

PRODUCT INFORMATION AND MANUAL

***Human sICAM-1
Matched Antibody Pairs
for ELISA***

BMS201MST

Module Set for the development of an enzyme-linked immunosorbent assay for quantitative detection of human sICAM-1.

For research use only.

Not for diagnostic or therapeutic procedures.



**Human sICAM-1
BMS201MST**

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READ BEFORE OPENING

- **Some vials contain small quantities of material, therefore centrifuge before use.**
- **This set of reagents is intended for use by persons experienced in the use of immunoassays. It is not suitable for use by inexperienced personnel.**
- **A sample protocol is included but please note that the protocol provided is a guideline. The type of substrate as well as all other reagents not included in the Module Set may influence assay performance.**

1 Reagents Provided

- 1 vial (3.85 ml) monoclonal **Coating Antibody** to human sICAM-1 (100 µg/ml)
- 1 vial (55 µl) **HRP-Conjugate** anti-human sICAM-1 monoclonal antibody
- 1 vial (50 µl) human sICAM-1 **Standard** protein, 10 µg/ml
- 2 vials (50 ml) **Sample Diluent**

2 Storage Instructions

Store kit components at -20 °C. Immediately after use remaining reagents should be returned to -20 °C storage, respectively. Avoid multiple freeze-thaw cycles. Aliquot reagents for repeated use at later dates. Reagents are labeled with expiration date.

Samples should be aliquoted and must be stored frozen at -20 °C to avoid loss of bioactive human sICAM-1.

3 Reagents and Materials Not Provided

Buffers and solutions can be prepared (see chapter 5) or ordered at Bender MedSystems. We recommend the use of BMS-reagents to ensure a high quality assay performance.

Reagents and Materials	BMS Order Numbers
Phosphate Buffered Saline (PBS)	BMS410.0010, 20x (10 ml)
Assay Buffer	BMS407.0050, 20x (50 ml):
Wash Buffer	BMS408.0500, 20x (500 ml):
Substrate Solution	BMS406.0100, ready to use (100 ml)
Stop Solution	BMS409.0100, ready to use (100 ml)
Microwell plate (We recommend F8 Immuno Modules Maxi sorb, NUNC Order Number 268667)	BMSPLATE

4 Precautions for Use

All reagents should be considered as potentially hazardous. We therefore recommend that this product is handled only by those persons who have been trained in laboratory techniques and that it is used in accordance with the principles of good laboratory practice. Wear suitable protective clothing such as laboratory overalls, safety glasses and gloves. Care should be taken to avoid contact with skin or eyes. In the case of contact with skin or eyes wash immediately with water. See material safety data sheet(s) and/or safety statement(s) for specific advice.

5 Preparation of Buffers and Solutions

Please note: The **quality of BSA** is a critical parameter for the test performance. We recommend Probumin™ Media Grade (e.g.: Millipore Order Number 81-068) or the use of the BMS-reagents (see chapter 3).

5.1 Phosphate Buffered Saline (PBS)

NaCl	8.00 g
KCl	0.20 g
Na ₂ HPO ₄ x 12 H ₂ O	2.85 g
KH ₂ PO ₄	0.20 g
H ₂ O _{dest}	adjust to 1 liter

5.2 Wash Buffer

Add 0.5 ml Tween 20 to 1 liter of PBS and mix well.

5.3 Assay Buffer

Bovine Serum Albumin (BSA)	5 g
Tween 20	0.5 ml
PBS	adjust to 1 liter

5.4 Fixing Buffer

Sucrose	75 g
PBS	adjust to 500 ml

5.5 Substrate Solution

1:2 mixture of H₂O₂ and Tetramethylbenzidine

5.6 Stop Solution

1M Phosphoric Acid (H₃PO₄)

6 Preparation of the Microwell Plate

6.1 Coating

Coating antibody final concentration is 3.5 µg/ml; 100 µl of the coating solution is added to each well. Dilute the coating antibody as following for one microtiter plate:

10.615 ml	PBS
0.385 ml	coating antibody (100 µg/ml)
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11 ml	coating solution (3.5 µg/ml)

Immediately after coating, seal the plate with an adhesive film and store at 2-8°C over night, allowing the binding process to take place. Aspirate the contents of the wells and wash once with 400 µl of Wash Buffer according the washing procedure described in the test protocol below (see 8.h).

6.2 Blocking and Fixing

6.2.1 Blocking

Add 250 µl of Assay Buffer to each well and incubate at room temperature for 2 hours. Alternatively the plate may be blocked over night at 2-8°C. Blocked plates can be stored at 2-8°C for up to one week. For immediate use after blocking wash the plate twice (see 8.h) before the samples are applied to the wells.

