

T-Select MHC class I human Tetramer

# HLA-A\*24:02 CA<sub>9</sub><sub>219-227</sub> Tetramer

## -EYRALQLHL (50 tests)

For Research Use Only. Not for use in diagnostic procedures.  
MBL manufactures and distributes these products under license from Beckman Coulter, Inc..

### Background

T lymphocytes play a central role in immune system. Total T cell and T cell subset counts are measured by detection of various cell surface molecules. Enumeration of CD8<sup>+</sup> antigen-specific T cells requires cognate recognition of the T cell receptor (TCR) by a class I MHC/peptide complex. This can be done using T-Select MHC class I Tetramers which are composed of four MHC class I molecules each bound to the specific peptide and conjugated with a fluorescent protein. Thus, T-Select MHC Tetramer assays allow quantitation of the total T cell population specific for a given peptide complexed in a particular MHC molecule. Furthermore, since binding does not depend on functional pathways, this population includes specific CD8<sup>+</sup> T cells regardless of functional status. Measurements may be performed in whole blood or isolated lymphocyte/mononuclear cell preparations. In some cases where frequency is low, it may be necessary to perform an *in vitro* cell expansion. Specific cell staining is accomplished by incubating the sample with the T-Select MHC Tetramer reagent, then washing away excess Tetramer. The number of Tetramer positive lymphocytes is then determined by flow cytometry.

This Tetramer reagent comprises human class I HLA-A\*24:02 and epitope peptide derived from Carbonic Anhydrase IX (CA<sub>9</sub>), and it can detect HLA-A\*24:02-restricted CA<sub>9</sub><sub>219-227</sub>-specific CD8<sup>+</sup> T cells by flow cytometry.

CA<sub>9</sub> plays a key role in maintaining cellular pH and is a specific molecular marker for renal cell carcinoma (RCC)<sup>1</sup>. The HLA-A\*24:02-restricted CA<sub>9</sub>-specific CTL epitope was initially identified by Dr. Kazuhiro Yoshikawa and his colleagues at Nara Medical University<sup>2</sup>. CA<sub>9</sub> is considered as new potential therapeutic target using several strategies including antibody-targeting, vaccine-based immunotherapy<sup>3</sup>, and small molecule inhibitors.

**HLA Restriction:** HLA-A\*24:02

### Origin and Sequence of CTL Epitope:

CA<sub>9</sub> (219-227 aa, EYRALQLHL)

### References for This Product

- 1) Pastorek J *et al. Oncogene* **9**: 2877-2888 (1994)
- 2) Shimizu K, *et al. Oncol Rep* **10**: 1307-1311 (2003)
- 3) Uemura H, *et al. Clin Cancer Res.* **12**: 1768-1775 (2006)

### High Specificity

The T cell surface CD8 enhances T cell antigen recognition by binding to HLA class I molecules. Therefore, MBL produced T-Select MHC class I human Tetramers with one point mutation at the HLA  $\alpha$ 3 domain known to alter the interaction with CD8. These mutated Tetramers showed a greatly diminished nonspecific binding but retained specific binding. Alterations of CD8 binding by mutation of the MHC greatly improved the specificity of MHC-peptide multimers, thus providing efficient tools to sort specific human T cells for immunotherapy.  
(French application Number; FR9911133)

### References for T-Select MHC Tetramer

- Altman JD, *et al. Science* **274**: 94-96 (1996)  
McMichael AJ, *et al. J Exp Med* **187**: 1367-1371 (1998)  
Bodinier M, *et al. Nat Med* **6**: 707-710 (2000)

### Reagents

T-Select MHC Class I Human Tetramer - 50 tests  
The Tetramer is dissolved in an aqueous buffer containing 0.5 mM EDTA, 0.2% BSA, 10 mM Tris-HCl (pH 8.0), 150 mM NaCl, and 0.09% NaN<sub>3</sub>.

### Conjugates

- TS-M112-1  
Streptavidin-Phycoerythrin (SA-PE)  
Excites at 486-580 nm  
Emits at 586-590 nm
- TS-M112-2  
Streptavidin-Allophycocyanin (SA-APC)  
Excites at 633-635 nm  
Emits at 660-680 nm

### Storage Conditions

Store at 2 to 8°C. Do not freeze. Minimize exposure to light. The expiration date is indicated on the vial label.

### Evidence of Deterioration

Any change in the physical appearance of this reagent may indicate deterioration and the reagent should not be used. The normal appearance is a clear, colorless to pink (SA-PE), or light blue (SA-APC).

