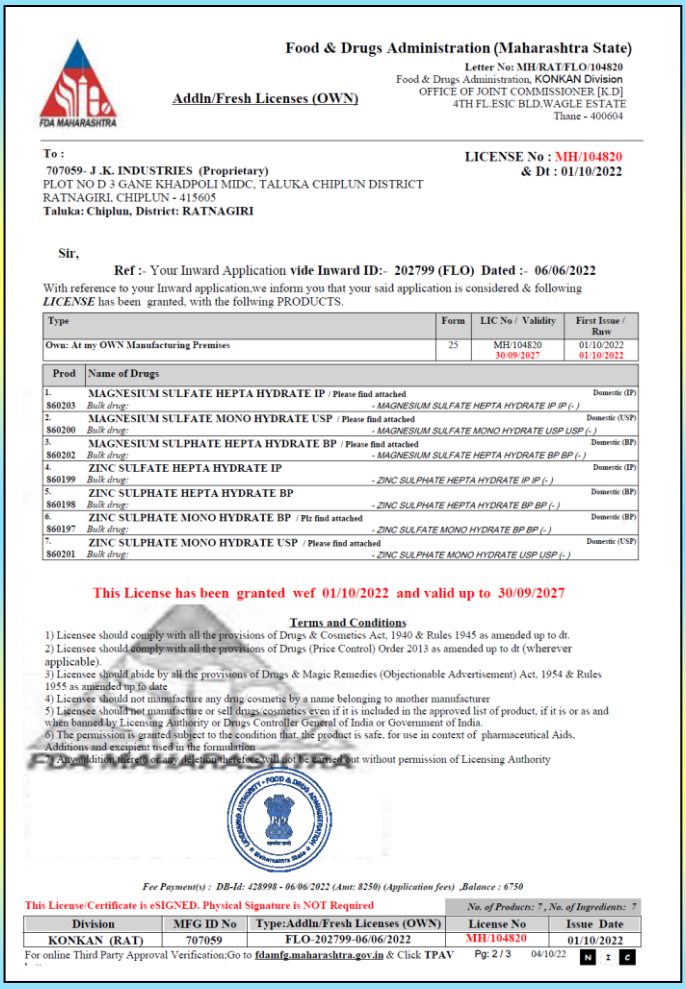
Abbreviations:

IP: Indian Pharmacopoeia

BP: British Pharmacopoeia

USP: United States Pharmacopoeia

Certificates:



Food & Drugs Administration (Maharashtra State)

FDA MAHARASHTRA

Addln/Fresh Licenses (OWN)

Letter No: MH/RAT/FLO/104820
Food & Drugs Administration, KONKAN Division
OFFICE OF JOINT COMMISSIONER [K.D]
4TH FL.ESIC BLD.WAGLE ESTATE
Thane - 400604

To :
707059- J .K. INDUSTRIES (Proprietary)
PLOT NO D 3 GANE KHADPOLI MIDC, TALUKA CHIPLUN DISTRICT
RATNAGIRI, CHIPLUN - 415605
Taluka: Chiplun, District: RATNAGIRI

LICENSE No : MH/104820
& Dt : 01/10/2022

Sir,

Ref :- Your Inward Application **vide Inward ID:- 202799 (FLO) Dated :- 06/06/2022**

With reference to your Inward application,we inform you that your said application is considered & following ***LICENSE*** has been granted, with the follwing PRODUCTS.

Type	Form	LIC No / Validity	First Issue / Renew
Own: At my OWN Manufacturing Premises	25	MH/104820 30/09/2027	01/10/2022 01/10/2022