6.2.2 Fixing

To store the coated plates for longer than one week aspirate Assay Buffer and add 150 µl Fixing Buffer to each well. Incubate for 1 hour at room temperature. Aspirate Fixing Buffer and dry plates over night at 28°C. When sealed with desiccant, the plates can be stored at 2-8°C for 2 months.

7 Preparation of Immunological Reagents

Please note: Centrifuge vials before opening to collect contents.

7.1 Preparation of Standard:

The concentrated **human sICAM-1 standard** must be diluted 1:100 with Assay Buffer just prior to use in a clean plastic test tube according to the following dilution scheme:

5 μ l	conc. Standard Protein (10 μ g/ml)
495 μ l	Assay Buffer
<hr/>	
500 μ l	Standard Protein (100 ng/ml)

Shake gently to mix. After usage remaining diluted standard cannot be stored and has to be discarded.

Aliquot the concentrated standard and store at -20 °C.

7.2 Preparation of HRP-Conjugate

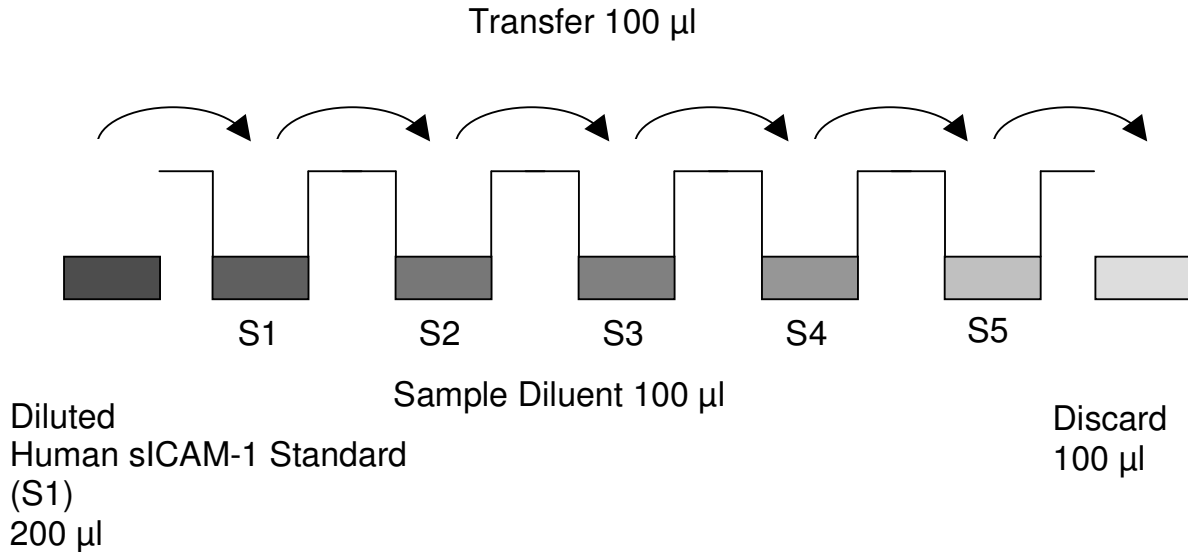
Dilute conc. **HRP-Conjugate** 1:1000 with Assay Buffer before use. Use within 30 min after preparation. For one microwell plate dilute the stock reagents as follows:

5.5 μ l	conc. HRP-Conjugate
5494.5 μ l	Assay Buffer
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5.5 ml	HRP-Conjugate

8 Test Protocol

- a. Add 100 μl of Sample Diluent in duplicate to **standard wells** B1/2-E1/2, leaving A1/A2 empty. Pipette 200 μl of diluted **standard** (see 7.1), (concentration of standard 1, S1 = 100 ng/ml) in duplicate into well A1 and A2. Transfer 100 μl to wells B1 and B2. Mix the contents of wells B1 and B2 by repeated aspiration and ejection, and transfer 100 μl to wells C1 and C2, respectively (see Figure 1). Take care not to scratch the surface of the microwells. Continue this procedure 2 times, creating two serially diluted columns of human sICAM-1 standard dilutions ranging from 100 to 6.25 ng/ml. Discard 100 μl from the last microwells (E1, E2). Final volume in all wells is 100 μl .

