

SPECTRUM

MEDICAL AND LAB SUPPLIES

www.spectrum-medlab.com



Company Profile

Spectrum for medical and lab supplies is an innovative sourcing & distribution company, involved in exclusive representation and global supply of a wide range of medical devices, commodities, and high-quality pharmaceuticals.

Based and founded in Jordan back in 2011 with offices and branches based in Iraq which is our main market we cover the entire need of the health care sector in the Iraqi market, we also have distribution centers in key cities to make sure the fast and cost-efficient delivery for our customers.


The founders of this company and their staff have worked for the Iraqi ministry of health and occupied high rank positions in the government whether that was for a short period or long it results in know how when it comes to successfully winning and fulfilling tenders and contracts.

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Tel: +962 6 5518791
Fax: +962 6 5518798

Khalda , Building 123 , cross two streets.
Ataba Ben Nafea and st.. Deir Al Asal (Jordanian Village)

Baghdad-Iraq
Tel: +964 78 19802020



Our new and improved Product Support Team composed of highly trained sales personnel, scientists and market researchers, who work closely with our approved GMP partner manufacturers and CRAMS organizations to successfully develop products from the earliest conception stage up till product commercialization.

We are an ISO9001:2015 certified company, specializing in providing full sourcing options and expert regulatory support for the widest range of the rarest and newest products supplied from manufacturing partners with full compliance to strict cGMP guidelines and regulated by leading authorities such as WHO, EDQM, TGA, US-FDA, UK-MHRA etc.



Every detail from manufacturing to packaging to shipping is handled with quality in mind..



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SPECTRUM TEAM

Our core team consists of highly qualified professionals with excellent academic accreditations who are fully experienced in various aspects of the medical, lab, and pharmaceutical industry.

Our employees share the vision of a better future for patients, as well as the progress and success of our organization. The performance acclimatized work culture and accountable working approach of our company attracts top professionals, who strive to strengthen the organization's image as a global leader in the healthcare industry.

Our main departments of medical devices, commodities, and pharmaceuticals are independent with their own managements and staff to ensure and fulfill the global regulations and high-quality standards.

Our mission is to become the preferred supply partner to our customers and achieve sustained growth, through consistent delivery of innovative, customer-centric, world-class quality products.

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SPECTRUM CENTERS

Our **Maintenance service centers** are highly qualified professionals with excellent academic accreditations who are fully experienced in various aspects of the medical & labrotary devices, Located in Baghdad , Najaf , and basrah

Our **private shops** 15 shops around Iraqi provinces , which are related with private & Government hospitals for direct sales the implemnted sales startegy and the highly skilled salesrepresntitives are the key factors of our high annual revenue

Our **Scientific Bureaus** 5 Offices, manage all tenders with Iraqi Ministry Of Health to follow up and fulfill the contracts according to the latest global regulations and high-quality standards which are implemented by the MOH



AL-Najaf Center



Baghdad Center

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SPECTRUM Group Com

LGM LGM
GROUP GROUP



KIVA
Fortress For Electrical Technology

شركة البيت الذهبي العراقي
IRAQ GOLDEN HOUSE
التجارة العامة GENERAL TRADING COMPANY

DENAMARK ELECTRIC



CERTIFICATE



This Certificate is granted to the organization

LGM
LGM INTERNATIONAL
MANUFACTURING GROUP INCORPORATED
Rt. 20, 275, 48 Courtland Avenue E.
Mississauga, Ontario,
Canada

Has been assessed and found to be in accordance with the standard requirements,

ISO 13485:2016
Medical Devices Quality Management System

For the full scope of certification

Provision of manufacturing including design, installation,
maintenance, trading of medical and dental device and
equipment, medical supplies and consumables.

Certificate No. : ICA/130000
Certificate Date : July 02, 2018
Valid Until : July 02, 2020
Registration No. : 2017/01/0000
Renewal Date : 17th April 2019

Signed on behalf of ICA by

President



EC Verification of Conformity

Request No.: 301P0004



This Certificate is granted to the organization

**LGM International
Manufacturing Group Incorporated**
Ontario Corporation: 00099924
Rt. 20, 275, 48 Courtland Avenue E.
Mississauga, Ontario,
Canada



Brand **LGM**
GROUP

We certify that the technical documentation of the product, drawn by the applicant, is in compliance with the specified Directives and Standards

Directive 93/42/EEC

In accordance ICA's products conformity assessment criteria and defined procedures for assess the application of Council of Directive for Medical Device Directive 93/42/EEC, directive and as referred to the information contained on our website (the description of the true scope and extended conformity and with regard to manufacturer's declaration of conformity this is to certify that the applicant's product(s), one aspect, are fulfill the essential requirements of EEC directive 93/42/EEC (amended by 2007/1/EC). Further, the product liability rest with the manufacturer or its representative in accordance with Council Directive 85/374/EEC.

Product(s) : ANKIS - 1
Mark : ANKIS - 1
Reference Standards : Directive 93/42/EEC
Medical devices annex II

The product liability rest with the manufacturer of the product(s) in accordance with Council Directive 85/374/EEC.
This document is issued by ICA on behalf of the manufacturer. The manufacturer shall accept the product liability rest with the manufacturer of the product(s) in accordance with Council Directive 85/374/EEC.

Certificate No. : ICA/EC130000
Verification Date : May 12, 2018
Certificate Date : May 12, 2019
Renewal Date : May 12, 2020
Registration No. : 2017/0000

Inspection/audit activities to verify compliance with the product(s) shall be conducted by ICA and the manufacturer shall, the manufacturer is required to provide the product(s) to the inspection/audit activities. The manufacturer shall be responsible for the product(s) in accordance with Council Directive 85/374/EEC.
Signed on behalf of ICA by

President

President

This certificate shall be reviewed every three (3) years or any other period of time as determined by ICA. The manufacturer shall be responsible for the product(s) in accordance with Council Directive 85/374/EEC.

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