

Human NG2/MCSP/CSPG4 ELISA Instructions Cat:EHY0162

Content

	CAT	Volume
CP (Coated Plate)	EHY0162CP	96 well
2 S (Standard)	EHY0162S1~S7,S0	8 vial
3 DA-H (Detect Antibody-HRP)	EHY0162DA-H	6 ml/bottle
4 AB (Assay Buffer 1×)	EAB01	12 ml/bottle
5 TS (TMB Substrate)	ETS01	12 ml/bottle
(3 SS (Stop Solution)	ESS01	12 ml/bottle
WB (Wash Buffer 10×)	EWB01	50 ml/bottle
8 SF (Sealer Film)	ESF01	6 pieces

NOTE: After the kit is opened, the stabilization period of each content is 30 days, so please use the kit within 30 days after opening.

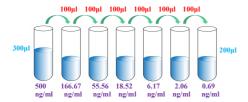
REAGENT PREPARATION

Washing Buffer (1×) Preparation

Pour entire contents (50 ml) of the Washing Buffer Concentrate (10×) into a clean 500 ml graduated cylinder. Bring to final volume of 500 ml with glass-distilled or deionized water. Transfer to a clean wash bottle and store at 2 to 25°C.

Standard Curve Preparation:

 ${\sf S1}$ to ${\sf S7}$ and ${\sf S0}$ is ready to use for serum and plasma.



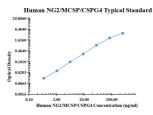
ASSAY PROCEDURE

Bring all reagents and samples to room temperature before use.

- 1 Prepare all reagents and working standards as directed in the previous sections.
- 2 Remove excess CP (Coated Plate) strips from the plate frame, return them to the foil pouch and reseal.
- 3 Add 50 μl of AB (Assay Buffer) to each well.
- 4 Add 50 μl or 10 μl of Standard or sample per well. Ensure reagent addition is uninterrupted and completed within 15 minutes.
- **5** Add 50 μl of **DA-H** (Detect Antibody-HRP) to each well.
- **6** Cover with an **SF** (Sealer Film). Incubate at room temperature (18 to 25°C) for 30 min on a microplate **shaker** set at 500 rpm.
- 8 Add 100 μl of TS (TMB Substrate) to each well. Incubate for 5-30 minutes at room temperature.
- 9 Add 100 μl of SS (Stop Solution) to each well.
- **10** Determine the optical density within 30 minutes, using microplate **reader** set to 450 nm corrected with 570 nm or 630 nm.



TYPICAL DATA



ng/ml	Ο.	O.D.		Corrected
0.00	0.0144	0.0150	0.0147	
0.69	0.0266	0.0269	0.0268	0.0121
2.06	0.0461	0.0456	0.0459	0.0312
6.17	0.1073	0.1186	0.1130	0.0983
18.52	0.2726	0.3004	0.2865	0.2718
55.56	0.8309	0.8644	0.8477	0.8330
166.67	2.0830	2.1660	2.1245	2.1098
500.00	3.8140	3.9010	3.8575	3.8428

SENSITIVITY

The minimum detectable dose (MDD) of human NG2/MCSP/CSPG4 is typically less than 0.004 ng/ml (50 μ l of sample volume) or 0.21 ng/ml (10 μ l of sample volume).

The MDD was determined by adding two standard deviations to the mean optical density value of ten zero standard replicates and calculating the corresponding concentration.

PRECISION

Intra-assay Precision (Precision within an assay) Three samples of known concentration were tested twenty times on one plate to assess intra-assay precision.

Inter-assay Precision (Precision between assays)

	Intra-assay Precision			Inter-assay Precision		
Sample	S1	S2	S3	S1	S2	S3
Number	22	22	22	6	6	6
Average (ng/ml)	1.7	8.6	30.9	1.6	8.9	31.6
Standard Deviation	0.1	0.4	2.0	0.1	0.6	1.8
Coefficient of Variation (%)	5.7	4.4	6.5	6.0	6.8	5.7

RECOVERY

The spike recovery was evaluated by spiking 3 levels of human NG2/MCSP/CSPG4 into health human serum sample. The un-spiked serum was used as blank in this experiment.

The recovery ranged from 99% to 117% with an overall mean recovery of 108%.

LINEARITY

To assess the linearity of the assay, five samples were spiked with high concentration of NG2/MCSP/CSPG4 in human serum and diluted with Sample Diluent to produce samples with values within the dynamic range of the assay.

The linearity ranged from 99% to 109% with an overall mean recovery of 103%.

SAMPLE VALUES

Serum/Plasma – Thirty samples from apparently healthy volunteers were evaluated for the presence of human NG2/MCSP/CSPG4 in this assay. No medical histories were available for the donors.

Sample Matrix	Sample Evaluated	Range (ng/ml)	Detectable %	Mean of Detectable (ng/ml)
Serum	30	59.50-274.06	100.0	129.52

n.d. = non-detectable. Samples measured below the sensitivity are considered to be non-detectable.