### Usage

This reagent is for use with standard flow cytometry methodologies.

### Statement of Warnings

1. This reagent contains 0.09% sodium azide. Sodium azide under acid conditions yields hydrazoic acid, an extremely toxic compound. Azide compounds should be flushed with running water while being discarded. These precautions are recommended to avoid deposits in metal piping in which explosive conditions can develop. If skin or eye contact occurs, wash excessively with water.
2. Specimens, samples and material coming in contact with them should be handled as if capable of transmitting infection and disposed of with proper precautions.
3. Never pipette by mouth and avoid contact of samples with skin and mucous membranes.
4. Minimize exposure of reagent to light during storage or incubation.
5. Avoid microbial contamination of reagent or erroneous results may occur.
6. Use Good Laboratory Practices (GLP) when handling this reagent.

### Materials Required But Not Supplied

- 12 x 75 mm polypropylene test tubes
- Transfer pipettes
- Pipettors and disposable pipette tips
- Vortex mixer
- Centrifuge capable of 150 x g or 400 x g
- Aspirator
- PBS
- MHC Tetramer Lyse Reagent, MBLI, PN T08002
- MHC Tetramer Fixative Reagent, MBLI, PN T08003
- Anti-CD8-FITC, Beckman Coulter, Inc., PN 6603861
- 7-AAD Viability Dye, Beckman Coulter, Inc., PN A07704
- Clear Back (human FcR blocking reagent), MBL, PN MTG-001

### Procedure for Whole Blood

1. Collect blood by venipuncture into a blood collection tube containing an appropriate anti-coagulant.
2. Add 10  $\mu$ L of T-Select MHC Tetramer to each 12 x 75 mm test tube.
3. Add 200  $\mu$ L of whole blood into each test tube.
4. Vortex gently.
5. Incubate for 30-60 minutes at 2-8°C protected from light.

6. Add any additional antibodies (e.g. anti-CD8) and vortex gently.
7. Incubate for 30 minutes at 2-8°C protected from light.
8. Lyse red blood cells using 2 mL of Lyse Reagent supplemented with 50  $\mu$ L Fixative Reagent per tube.
9. Vortex for 5 seconds immediately after the addition of the Lyse/Fixative solution.
10. Incubate for a minimum of 10 minutes at room temperature protected from light.
11. Centrifuge tubes at 150 x g for 5 minutes.
12. Aspirate or decant the supernatant.
13. Add 3 mL of PBS and centrifuge tubes at 150 x g for 5 minutes.
14. Aspirate or decant the supernatant.
15. Resuspend the pellet in 500  $\mu$ L of PBS with 0.1% formaldehyde. (12.5  $\mu$ L Fixative Reagent/1 mL PBS).
16. Store prepared samples at 2-8°C protected from light for a minimum of 1 hour (maximum 24 hours) prior to analysis by flow cytometry.

### Procedure for Peripheral Blood Mononuclear Cells

1. Prepare peripheral blood mononuclear cells (PBMC) according to established procedures. Cells should be re-suspended at a concentration of  $2 \times 10^7$  cells/mL. 50  $\mu$ L of sample is required for each T-Select MHC Tetramer determination.
2. Add 10  $\mu$ L of Clear Back (human FcR blocking reagent, MBL, PN MTG-001) to each 12 x 75 mm test tube.
3. Add 50  $\mu$ L PBMC into each test tube (e.g.  $1 \times 10^6$  cells per tube).
4. Incubate for 5 minutes at room temperature.
5. Add 10  $\mu$ L of T-Select MHC Tetramer and vortex gently.
6. Incubate for 30-60 minutes at 2-8°C protected from light.
7. Add any additional antibodies (e.g. anti-CD8) and vortex gently.
8. Incubate for 30 minutes at 2-8°C protected from light.
9. Add 3 mL of PBS or FCM buffer (2% FCS/0.09% NaN<sub>3</sub>/PBS).
10. Centrifuge tubes at 400 x g for 5 minutes.
11. Aspirate or decant the supernatant.
12. Resuspend the pellet in 500  $\mu$ L of PBS with 0.5% formaldehyde. (62.5  $\mu$ L Fixative Reagent/1 mL PBS).
13. Store prepared samples at 2-8°C protected from light for a minimum of 1 hour (maximum 24 hours) prior to analysis by flow cytometry.

### Limitations

1. For optimal results with whole blood, retain specimens in blood collection tubes at room temperature, while rocking, prior to staining and analyzing. Refrigerated specimens may give aberrant results.