Prod	Name of Drugs	
1. S60203	MAGNESIUM SULFATE HEPTA HYDRATE IP / Please find attached *Bulk drug:*	Domestic (IP) *- MAGNESIUM SULFATE HEPTA HYDRATE IP IP (-)*
2. S60200	MAGNESIUM SULFATE MONO HYDRATE USP / Please find attached *Bulk drug:*	Domestic (USP) *- MAGNESIUM SULFATE MONO HYDRATE USP USP (-)*
3. S60202	MAGNESIUM SULPHATE HEPTA HYDRATE BP / Please find attached *Bulk drug:*	Domestic (BP) *- MAGNESIUM SULFATE HEPTA HYDRATE BP BP (-)*
4. S60199	ZINC SULFATE HEPTA HYDRATE IP *Bulk drug:*	Domestic (IP) *- ZINC SULPHATE HEPTA HYDRATE IP IP (-)*
5. S60198	ZINC SULPHATE HEPTA HYDRATE BP *Bulk drug:*	Domestic (BP) *- ZINC SULPHATE HEPTA HYDRATE BP BP (-)*
6. S60197	ZINC SULPHATE MONO HYDRATE BP / Plz find attached *Bulk drug:*	Domestic (BP) *- ZINC SULFATE MONO HYDRATE BP BP (-)*
7. S60201	ZINC SULPHATE MONO HYDRATE USP / Please find attached *Bulk drug:*	Domestic (USP) *- ZINC SULPHATE MONO HYDRATE USP USP (-)*

This License has been granted wef 01/10/2022 and valid up to 30/09/2027

Terms and Conditions

1) Licensee should comply with all the provisions of Drugs & Cosmetics Act, 1940 & Rules 1945 as amended up to dt.
2) Licensee should comply with all the provisions of Drugs (Price Control) Order 2013 as amended up to dt (wherever applicable).
3) Licensee should abide by all the provisions of Drugs & Magic Remedies (Objectionable Advertisement) Act, 1954 & Rules 1955 as amended up to date
4) Licensee should not manufacture any drug/cosmetic by a name belonging to another manufacturer
5) Licensee should not manufacture or sell drugs/cosmetics even if it is included in the approved list of product, if it is or as and when banned by Licensing Authority or Drugs Controller General of India or Government of India.
6) The permission is granted subject to the condition that, the product is safe, for use in context of pharmaceutical Aids, Additions and excipient used in the formulation.
7) Any addition, deletion or any alteration in the above will not be carried out without permission of Licensing Authority

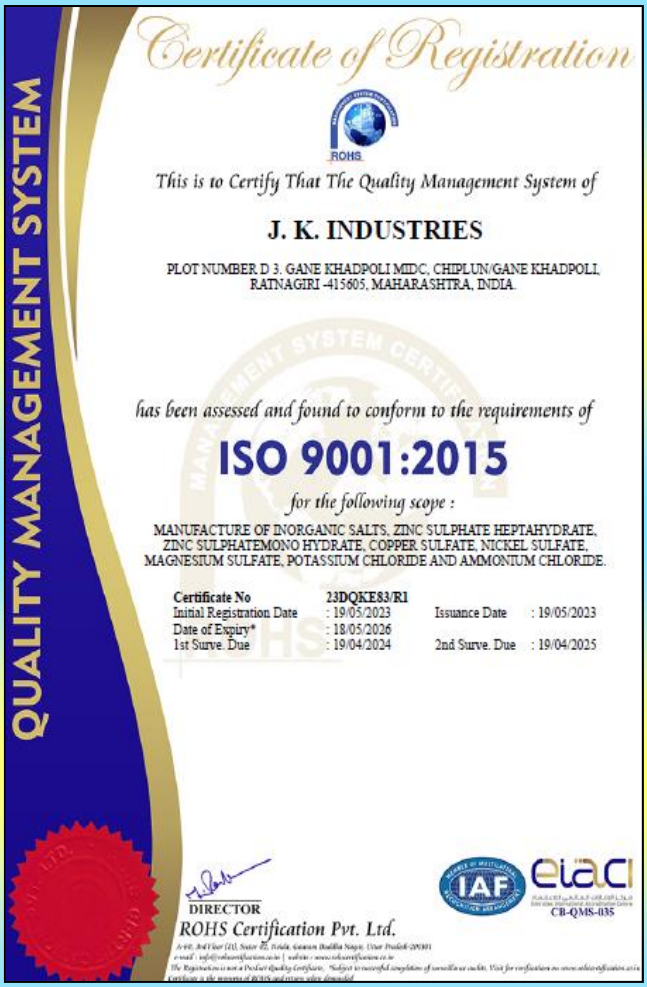
Fee Payment(s) : DB-Id: 428998 - 06/06/2022 (Amt: 8250) (Application fees) ,Balance : 6750

