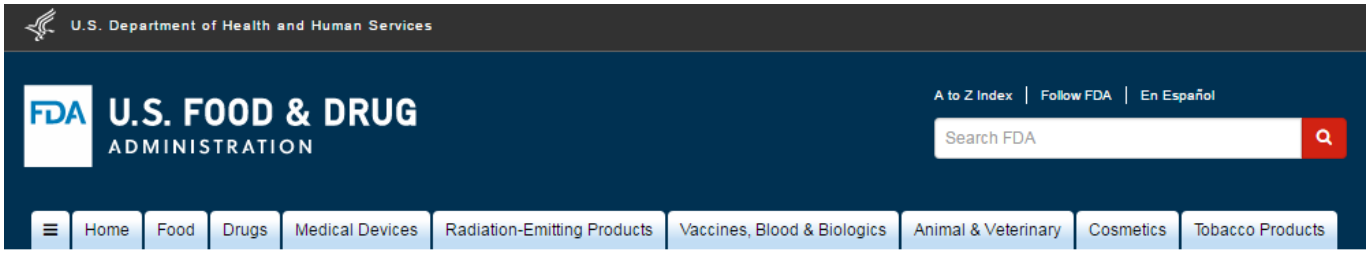
Pfungstadt, 19. October 2016

VEPRO *FDA Certification 2017.*

As an important acknowledgment for its successful quality assurance, VEPRO received the extension of its worldwide certification by FDA*.

This certification differentiates us from many other suppliers and document the high standard of our products.



US FDA 510 (k) Device Registration

Facility Information

Registration Number:	9614981
Facility Name:	VEPRO AG
Address:	MAX-PLANCK-STR. 1-3, PFUNGSTADT, Hessen, 64319, GERMANY
Device Name:	PACS - Radiological image processing
Listing Number:	D061761
Premarket Subm.no.:	K972215
Product Code:	LLZ

Registration and listing information for 2017 wass successfully updated and will be valid through Dec 31, 2017.

The next registration renewal period is October 1 – December 31, 2017.

* The FDA (Food and Drug Administration) is a department of the U.S. Department of Health. It is the "official food inspection and Drug Administration" for the United States and serves to protect the public health.

About VEPRO AG Germany

VEPRO AG is a leading international healthcare company with its own product and software development, "Made in Germany". With over 30 years of national and international presence in the solutions business, VEPRO AG is one of the most experienced providers in the healthcare industry.

The aim of the VEPRO concept is cost minimization in the hospital and hospital administration while optimizing patient care. This is achieved through a wide range of processes to improve the archiving of image, film and health report data. All information is available via internal networks, but also in external networks (cloud) within seconds.

Worldwide with its own subsidiaries or affiliates more than 4,000 clients are served around the clock. The company is headquartered in Pfungstadt close to Frankfurt (Germany).

Responsible for press releases

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