



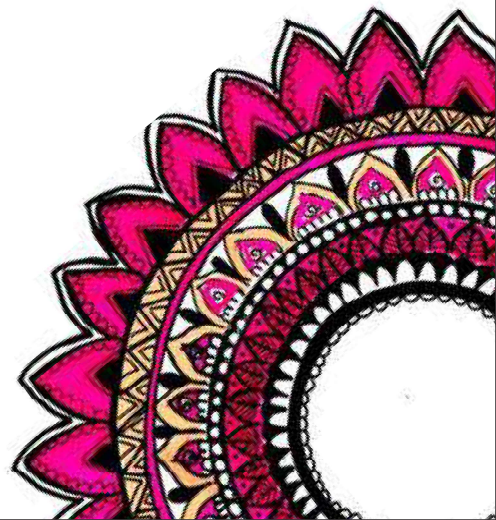
MMC
Healthcare Ltd

A SURAKSHA Group Company



We Care for Life...

Art by : Dr.Padmaja Manepalli



SURAKSHA

We Care for Life...

MANAGEMENT



FOUNDER
MANEPALLI
NAGESWARA RAO



CHAIRMAN
M.V.K. NAGESWARA RAO



MANAGING DIRECTOR
Dr.M. NAGESWARA RAO



DIRECTOR
M. SOMESWARA RAO



DIRECTOR
Dr. M. PADMAJA RANI



DIRECTOR
M.R.T. ASWANI



DIRECTOR
M. RAJ ARYAN



DIRECTOR
M. RISHIKA

GROUP



DIVISIONS



ACCREDITATIONS & AFFILIATIONS



Indian Pharma
Graduates' Association



Indian Drug
Manufacturers Association



Federation of
Indian Export Organisations



The Indian Pharmaceutical
Association



Andhra
Chamber of Commerce



Pharmaceuticals
Export Promotion Council



WHO GMP



NHP
CANADA



SCHEDULE
M GMP



FSSAI



AYUSH



NABL



ISO



HALAL



HACCP



KOSHER

SERVICES

IN HOUSE

FRD · ARD · Pellets
Folds · Designing

SPECIALITIES

Psychiatric · Neuro · Diabetic
· Cardiac · Gynaec · Pediatric
Ortho and General Medicines

CONSULTATION

DPCO · DCGI
Branding · Narcotic
Trademark

SECTIONS

Beta Lactam
Non Beta Lactam



Generations
of Expertise



Companies



Divisions



States India
Presence



Countries Global
Presence



Global Customers



Export
Own Labels



Employees



Domestic
Own Labels



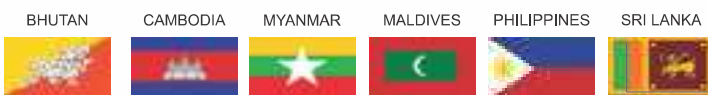
India
Customers



Domestic
Private Labels

GLOBAL PRESENCE

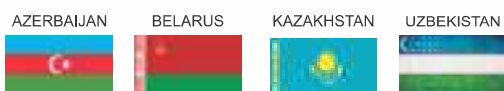
ASIA



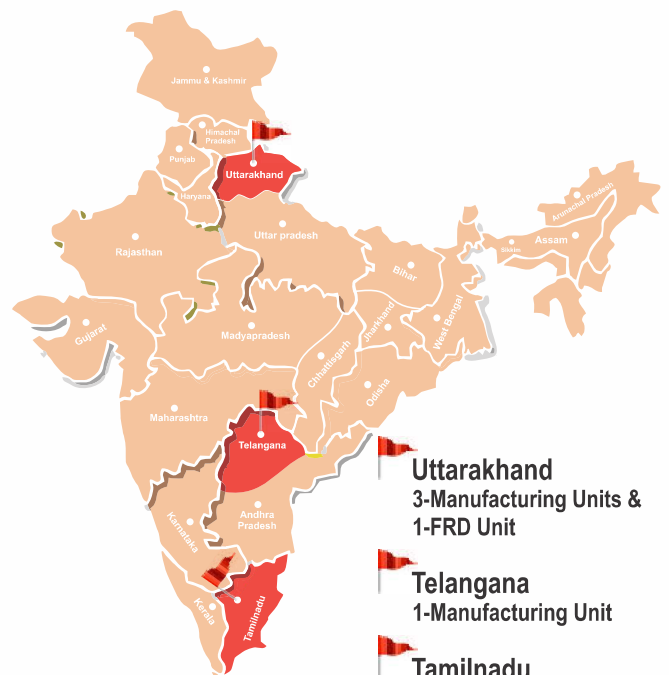
AFRICA & MIDDLE EAST



CIS



NORTH AMERICA & LATAM



Global Divisions



Vitalis is a brand of dietary supplements designed to offer various health benefits. These supplements are intended for oral administration and serve as valuable sources of nutrients, contributing to improved health and overall wellness.