This License/Certificate is eSIGNED. Physical Signature is NOT Required

No. of Products: 7 , No. of Ingredients: 7

Division	MFG ID No	Type:Addln/Fresh Licenses (OWN)	License No	Issue Date
KONKAN (RAT)	707059	FLO-202799-06/06/2022	MH/104820	01/10/2022

For online Third Party Approval Verification;Go to **fdamfg.maharashtra.gov.in** & Click **TPAV** Pg: 2 / 3 04/10/22



QUALITY MANAGEMENT SYSTEM

Certificate of Registration

ROHS

This is to Certify That The Quality Management System of

J. K. INDUSTRIES

PLOT NUMBER D 3, GANE KHADPOLI MIDC, CHIPLUN/GANE KHADPOLI, RATNAGIRI -415605, MAHARASHTRA, INDIA.

has been assessed and found to conform to the requirements of

ISO 9001:2015

for the following scope :

MANUFACTURE OF INORGANIC SALTS, ZINC SULPHATE HEPTAHYDRATE, ZINC SULPHATEMONO HYDRATE, COPPER SULFATE, NICKEL SULFATE, MAGNESIUM SULFATE, POTASSIUM CHLORIDE AND AMMONIUM CHLORIDE.

Certificate No	**23DQKE83/R1**		
Initial Registration Date	: 19/05/2023	Issuance Date	: 19/05/2023
Date of Expiry*	: 18/05/2026		
1st Surve. Due	: 19/04/2024	2nd Surve. Due	: 19/04/2025

DIRECTOR
ROHS Certification Pvt. Ltd.

IAF eIAC CB-QMS-035

Food & Drugs Administration (Maharashtra State)

Letter No: MH/RAT/PRP/1118409
Food & Drugs Administration, KONKAN Division
OFFICE OF JOINT COMMISSIONER [K.D]
4TH FL.ESIC BLD,WAGLE ESTATE
Thane - 400604

Additional Product Permission

PERMISSION No : 1118409
Dt : 18/04/2024

707059- J.K. INDUSTRIES
PLOT NO D 3 GANE KHADPOLI MIDC, TALUKA CHIPLUN DISTRICT RATNAGIRI,
CHIPLUN - 415605
Taluka: Chiplun, District: RATNAGIRI

Ref :- Your Inward Application**vide Inward ID :- 253493 Dated :- 23/01/2024**

With reference to your Inward application,we have to inform you that your said application is considered & following *PRODUCTS PERMISSION* have been granted, under the following *LICENSES.*

Type	Form	LIC No / Validity	First Issue / Raw
Own: At my OWN Manufacturing Premises	25	MH/104820 30/03/2027	01/10/2022 01/10/2022

Prod	Name of Drugs		
1. 918632	AMMONIUM CHLORIDE BP	- AMMONIUM CHLORIDE BP (-)	Domestic (BP)
2. 918631	AMMONIUM CHLORIDE IP	- AMMONIUM CHLORIDE IP (-)	Domestic (IP)
3. 918633	AMMONIUM CHLORIDE USP	- AMMONIUM CHLORIDE USP (-)	Domestic (USP)

Terms and Conditions

1) Licensee should comply with all the provisions of Drugs & Cosmetics Act, 1940 & Rules 1945 as amended up to dt.
2) Licensee should comply with all the provisions of Drugs (Price Control) Order 2013 as amended up to dt (wherever applicable).
3) Licensee should abide by all the provisions of Drugs & Magic Remedies (Objectionable Advertisement) Act, 1954 & Rules 1955 as amended up to date
4) Licensee should not manufacture any drug/cosmetic by a name belonging to another manufacturer
5) Licensee should not manufacture or sell drugs/cosmetics even if it is included in the approved list of product, if it is or as and when banned by Licensing Authority or Drugs Controller General of India or Government of India.
6) The permission is granted subject to the condition that, the product is safe, for use in context of pharmaceutical Aids, Additions and excipient used in the formulation
7) Any addition thereto or any deletion therefore will not be carried out without permission of Licensing Authority

eSign
Digitally Signed
NARENDRA PURUSHOTTAM SUPE
e-Signed on 18-04-2024 12:40
TPAV # WLN2QE4SVS

