

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 ZS 1953  2021-12

Reagents	LOT		Concentration	Range
Standards	a	AS 1131	2021-12	13.2 ng/mL
Standards	b	AS 1132	2021-12	10.3 ng/mL
Standards	c	AS 1133	2021-12	7.3 ng/mL
Standards	d	AS 1134	2021-12	4.0 ng/mL
Standards	e	AS 1135	2021-12	0.7 ng/mL
Curve Control	AS 1136	2021-12	2.5 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: 2020-07-22


 Quality Control


 Quality Assurance

