

DATA SHEET

Factor Xa 71 NKAT

PROPERTIES

Factor X is purified from a barium citrate elution of bovine plasma and it is activated to FXa by a Russell's Viper Venom matrix.

COMPOSITION

Each vial contains:

Art. No. 82 0985 39

Factor Xa:	71±10 nkat	
Bovine Albumin:	20 mg	S-2222
Polyethyleneglycol 8000:	10 mg	
Sodium Chloride:	10 mg	
Tris (hydroxymethyl) aminomethane:	6.4 mg	Tris (hydroxymethyl) aminomethane HCl: 69 mg

WARNINGS



Danger

Hazard class: H334, Resp Sens. 1

Hazard statements: H334: May cause allergy or asthma symptoms or breathing difficulties if inhaled.

Precautionary statements: P261: Avoid breathing dust/fume. P284: [In case of inadequate ventilation] wear respiratory protection. P304+P340: IF INHALED: Remove person to fresh air and keep comfortable for breathing. P342+P311: If experiencing respiratory symptoms: Call a POISON CEN- TER/doctor. P501: Dispose of contents/container in accordance with local/ regional/national/international regulation.

Supplemental hazard information: Contains Factor Xa. Up to 11.3% of the mixture consists of components of unknown acute toxicity (oral, dermal, inhalation) for the human health. Up to 7.4% of the mixture consists of components of unknown hazard to the aquatic environment.

STORAGE

The lyophilized powder is stable at 2-8°C until the expiry date printed on the label.

RECONSTITUTION

Reconstitute with 10 ml sterile water to obtain an activity of 7 nkat/ml (reaction buffer: 0.05 mol/l Tris buffer pH 8.4).

STABILITY

The reconstituted Factor Xa is stable for 1 month at 2-8°C or 6 months at

-20°C when used for the determination of Heparin, according to the test conditions of the Heparin kit.

APPLICATION

For the determination of Heparin in plasma.

ENZYME ACTIVITY DETERMINATION

The activity of FXa can be measured using the following chromogenic substrates:

Substrates

S-2222, 25 mg vial

S-2765, 25 mg vial

S-2772, 26 mg vial

For In-Vitro Diagnostic Use. Not For Human Or Animal Consumption.