2. Recommended cell viability for venous blood specimens is > 90%.
3. Prolonged exposure of cells to lytic reagents may cause white blood cell destruction and loss of cells in the population of interest.
4. All red blood cells may not lyse under the following conditions: nucleated red blood cells, abnormal protein concentration or hemoglobinopathies. This may cause falsely decreased results due to unlysed red blood cells being counted as leukocytes.

## Technical Hints

- A. If PBMC culture is needed, we recommend the use of heparin as an anti-coagulant.
- B. Clear Back reagent (human FcR blocking reagent) may effectively block non-specific binding caused by macrophages or endocytosis, resulting in clear staining when cells are stained with MHC Tetramer and antibodies. Please refer to the data sheet (MBL, PN MTG-001) for details.
- C. A Tetramer that is constructed with the same allele of interest and an irrelevant peptide may be used as a negative control.
- D. We recommend the use of anti-CD8 antibody, clone SFC121Thy2D3 (T8, Beckman Coulter, Inc.), because some anti-CD8 antibodies inhibit Tetramer-specific binding to TCR.
- E. The use of CD45 antibody and gating of the lymphocyte population are recommended in order to reduce contamination of unlysed or nucleated red blood cells in the gate.
- F. Apoptotic, necrotic, and/or damaged cells are sources of interference in the analysis of viable cells by flow cytometry. Cell viability should be determined by 7-aminoactinomycin D (7-AAD) staining; intact viable cells remain unstained (negative).
- G. Cells do not require fixation prior to analysis if the stained cells are analyzed by flow cytometry within several hours.

## Related Products

### T-Select Human Tetramers

#### Cancer

TS-M112-1 HLA-A\*24:02 CA9 Tetramer-EYRALQLHL-PE  
TS-M080-1 HLA-A\*02:01 CEA Tetramer-YLSGADLNL-PE  
TS-M084-1 HLA-A\*02:01 EphA2 Tetramer-TLADFDP RV-PE  
TS-0014-1C HLA-A\*02:01 gp100 Tetramer-ITDQVPFSV-PE  
TS-0013-1C HLA-A\*02:01 gp100 Tetramer-IMDQVPFSV-PE  
TS-0035-1C HLA-A\*02:01 gp100 Tetramer-KTWGQYWQV-PE  
TS-M082-1 HLA-A\*02:01 gp100 Tetramer-YLEPGPVTA-PE  
TS-M089-1 HLA-A\*24:02 gp100 Tetramer-VYFFLPDHL-PE  
TS-0015-1C HLA-A\*02:01 Her-2/neu Tetramer-KIFGSLAFL-PE  
TS-0016-1 HLA-A\*02:01 Her-2/neu Tetramer-RLLQETELV-PE  
TS-M083-1 HLA-A\*02:01 HM1.24 Tetramer-KLQDASAEV-PE  
TS-M010-1 HLA-A\*24:02 hTERT Tetramer-VYGFVRACL-PE  
TS-M070-1 HLA-A\*02:01 MAGE-A1 Tetramer-KVLEYVIKV-PE  
TS-M071-1 HLA-B\*07:02 MAGE-A1 Tetramer-RVRFFPSL-PE

TS-M072-1 HLA-A\*02:01 MAGE-A2 Tetramer-YLQLVFGIEV-PE  
TS-M073-1 HLA-A\*24:02 MAGE-A2 Tetramer-EYLQLVFGI-PE  
TS-M074-1 HLA-A\*01:01 MAGE-A3 Tetramer-EVDPIGHLY-PE  
TS-M075-1 HLA-A\*02:01 MAGE-A3 Tetramer-KVAELVHFL-PE  
TS-M076-1 HLA-A\*02:01 MAGE-A3 Tetramer-FLWGPRLV-PE  
TS-M077-1 HLA-A\*24:02 MAGE-A3 Tetramer-IMPKAGLLI-PE  
TS-M078-1 HLA-A\*02:01 MAGE-A10 Tetramer-GLYDGMEHL-PE  
TS-0009-1C HLA-A\*02:01 Mart-1 Tetramer-ELAGIGILTV-PE  
TS-M091-1 HLA-A\*24:02 MCPyV Tetramer-EWWRSGGFSF-PE  
TS-M088-1 HLA-A\*02:01 MUC1 Tetramer-LLLTLVLTV-PE  
TS-M011-1 HLA-A\*02:01 NY-ESO-1 Tetramer-SLLMWITQC-PE  
TS-M109-1 HLA-B\*07:02 P2X5 Tetramer-TPNQRQNV-PE  
TS-M081-1 HLA-A\*02:01 p53 Tetramer-LLGRNSFEV-PE  
TS-M107-1 HLA-A\*02:01 PAP Tetramer-ALDVYNGLL-PE  
TS-0017-1 HLA-A\*02:01 PR-1 Tetramer-VLQELNVTV-PE  
TS-M087-1 HLA-A\*02:01 PSA Tetramer-KLQCVDLHV-PE  
TS-M079-1 HLA-A\*02:01 SSX-2 Tetramer-KASEKIFYV-PE  
TS-M085-1 HLA-A\*02:01 Survivin Tetramer-LMLGEFLKL-PE  
TS-M025-1 HLA-A\*24:02 survivin-2B Tetramer-AYACNTSTL-PE  
TS-0019-1C HLA-A\*02:01 tyrosinase Tetramer-YMDGTMSQV-PE  
TS-M090-1 HLA-A\*24:02 tyrosinase Tetramer-AFLPWHRFL-PE  
TS-M014-1 HLA-A\*24:02 WT1 (mutant) Tetramer-CYTWNQMNL-PE  
TS-M016-1 HLA-A\*02:01 WT1 Tetramer-RMFPNAPYL-PE