Healthy habits, Happy life...



The Magic Power House...

Hemp Tribe, we are passionate about unlocking the full potential of hemp and bringing its benefits to everyone. Our mission is to lead the Hemp Revolution by offering a wide range of HemPicked products that are carefully curated to deliver the finest hemp-based solutions. We believe in the power of this versatile plant and its ability to transform lives for the better.

Life is better with Hemp...



Innovation with Tradition...

Suraksha Naturals Combines Modern Innovation with Traditional Wellness practices in Revolutionary "Ketoveyda" Product Line

Ketoveyda Product-Line Works Synergistically with a keto diet, and is created utilizing traditional Ayurvedic Practices, enhanced by modern innovation. Suraksha Naturals has pioneered a line of Products called ketoveyda, with meal options and supplements that help the body recover the nutrients the might be lost to a conventional keto diet

Natural Approach Towards Health...



Nextra is a niche Nutraceutical line of Suraksha Natural Inc. Nextra Provides Nutritional and Speciality Health supplements which energizes both Mind and Body

Nextra ensures Wellness...

ACCREDITATIONS

L. No. 96.8732X-I/2004
MPL/CAs/19899

Office of the Director of Drugs Control,
Tamil Nadu, Chennai - 600 006
Date: 26.04.2004

This is to certify that **M/s. MMC Healthcare Limited**, No.248, State Industrial Estate, Thirumachilai, Chennai - 600 124, is holding license in **Forme 25 & 25** bearing Nos **5887987 & 3577987** dated: **01.04.2004**. It is also certified that they have renounced the license absolutely from the further period from **01.04.2004** to **31.03.2009** as per Drugs Rules 1945 for retaining the License.

Yours faithfully,
Deputy Director

M/s. MMC Healthcare Limited,
No.248, State Industrial Estate,
Thirumachilai, Chennai - 600 124

Signature valid
Date: 26.04.2004
DDC/26.04.2004/400




**DEPARTMENT OF FOOD SAFETY AND DRUGS CONTROL, ADMINISTRATION
 GOVERNMENT OF TAMILNADU**
 No. Anna Salai, Chennai - 600 006, Tamil Nadu.

VERIFICATION OF GOOD MANUFACTURING PRACTICES

Confirmed No. & the No. of Samples taken. Date: 01/01/2024

On the basis of the inspection carried out on 27/01/2024 & 28/01/2024 and 26/02/2024 we certify that the site included in the certificate complies with Good Manufacturing Practices for the storage forms, categories and activities listed in Table '1'.

1. Name and address of site: M/S MNC Healthcare Limited
 30-8 Sales Indu Street
 Thiruvallur, Chennai-602039

2. Manufacturer's license number: Form 22 Binding No: 60019607, dated 01/04/2024
 Form 22 Binding No: 60719607, dated 01/04/2024

3. Table '1'

Storage Form	Category/Product	Authorized
Tablets, Hard Capsules, Capsules, Lozenges and Bulk Drugs (Pharmaceutical Grade) Pharmaceutical Ingredients (GMP Warehouse)	General	Manufacturer

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured from this process will rest with the manufacturer.

This certificate remains valid until 31.12.2024 & becomes invalid if the activities under categories certified herewith are straggled or if the site is no longer associated to its in compliance with GMP.

Name and function of responsible person: M. N. SUNDAR, B. Pharm.



