2020

CERTIFICATE OF REGISTRATION

This certifies that:

VEPRO AG
MAX-PLANCK STR. 1-3
PFUNGSTADT Hessen, GERMANY 64319

is registered with the U.S. Food and Drug Administration for FY 2020 pursuant to Title 21, 807 et seq. of the United States Code of Federal Regulations:

Establishment Registration:	**9614981**
DUNS No.:	**32-166-8584**
Device Classification Name:	**SYSTEM, IMAGE PROCESSING, RADIOLOGICAL**
Product Code:	**LLZ**
Regulation Number:	**892.2050**
Official Correspondent and U.S. Agent:	**Registrar Corp** 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until the end of the year stated above, unless said registration is terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. Registrar Corp assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.39, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding."

The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lennarz
Executive Director
Registrar Corp
Dated: November 19, 2019

©2005-2020 Registrar Corp

Establishment Registration & Device Listing

New Search | Back To Search Results

Establishment:
VEPRO AG
MAX-PLANCK STR. 1-3
PFUNGSTADT Hesse, DE 64319
Registration Number: 9614981
FEI Number*: 3004562407
Status: Active
Date Of Registration Status: 2024

Owner/Operator:
VEPRO AG
MAX-PLANCK STR. 1-3
PFUNGSTADT, DE 64319
Owner/Operator Number: 9030770

Official Correspondent:
David Lennarz
Registrar Corp
144 Research Drive
Hampton, VA 23666
Phone: 1-757-2240177

US Agent:
David Lennarz
Registrar Corp
144 Research Drive
Hampton, VA US 23666
Phone: 757 2240177 Ext
Fax: 757 2240179
Email: David.Lennarz@Registrarcorp.Com

* Firm Establishment Identifier (FEI) should be used for identification of entities within the imports message set

Vepro Ag

FDA Filings

This page includes the latest FDA filings for Vepro Ag. Currently, you will find the latest 100 filings for Premarket Notifications, Premarket Applications, De Novo Applications, and GUDID registrations.

FDA Registration(s)

Registration Number	9614981
FEI Number	3004562407
Name	VEPRO AG
Owner & Operator	VEPRO AG
Contact Address	MAX-PLANCK STR. 1-3 -- PFUNGSTADT DE-NOTA 64319 DE
US Agent	• David Lennarz • Registrar Corp • 757 2240177 • 757 2240179 • david.lennarz@registrarcorp.com
Importing Logistics Registration	Importer: VEPRO North America, Inc. Address: 8459 Little Rock Way Unit 204 Highlands Ranch, CO 80126 UNITED STATES Importer Type: Agent
Registration Status	1
Initial Importer	N
Registration Expiration	2020-04-25
Registration Address	MAX-PLANCK STR. 1-3 PFUNGSTADT Hessen, 64319 DE
Establishment Type	Manufacture Medical Device

FDA Filings

Device	Company	Device	Date
PMN K972215 | LLZ	VEPRO AG	EMR Manager	2008-10-31

FDA.report PMA PMN De Novo MAUDE GUDID NDC DailyMed Drug Applications

MEDIMAGE

System, Image Processing, Radiological

VEPRO - COMPUTERSYSTEME GMBH

The following data is part of a premarket notification filed by Vepro - Computersysteme Gmbh with the FDA for Medimage.

Pre-market Notification Details

Device ID	K972215
510k Number	K972215
Device Name:	MEDIMAGE
Classification	System, Image Processing, Radiological
Applicant	VEPRO - COMPUTERSYSTEME GMBH AN DER TUCHBLEICHE 26 Pfungstadt, DE D-64319
Contact	Harald G Roth
Correspondent	Harald G Roth VEPRO - COMPUTERSYSTEME GMBH AN DER TUCHBLEICHE 26 Pfungstadt, DE D-64319
Product Code	LLZ
CFR Regulation Number	892.2050 [🔎]
Decision	Substantially Equivalent (SESE)
Type	Traditional
3rd Party Reviewed	No
Combination Product	No
Date Received	1997-06-13
Decision Date	1997-11-19
Summary:	summary