Figure 1



- b. Add 100 μ l of **Sample Diluent** in duplicate to **blank wells**.
- c. Add 90 μ l of **Sample Diluent** to the **sample wells**.
- d. Add 10 μ l of each **sample** in duplicate to the **sample wells**.
- e. Prepare **HRP-Conjugate** (see 7.2)
- f. Add 50 μ l of prepared **HRP-Conjugate** to all wells.
- g. Cover with an adhesive film and incubate at room temperature (18 to 25°C) for 1 hour, on a microplate shaker set at 100 rpm if available.
- h. Remove adhesive film and empty wells. Wash microwell 3 times with approximately 400 μ l **Wash Buffer** per well, with thorough aspiration of microwell contents between washes. Allow the Wash Buffer to sit in the wells for about **10 – 15 seconds** before aspiration. Take care not to scratch the surface of the microwells.
After the last wash step, empty wells and tap microwell strips on absorbent pad or paper towel to remove excess Wash Buffer. Use the microwell plate immediately after washing. Alternatively microwell plate can be placed upside down on a wet absorbent paper for no longer than 15 minutes. **Do not allow wells to dry.**
- i. Pipette 100 μ l of **Substrate Solution** to all wells.
- j. Incubate the microwell strips at room temperature (18° to 25°C) for about 10 min. Avoid direct exposure to intense light.

Monitor the colour development on the plate. The substrate reaction should be stopped (see next point of this protocol) before positive wells are no longer properly recordable. Determination of the ideal time period for colour development has to be done individually for each assay.

Add the stop solution when the highest standard has developed a dark blue colour. Alternatively the colour development can be monitored on a plate reader at 620 nm. The substrate reaction should be stopped as soon as Standard 1 has reached an OD of 0.9 – 0.95.

- k. Stop the enzyme reaction by quickly pipetting 100 μ l of **Stop Solution** into each well. It is important that the Stop Solution is spread quickly and uniformly throughout the microwells to completely inactivate the enzyme. Results must be read immediately after the Stop Solution is added, or within one hour if the microwell strips are stored at 2 - 8°C in the dark.
- l. Read absorbance of each microwell on a spectro-photometer using 450 nm as primary wave length (you can use 620 nm as reference wave length; 610 nm to 650 nm is acceptable). Blank the plate reader according to the manufacturer's instructions by using the blank wells. Determine the absorbance of both the samples and the human sICAM-1 standards.

9 Calculation of Results

- Calculate the average absorbance values for each set of duplicate standards and samples. Duplicates should be within 20 per cent of the mean.
- Create a standard curve by plotting the mean absorbance for each standard concentration on the y-axis, against the human sICAM-1 concentration on the x-axis. Draw a best fit curve through the points of the graph (a 5-parameter curve fit is recommended).
- To determine the concentration of soluble human sICAM-1 for each sample, first calculate the mean absorbance value for the duplicate wells of the sample, then extend a horizontal line from this point on the y-axis to the standard curve. At the point of intersection, extend a vertical line to the x-axis and read the corresponding human sICAM-1 concentration.
- **If instructions in this protocol have been followed samples have been diluted 1:10 (10 μ l sample + 90 μ l Sample Diluent), the concentration read from the standard curve must be multiplied by the dilution factor (x 10).**
- **Calculation of samples with a concentration exceeding that of standard 1 will result in inaccurate, low human sICAM-1 levels (Hook Effect). Such samples require further external predilution according to expected human sICAM-1 values with Sample Diluent in order to precisely quantitate the actual human sICAM-1 level.**
- Each testing facility should establish a control sample of known human sICAM-1 concentration and run this additional control with each assay. If the values obtained are not within the expected range of this control, the assay results may be invalid.

A basic understanding of immunoassay development and technical experience in ELISA performance are conditional for the successful use of this Module Set.

The protocol provided is just a guideline. The type of substrate as well as all other reagents not included in the Module Set may influence the test characteristics.

10 Human sICAM-1 Module Set Characteristics

10.1 Specificity

The assay detects both natural and recombinant human sICAM-1. The interference of circulating factors of the immune system was evaluated by spiking these proteins at physiologically relevant concentrations into human sICAM-1 positive serum.

There was no cross reactivity detected with sTNF-R (60 kDa), sTNF-R (80 kDa), IL-8/NAP-1, TNF α , TNF β , IFN γ , IFN α 2C, IFN ω , IL-6, IL-2R, ELAM-1 and L-selectin.

10.2 Expected Values

A panel of 50 sera samples from randomly selected apparently healthy donors (males and females) was tested for human sICAM-1.

The detected human sICAM-1 levels ranged between 129.9 and 297.4 ng/ml with a mean level of 230.3 ng/ml and a standard deviation of 47.4 ng/ml.

Normal human sICAM-1 levels may vary depending on the serum collective used ranging up to 400 ng/ml.

11 Additionally available Components

Additional components of the human sICAM-1 MST Kit are available:

- Standard Protein (S): BMS201MSTS (for 10 assays) or BMS201S (for 1 assay)
- Coating Antibody (CA): BMS201MSTCA
- Sample Diluent (SD): BMS201MSTSD
- HRP-Conjugate (K): BMS201MSTK

Please contact Bender MedSystems at techserv@bendermedsystems.com or your local distributor.

12 Ordering Information

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