 **STATE LICENSING AUTHORITY (SLA)
 CONTROLLERS AUTHORITY**
 1, Anna Salai

M. N. Sundar, B. Pharm.
 Director, State Licensing Authority
 1, Anna Salai, Chennai-600 006

No. of Samples taken: M/S. MNC Healthcare Limited
 30-8 Sales Indu Street
 Thiruvallur, Chennai-602039
 600, Chennai-600 039

This certificate is issued on 01/04/2024

 Form C Government of India Food Safety and Standards Authority of India (Enacted under FSS Act, 2006)		 भारत सरकार
एप्लिकेशन / Application Form		
1. Name of Applicant (FFO or address of Applicant): (applicable to college students as well)	NAME: Ms. T. CHANDRASEKHAR STATE: Kerala / District: Thiruvananthapuram Registration No./College Name: Thiruvananthapuram / Thiruvananthapuram	
2. Address of Applicant (Residence or College Office as per)	No. 4, 10007, Industrial Estate, The Industrial Estate, KADAPPA, K. Nagar, Puzhuvayal, K. Nagar, Changanassery, Thiruvananthapuram	
3. Kind of Applicant (College student)	Manufacturer / Food or Food Supplement and Technological Unit Manufacturer / Supplier / Manufacturer / Distributor / Supplier	
4. Date of Birth (Date of Birth of Applicant)	NA	
5. Category of Applicant (College as per)	Regular student	
We warrant that the information provided in this form is true and correct to the best of our knowledge and belief. We warrant that the information provided in this form is true and correct to the best of our knowledge and belief.		
Name of user: Chandrasekhar Contact No./Mobile: 94 47 22 24 24 / Email address: chandrasekhar@ssai.gov.in Email address / Mobile: 94 47 22 24 24 / Email address: chandrasekhar@ssai.gov.in	Signature of user  Date: 01/07/2024 / Time: 10:00:00 Location: Thiruvananthapuram / Date: 01/07/2024	
Remarks: 1. Product category 2. Category of product 3. Registration number 4. Conditions of use		

LURG CERTIFICATIONS
ISO 9001:2015

Certificate of Registration

ISO Certification body certifies that the Quality Management System of:

MMC HEALTHCARE LTD
B-94/B, 3000 Karam Road, Newmarket, Ontario M3J 4A, Canada, India

has been assessed and found to comply to complete and meets the requirements of following standard:

ISO 9001:2015

For the scope of:

Manufacturing & Exporting of Various Operating and Prosthetic, Dental and Prosthetic Supplies

Initial date of certification:	May 14, 2023
Current date of certification:	May 14, 2023
Date of expiry:	May 13, 2026

(Subject to surveillance visits as per ISO 9001:2015)

1 st Surveillance Date:	Jun 2023
2 nd Surveillance Date:	Jul 2023

Certificate Number: ISO9001/2015/12400
ISO Code: 12400

ISO Certification:
Mr. Ajay Singh
Registrar/Signatory

LURG CERTIFICATIONS
ACCREDITED

ISO 9001:2015

Registration and certification are subject to surveillance visits as per ISO 9001:2015

WRO
WORLDWIDE
REGISTRATION
ORGANIZATION

**CERTIFICATE
OF EXCELLENCE**

WRO Certification hereby certifies that the Health Management System of

MMC HEALTHCARE LTD

914 H, 999 P, Changanacherry, Thiruvananthapuram, Kerala 695 022, Appal Nalla India

has been assessed and found to operate in compliance with the requirements of following standard

HALAL CERTIFICATE

For the scope of

Management of Health-care System, Medication Safety, Medication Storage, Medication Production & Supply

Issue date of certificate: 15 July 2010
 Expiry date of Certificate: 15 July 2011
 Issuing Agency: Dr. Jay. Jeyaraj
 Issuing Institution: WRO at MMC Healthcare Ltd
 Issued To: MMC Ltd
 Issued on: 15 July 2010
 Issued By: Dr. Jay. Jeyaraj

Dr. Jay. Jeyaraj

WRO CERTIFICATION

WRO
WORLDWIDE
REGISTRATION
ORGANIZATION

ACCREDITED



HAZARD ANALYSIS & CRITICAL CONTROL POINTS

Certificate Of Excellence

WRG Certification hereby certifies that the HACCP System of

MMC HEALTHCARE LTD

XXXX XXXX Industrial Estate, Portsmouth PO15 019A, Hampshire, UK

has been successfully found to comply with the requirements of the HACCP system as set out in the HACCP standard.

