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Annual Registration Successful

Facility: KEISEI MEDICAL INDUSTRIAL CO., LTD., BUNKYO-KU, Tokyo, JAPAN

You have successfully updated your registration and listing information for 2024.

Your registration will be valid through Dec 31, 2024.

Be sure to print this page for your records.

The next registration renewal period is October 1 - December 31, 2024.

Registering your facility and listing devices does not, in any way, constitute FDA approval of your facility or devices.

You may contact the FDA with any questions at reglist@cdrh.fda.gov.

The Owner/Operator Number for this Registration is: 8031595.

Facility Information

Registration Number:

9611484

Initial Importer:

N

Facility Name:

KEISEI MEDICAL INDUSTRIAL CO., LTD.

Legal Name:

Address:

19-6, HONGO SANCHOME,

BUNKYO-KU, Tokyo, 113-0033, JAPAN

DUNS Number:

Foreign Trade Zone:

Ν

Facility URL:

www.kelseimed.com

Other Business Trade Name(s):

Establishment located on a campus:

Owner/Operator Information

Owner/Operator Number:

8031595

Contact Name:

SHIGERU - UETAKE

Company:

KEISEI MEDICAL INDUSTRIAL CO., LTD.

Address:

19-6, Hongo Sanchome

Bunkyo-ku, TOKYO, 113-0033, JAPAN

Telephone:

81 - 3 - 38165851

Fav

81 - 3 - 38162941

E-mail:

int@keiseimed.com

DUNS Number:

Official Correspondent Information

Contact Name: Junya Ohyama

Company:

KEISEI MEDICAL INDUSTRIAL CO., LTD.

Address:

3-19-6 Hongo

Bunkyo-ku, TOKYO, 1130033, JAPAN

Telephone:

81 - 3 - 38165851

Fax

81 - 3 - 38162941

E-mail:

johyama@keiseimed.com

DUNS Number:

United States Agent Information

Contact First Name

Alan

Contact Last Name

Schwaartz

Contact Title:

Mr

Business Name:

mdi Consultants, Inc..

Address:

55 Northern Blvd

Great Neck, California, 11021, UNITED STATES

Phone:

516 - 4829001

Fax:

516 -4820186

DUNS Number:

E-mail:

Alan@mdiconsultants.com

Device Listings

Listing Number	Premarket Submission Number	Premarket Submission Type	Product Code(s)	Device Name(s)	Activities	Importers
R022361		510(k) exempt	KDC	INSTRUMENT, SURGICAL, DISPOSABLE	Manufacturer	
A431019		510(k) exempt	INX	BED, AIR FLUIDIZED	Manufacturer	
D013619	K993970		GAR	Suture, nonabsorbable, synthetic, polyamide	Manufacturer	
E173215		510(k) exempt	EPT	Microscope, surgical	Manufacturer	
D116286		510(k) exempt	GFD	Dermatome	Manufacturer	