ViroGates

K-13-0342-01

Certificate of Analysis

For Standards and Curve Control used in suPARnostic® AUTO Flex ELISA, code no. E001

Intended use
For in vitro diagnostic use. For professional use only.
suPARnostic® standards are intended for establishing standard curve for the quantitative immunological determination of human suPAR (soluble urokinase plasminogen activator receptor) by ELISA.

Standards a-e
Recombinant suPAR (soluble urokinase plasminogen activator receptor) in PBS buffer with proprietary additives and 0.05% Bronidox® as preservative. The standards are intended to be used in combination with the reagents provided with suPARnostic AUTO Flex ELISA code E001.

KIT LOT XS 2473 2020-09

Reagents		LOT		Concentration	Range
Standards	a	XS 2682	2020-09	14.3 ng/mL	
Standards	b	XS 2683	2020-09	11.2 ng/mL	
Standards	c	XS 2684	2020-09	7.9 ng/mL	
Standards	d	XS 2685	2020-09	4.4 ng/mL	
Standards	e	XS 2686	2020-09	0.7 ng/mL	
Curve Control		XS 2687	2020-09	2.7 ng/mL	2.0-3.4 ng/mL

PRECAUTIONS AND RECOMMENDATIONS
Do not use components beyond indicated expiration date.
Do not mix reagents from different kits.

Calculation of suPAR values from Raw OD´s: Use the Calculation tool software available on www.virogates.com or contact info@virogates.com

Date of approval: 2018-11-14

Quality Control

Quality Assurance

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