HACCP

for the scope of:

**Manufacturing & Supply of Various Capsules and Tablets
Chlorinated & Plain Tablets**

Issue Date of certificate	Mar 24, 2020	
Current Date of certification	Mar 24, 2020	
Date of expiry	Mar 23, 2028	

1. Surveillance Date

Feb 2025

2. Surveillance Date

Feb 2025

1. Surveillance Date

Feb 2025

2. Surveillance Date

Feb 2025

Certificate No. : WRG/HA/00000001/00000001

HACCP Code : 0126-02




Manufacturing and Supply of Various Capsules and Tablets
Chlorinated & Plain Tablets

Manufacturing and Supply of Various Capsules and Tablets
Chlorinated & Plain Tablets



MMC Healthcare Ltd.

(WHO-GMP & ISO 9001:2015 Certified Company)

QUALITY POLICY

MMC Healthcare Limited is a WHO GMP Certified and ISO 9001:2015 accredited facility.

MMC Healthcare Limited policy is to preserve and improve patient health by consistently delivering high quality, safe and effective specialty pharmaceutical formulations .

While doing so MMC Healthcare Limited shall keep in mind, the safety of our employees, all work processes as well as our natural environment.

MMC Healthcare Limited aims to improve the processes, products and services continuing to deliver with highest professional standards, thereby providing a competitive advantage to the customers and end -users.

MMC Healthcare Limited will ensure maximum customer satisfaction and continuing quality improvement at all times.

For MMC Healthcare Limited.,

(Dr.M. Nageswara Rao Gupta)
Chief Executive Officer

DOSAGE FORMS

- Bilayered Tablets
- Immediate Release Tablets
- Sustained Release Tablets
- Extended Release Tablets
- Modified Release Tablets
- Pellets In Capsules
- Powder In Capsules
- Taste Masked Oral Liquids
- Bilayered Coated Tablets
- Mouth Dissolving Tablets
- Film Coated Tablets
- Chewable Tablets
- Liposomal Technology
- Enteric Coated Capsules
- Sachets
- Oral & Sublingual Sprays

WE SUPPORT

SECTIONS :

- Non Beta Lactam
- Beta Lactam
- FSSAI

SPECIALITIES :

- Psychiatric
- Neuro
- Cardiac
- Diabetic
- Gynaec
- Pediatric
- Ortho and General Medicines

CONSULTATION :

- DPCO
- DCGI
- Branding
- Narcotic
- Trademark

IN HOUSE :

- FRD
- ARD
- Pellets
- Foils
- Designing

GMP PLANT APPROVALS

 Sri Lanka Medical Council ශ්‍රී ලංකා වෛද්‍ය පාලන සභාව		Application Form අයදුම් පත්‍රය	
The Managing Director Health Services Division No. 100/1, New Road, Nugegoda, Colombo 06 10000-01		Application Form 2019/01/01	
Attention: Manager, Regulatory			
LETTER OF APPROVAL OF FOREIGN PHARMACEUTICAL MANUFACTURING SITE OF FINISHED DOSEAGE FORMS			
Dear Sir/Madam,			
Advise you as the application dated 2019-01-01 for approval of manufacturing site of [Select Medicines] are			
Approved of the Office of the Registrar, No. 10/0, 10/001, Industrial Estate, Pannipitiya, Colombo, 10000, Sri Lanka			
Approved of the Manufacturing Site, No. 10/0, 10/001, Industrial Estate, Pannipitiya, Colombo, 10000, Sri Lanka			
Types: Category (a) and (b) and (c) of products, namely, manufactured as			
Category of Product:		Dosage:	
Strength/Concentration of Product:		Tablet/Capsule/Injection/Suppository	
Product Name:		Name:	
Evaluation of application for pharmaceutical manufacturing site has been completed.			
Approval is granted for the above manufacturing site for the categories of products and Dosage Form mentioned above.			
As per it is approved meeting manufacturing site, approval must be granted.			
Yours faithfully,			
 Registrar			
(Sd/-) If you wish to add more types of categories/dosage strength of products, advise them separately.			
(Sd/-) If you wish to add more types of categories/dosage strength of products, advise them separately.			
(Sd/-) If you wish to add more types of categories/dosage strength of products, advise them separately.			

