

## Certificate of CE-Notification

This is to certify that, in accordance with the *In Vitro* Diagnostic Medical Device Directive 98/79/EC, **CEpartner4U BV** agrees to perform all duties and responsibilities as the Authorized Representative for

### **CJSC EKOlabor**

1 Budennogo Str., Elektrogorsk, Moscow region, 142530, Russia

as stipulated and demanded by the aforementioned Directive. The Dutch Competent Authorities have accepted the manufacturer's medical device registrations by CEpartner4U as listed on the product list attached to the manufacturer's Declaration of Conformity:

### **Device group: Rabbit plasma**

**IVD devices were registered under number:**

**Registration number Rabbit plasma: NL-CA002-2017-43242**

**with Dutch Competent Authorities as a consequently this IVD devices were entered in EUDAMED by Dutch Competent Authorities**

The manufacturer has provided CEpartner4U with all necessary documentation, together with an appropriate Declaration of Conformity that the IVD medical devices fulfil the essential requirements of Directive 98/79/EC.

2017-12-18



Olga Teirlinck  
Consultant CEpartner4U BV

**C e p a r t n e r 4 U**

**Esdoornlaan13  
3951 DB Maarn NL  
tel: +31 (0)343 442 524  
[www.cepartner4u.nl](http://www.cepartner4u.nl)**