N. P. SUPE
State Licensing Authority
Food & Drugs Administration
KONKAN Division, Maharashtra State

FDA MAHARASHTRA

For Payment(s) : DD-Id: 494463 - 23/01/2024 (Amt: 2250) (ADDITIONAL PRODUCT FEES) Balance : 1600
COL-Colour, FLV-Flavour,PRV-Preservative,REA-Reagent

This License/Certificate is eSIGNED. Physical Signature is NOT Required

Division	MFG ID No	Type:Additional Product Permission	PERMISSION No	Issue Date
KONKAN (RAT)	707059	PRP-253493-23/01/2024	1118409	18/04/2024

For online Third Party Approval Verification Go to fdamfg.maharashtra.gov.in & Click TPAVisit Pg: 1 / 1 (18/04/24)

Food & Drugs Administration (Maharashtra State)

Letter No: MH/RAT/PRP/1118326
Food & Drugs Administration, KONKAN Division
OFFICE OF JOINT COMMISSIONER [K.D]
4TH FL.ESIC BLD,WAGLE ESTATE
Thane - 400604

Additional Product Permission

PERMISSION No : 1118326
Dt : 16/04/2024

707059- J.K. INDUSTRIES
PLOT NO D 3 GANE KHADPOLI MIDC, TALUKA CHIPLUN DISTRICT RATNAGIRI,
CHIPLUN - 415605
Taluka: Chiplun, District: RATNAGIRI

Ref :- Your Inward Application **vide Inward ID :- 253498 Dated :- 30/01/2024**

With reference to your Inward application,we have to inform you that your said application is considered & following *PRODUCTS PERMISSION* have been granted, under the following *LICENSES.*

Type	Form	LIC No / Validity	First Issue / Raw
Own: At my OWN Manufacturing Premises	25	MH/104820 30/03/2027	01/10/2022 01/10/2022

Prod	Name of Drugs		
1. 918488	POTASSIUM CHLORIDE BP	- POTASSIUM CHLORIDE BP (-)	Domestic (BP)
2. 918489	POTASSIUM CHLORIDE IP	- POTASSIUM CHLORIDE IP (-)	Domestic (IP)
3. 918487	POTASSIUM CHLORIDE USP	- POTASSIUM CHLORIDE USP (-)	Domestic (USP)

Terms and Conditions

1) Licensee should comply with all the provisions of Drugs & Cosmetics Act, 1940 & Rules 1945 as amended up to dt.
2) Licensee should comply with all the provisions of Drugs (Price Control) Order 2013 as amended up to dt (wherever applicable).
3) Licensee should abide by all the provisions of Drugs & Magic Remedies (Objectionable Advertisement) Act, 1954 & Rules 1955 as amended up to date
4) Licensee should not manufacture any drug/cosmetic by a name belonging to another manufacturer
5) Licensee should not manufacture or sell drugs/cosmetics even if it is included in the approved list of product, if it is or as and when banned by Licensing Authority or Drugs Controller General of India or Government of India.
6) The permission is granted subject to the condition that, the product is safe, for use in context of pharmaceutical Aids, Additions and excipient used in the formulation
7) Any addition thereto or any deletion therefore will not be carried out without permission of Licensing Authority

eSign
Digitally Signed
NARENDRA PURUSHOTTAM SUPE
e-Signed on 16-04-2024 15:21
TPAV # 6UR2975JTE

N. P. SUPE
State Licensing Authority
Food & Drugs Administration
KONKAN Division, Maharashtra State

FDA MAHARASHTRA

For Payment(s) : DD-Id: 493054 - 30/01/2024 (Amt: 2250) (ADDITIONAL PRODUCT FEES) Balance : 1600
COL-Colour, FLV-Flavour,PRV-Preservative,REA-Reagent

This License/Certificate is eSIGNED. Physical Signature is NOT Required

Division	MFG ID No	Type:Additional Product Permission	PERMISSION No	Issue Date
KONKAN (RAT)	707059	PRP-253498-30/01/2024	1118326	16/04/2024

For online Third Party Approval Verification Go to fdamfg.maharashtra.gov.in & Click TPAVisit Pg: 1 / 1 (16/04/24)