Sri Lanka

[illegible]

Cambodia

[illegible]

Ghana

2004377


NATIONAL DRUG AUTHORITY

CERTIFICATE OF COMPLIANCE WITH GOOD MANUFACTURING PRACTICE GUIDELINES

THE NATIONAL DRUG POLICY AND AUTHORITY ACT, CAP 209
Enacted under Regulation 20(5) of the Pharmacy Drug Policy and Authority (Establishment) Regulations, 2014

(Certificate No. 945/2022)

This is to certify that the third manufacturing facility

Name of facility: **ABC Healthcare Limited**

Physical address of facility: **34/35, 95000 Industrial Estate, Torranceville, Chennai - 605 004, India**

Licence number of the manufacturer: **Licence No. 528 in Types 301 and 302 in Form 28**
 valid till 30th September 2025, Subject to the
 Conditions of Licence Extension, Latest Valid, India

Has been inspected by the Authority for compliance with the Good Manufacturing Practice Guidelines.

On the basis of the inspection carried out on 28th and 29th November 2024, it is certified that the facility complies on key aspects stipulated with Good Manufacturing Practice for storage items held in Table 1 below.

Table 1. Approved list.

No.	Storage Item	Category	Activities
1	Tablets, Capsules & Uncoated Hard Gelatin Capsules	Non-Rate Locked	Manufacture of Finished Pharmaceutical (Self-Used Product)

The responsibility for the validity of the individual activities of the drugs manufactured through this facility lies with the Manufacturer.

This certificate remains valid till 28th November 2024. It remains invalid if the activities of the categories stipulated change or if the facility is no longer compliant for its compliance with GMP.

Issued Date: 02nd September 2022.


NATIONAL DRUG AUTHORITY
 Dr. Ganes Moorthy
 Vice-Chief Authority

Uganda

[illegible]

Yemen

[illegible]

Nigeria



THE UNITED REPUBLIC OF TANZANIA

MINISTRY OF INDUSTRY AND TRADE

TANZANIA BUREAU OF STANDARDS (TBS)



Address: P.O. Box 35890, Dar es Salaam
 E-mail: info@tbs.go.tz
 Tel: +255 (0)22 261 2000
 Fax: +255 (0)22 261 2001

Ref No. : TB/0000000000 **Date: 2024-12-15**

MEMO TO: **Mr. [Name]**
 To: **Mr. [Name]**
 From: **Mr. [Name]**
 Subject: **[Subject]**

THE GOOD MANUFACTURING PRACTICE INSPECTION OF MEAT HEALTH CARE

Reference is made to the above captioned subject and the inspection which was conducted at your food manufacturing plant on 10/11/2024 by the Tanzania Bureau of Standards (TBS). This inspection was conducted as a result of your application for registration of your food products in Tanzania.

1. Based on the information provided and the inspection visit for inspection food products you are informed that the plant complies with the Good Manufacturing Practice (GMP) minimum requirements as per Criteria for Good manufacturing practice for high risk food products. Therefore the validity of the GMP certificate is five years from the date of inspection of the facility.

2. TBS further advises on the result of your company to further improve the areas listed in the Form TB/0000000000 and continue adhering to the requirements of Good Manufacturing Practice.

3. Thank you for your cooperation.


 Dr. [Name]
 DIRECTOR GENERAL

RECEIVED

DATE _____

TIME _____

BY _____

FOR _____

RECEIVED

DATE _____

TIME _____

BY _____

FOR _____

Tanzania



PHARMACY, MEDICINES & POISONS BOARD
(Act No. 75 of 1986)

GMP CERTIFICATE

This is to certify that, **MNC Healthcare Limited**, is a manufacturing plant situated at **Chennai, INDIA**. The plant was certified by the Pharmacy Medicines and Poisons Board Inspectors, in strict compliance with the provisions of the Pharmacy, Medicines and Poisons Act (75/86), particularly at points of processes in the manufacturing of drugs from suitability of equipment, process controls and the overall quality assurance system.

First Inspection: **24th February, 2019**
 Registration No: **PMPC-GMP/01/2019/04**
 Compliance with GMP: **Compliment**
 Valid up to: **30th June, 2022**


 (DIRECTOR GENERAL)


 (DIRECTOR GENERAL)

Date of Issue: **1st July, 2019**



Malawi

[illegible]

Ethiopia

PRODUCT REGISTRATION CERTIFICATES

National Medicines Regulatory Authority
Sri Lanka

CERTIFICATE OF REGISTRATION OF A MEDICINE

Registration No: SL/2019/001234

Product Name: Paracetamol Tablets

Manufacturer: ABC Pharmaceuticals Ltd, Colombo, Sri Lanka

Valid from: 01/01/2020 to 31/12/2025

Signature: [Signature]

