



EVPU[®]

NOTIFIED BODY No. 1293

EC Certificate Full Quality Assurance System

Directive 98/79/EC on In Vitro Diagnostic Medical Devices of the European Parliament and of the Council (IVDD), **Annex IV excluding (4, 6) (Module H)** transposed into "Slovak government decree No. 569/2001 Collection of Laws" as amended

No. 40012/101/1/2014/CE

EVPU a.s., Notified Body No. 1293, has audited the quality system in accordance with IVDD Annex IV and found that quality system meets the requirements of IVDD Annex IV.

Manufacturer and Facility

Beijing Wantai Biological Pharmacy Enterprise Co., Ltd.
No. 31 Life Science Park Road, Changping District
Beijing 102206, China

Device(s)

In Vitro Diagnostic Medical Device
Human Immunodeficiency Virus 1+2 Antibody Elisa Kit
(3rd generation)

Type(s)

Aid[™] anti-HIV 1+2 ELISA

Product code(s)

WI-4396 / WI-43480

Device(s)

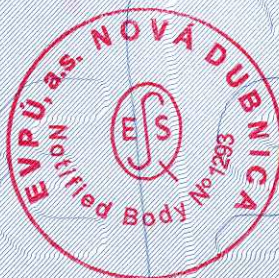
in List A

Relevant report(s)

40012/2014/C

Audit report(s)

M030/10-4




Marek Hudák

Issued on May 6th, 2014

Valid until May 5th, 2019

Manufacturer can affix the CE mark with number of Notified Body only in case devices are in comply with all relevant and effective Directives of European Parliament and of the Council. For placing on the market of List A devices covered by this certificate an EC Design-Examination Certificate according to IVDD Annex IV (4) is required. Surveillance audits according to IVDD, Annex IV (5) will be held to verify the validity of this Certificate.

The manufacturer must inform EVPU a.s. of any plan for substantial changes in the design of the device(s), in construction of the device(s) or in the quality system of production in order to examine whether this Certificate remains valid.

This Certificate is valid until the date specified. Any significant changes in the design of the device(s), in construction of the device(s), in the quality system or amendments to the Directive 98/79/EC may render this Certificate invalid at an earlier date. The product liability rests with the manufacturer or his representative in accordance with the Directive 85/374/EEC.

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