#### Control

TS-0029-1C HLA-A\*02:01 Negative Tetramer-PE  
TS-0029-2C HLA-A\*02:01 Negative Tetramer-APC  
TS-M007-1 HLA-A\*24:02 HIV env Tetramer-RYLRDQQLL-PE  
TS-M007-2 HLA-A\*24:02 HIV env Tetramer-RYLRDQQLL-APC  
TS-M007-3 HLA-A\*24:02 HIV env Tetramer-RYLRDQQLL-FITC  
TS-M054-1 HLA-B\*07:02 HIV nef Tetramer-TPGPGVRYPL-PE  
TS-M054-2 HLA-B\*07:02 HIV nef Tetramer-TPGPGVRYPL-APC

#### Others

4844 IMMUNOCYTO CD107a Detection Kit  
8223 IMMUNOCYTO IFN- $\gamma$  ELISPOT Kit  
AM-1005 IMMUNOCYTO Cytotoxicity Detection Kit  
TS-8002 T-Select MHC Tetramer Lyse  
TS-9004 T-Select Antibody Gating Kit  
TS-9017 T-Select MHC Tetramer T Cell Typing Kit  
6603861 CD8-FITC (T8)  
6607011 CD8-PC5 (T8)  
A07704 7-AAD Viability Dye  
IM-1400 OptiLyse B  
A11895 OptiLyse C  
MTG-001 Clear Back (Human FcR blocking reagent)

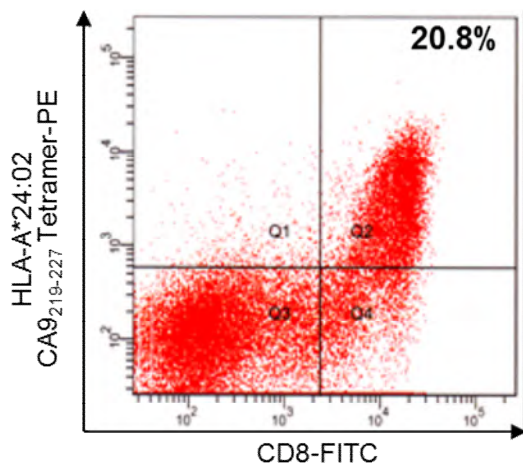
Please check our web site (<http://ruo.mbl.co.jp>) for up-to-date information on products and custom MHC Tetramers.

## Experimental Data

Data kindly provided by Dr. Kazuhiro Yoshikawa,  
Cell Therapy Center, Aichi Medical University  
Hospital

Venipuncture whole blood from healthy adult volunteers after receiving vaccines including epitope peptide was collected in sodium heparin blood collection tube. Peripheral blood mononuclear cells (PBMCs) were isolated from whole blood according to standard methods. Aliquots of PBMCs were incubated in a flask in the presence of a synthetic peptide (EYRALQLHL, 5 µg/mL), 100 IU/mL interleukin-2 (IL-2) and 5% (v/v) autologous plasma. After 72 hours, half of the medium was changed with medium containing 5 µg/mL of a synthetic peptide, 100 IU/mL IL-2 and 5% (v/v) autologous plasma, and every 3 days thereafter, half of the medium was changed. After 3 days, aliquots of these cells were stained with MHC Tetramers and CD8 antibody.

The number in the top right quadrants represents the percentage of tetramer-positive cells in the total cells.



250901-2