Official Seal: [Seal]

Sri Lanka

Ministry of Health
Kingdom of Cambodia

CERTIFICATE OF REGISTRATION

Registration No: KH/2018/005678

Product Name: Chloroquine Phosphate

Manufacturer: DEF Pharma Co., Phnom Penh, Cambodia

Valid from: 15/06/2019 to 14/06/2024

Signature: [Signature]

Official Seal: [Seal]

Cambodia

FOOD AND DRUGS AUTHORITY
Ghana

CERTIFICATE OF REGISTRATION OF A DRUG

Registration No: GH/2017/009012

Product Name: Chloroquine Phosphate

Manufacturer: GHI Pharmaceuticals Ltd, Accra, Ghana

Valid from: 01/01/2018 to 31/12/2023

Signature: [Signature]

Official Seal: [Seal]

Ghana

Food and Drug Administration
Philippines

CERTIFICATE OF PRODUCT REGISTRATION

Registration No: PDA/2019/001234

Product Name: Paracetamol Tablets

Manufacturer: JKL Pharmaceuticals Inc., Manila, Philippines

Valid from: 01/01/2020 to 31/12/2025

Signature: [Signature]

Official Seal: [Seal]

Philippines

TANZANIA BUREAU OF STANDARDS
Tanzania

PRODUCT REGISTRATION CERTIFICATE

Registration No: TB/2018/005678

Product Name: Paracetamol Tablets

Manufacturer: MNO Pharmaceuticals Ltd, Dar es Salaam, Tanzania

Valid from: 01/01/2019 to 31/12/2024

Signature: [Signature]

Official Seal: [Seal]

Tanzania

NATIONAL AGENCY FOR FOOD AND DRUG ADMINISTRATION AND CONTROL
Nigeria

Certificate of Registration

Registration No: NAFDAC/2017/009012

Product Name: Paracetamol Tablets

Manufacturer: PQR Pharmaceuticals Ltd, Lagos, Nigeria

Valid from: 01/01/2018 to 31/12/2023

Signature: [Signature]

Official Seal: [Seal]

Nigeria

Ministry of Health
Uzbekistan

CERTIFICATE OF REGISTRATION

Registration No: UZ/2019/001234

Product Name: Paracetamol Tablets

Manufacturer: RST Pharmaceuticals Ltd, Tashkent, Uzbekistan

Valid from: 01/01/2020 to 31/12/2025

Signature: [Signature]

Official Seal: [Seal]

Uzbekistan

Ministry of Health
Myanmar

DRUG REGISTRATION CERTIFICATE

Registration No: MY/2018/005678

Product Name: Paracetamol Tablets

Manufacturer: TUV Pharmaceuticals Ltd, Yangon, Myanmar

Valid from: 01/01/2019 to 31/12/2024

Signature: [Signature]

Official Seal: [Seal]

Myanmar

Ministry of Health
Kyrgyzstan

CERTIFICATE OF REGISTRATION

Registration No: KG/2017/009012

Product Name: Paracetamol Tablets

Manufacturer: VWX Pharmaceuticals Ltd, Bishkek, Kyrgyzstan

Valid from: 01/01/2018 to 31/12/2023

Signature: [Signature]

Official Seal: [Seal]

Kyrgyzstan

PRODUCT REGISTRATION CERTIFICATES



Kazakhstan



Tajikistan



Guatemala



Maldives



Zambia



Nigeria



Jordan



Georgia



Ivory Coast

SITE REGISTERED COUNTRIES

AFRICAN MARKETS

Nigeria
Ethopia - Nutra
Ghana
Ivory Coast
Malawi
Tanzania - Nutra
Uganda
Zambia

ASIAN MARKETS

Cambodia
Sri Lanka
Maldives
Philippines

EXPORTING COUNTRIES

Burundi
Burkina Faso
Cambodia
DR Congo
Sri Lanka
Tajikistan
Uganda
Ghana
Guatemala
Jordan
Kazakhstan
Kyrgyzstan
Maldives
Mali
Mauritius
Myanmar
Nigeria
Philippines
Uzbekistan
Zambia

DEEMED EXPORTS

Colombia
South Africa
Kenya
Rwanda
Trinidad
Armenia
Botswana
Jamaica
Laos
Ecuador
El Salvador & Honduras
Ethopia
Peru
Sierra Leone
Tanzania
Dominican Republic
UAE