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Chandirasegara Massilamany
University of Nebraska-Lincoln, cmassilamany@unl.edu

Bharathi Krishnan
University of Nebraska-Lincoln, bkrishnan2@unl.edu

Jay Reddy
University of Nebraska-Lincoln, jayreddy@unl.edu

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Major histocompatibility complex class II dextramers: New tools for the detection of antigen-specific, CD4 T cells in basic and clinical research

Chandirasegaran Massilamany, Bharathi Krishnan, and Jay Reddy

School of Veterinary Medicine and Biomedical Sciences, University of Nebraska-Lincoln, Lincoln, NE 68583

Abstract

The advent of major histocompatibility complex (MHC) tetramer technology has been a major contribution to T cell immunology, because tetramer reagents permit detection of antigen-specific T cells at the single-cell level in heterogeneous populations by flow cytometry. However, unlike MHC class I tetramers, the utility of MHC class II tetramers has been less frequently reported. MHC class II tetramers can be used successfully to enumerate the frequencies of antigen-specific CD4 T cells in cells activated *in vitro*, but their use for *ex vivo* analyses continues to be a problem, due in part to their activation dependency for binding with T cells. To circumvent this problem, we recently reported the creation of a new generation of reagents called MHC class II dextramers, which were found to be superior to their counterparts. In this review, we discuss the utility of class II dextramers vis-a-vis tetramers, with respect to their specificity and sensitivity, including potential applications and limitations.

Introduction

Historically, studies related to the detection and functionalities of antigen-specific T cells at the single-cell level have been limited because the appropriate reagents and tools were not available. Commonly employed readouts included T cell proliferation assays based on incorporation of tritiated $^3\text{[H]}$ -thymidine or 5-bromo-2'-deoxyuridine (BrdU); Carboxyfluorescein succinimidyl ester (CFSE)-labelling; enzyme-linked immunospot (ELISPOT) assays; limiting-dilution analysis (LDA); and intracellular cytokine analysis [1–5]. Although most of these assays are helpful in ascertaining antigen-specific T cell responses in mixed cell cultures at the population level, accurate enumeration of the frequencies of antigen-specific T cells at the single-cell level has been a major limitation. Even assays like ELISPOT or cytokine-analysis can be prone to errors because it is difficult to eliminate the contribution of bystander T cells that can be non-antigen-specifically activated, leading to the possibility of overestimating the antigen-specific T cells [1]. Similarly, although LDA permits analysis of antigen-specific T cells at the single cell-level, this assay can never be routinely practical due to both the need to repeatedly activate the cells and the laborious nature of the assay [1, 6].

Correspondence address: Jay Reddy MVSc., PhD, Room 202, Bldg VBS, School of Veterinary Medicine and Biomedical Sciences, University of Nebraska-Lincoln, Lincoln, NE 68583, Phone: (402) 472 8541, nreddy2@unl.edu, Web: <http://jayreddy.unl.edu>.

These circumstances changed, however, with the publication of a 1996 landmark paper by Altman et al. describing the creation of major histocompatibility complex (MHC) class I tetramer technology; the tetramer reagents were found to be valuable in enumerating the frequencies of antigen-specific CD8 T cells by flow cytometry [7]. In 1998, Kappler's group provided a similar platform for CD4 T cells by generating peptide-tethered MHC class II tetramers [8]. These discoveries enabled researchers to determine the specificity of antigen-responsive T cells, particularly with respect to their appearance, disappearance and/or persistence in both basic and clinical research investigations (Table 1). MHC tetramers can be defined as artificially created soluble fluorochrome-conjugated MHC molecules assembled with peptides of interest. Their binding to antigen-specific T cells is captured by flow cytometry using the signals emitted by fluorochromes as readouts. However, some issues have continued to persist related to the inherent inability of MHC class II tetramers to bind CD4 T cells, especially low-affinity T cell receptor (TCR)-bearing autoreactive T cells, in spite of the fact that they are antigen-specific [9, 10]. To alleviate this problem, we created a newer version of tetramers called MHC class II dextramers for various autoantigens and successfully tested their utility in several experimental autoimmune and infectious disease models [11–15]. In this review, we discuss the utility of MHC class II dextramer reagents, most importantly their advantages over tetramers, as well as potential applications and limitations (Table 2). Nonetheless, for extensive details on the derivation and use of MHC class II tetramers, readers are encouraged to consult other excellent reviews published by various groups [9, 10, 16, 17].

What are MHC class II tetramers and dextramers and how are they created?

To understand the derivation of MHC class II tetramers and dextramers, it is useful to understand how MHC molecules display peptides for recognition by T cells. In contrast to MHC class I molecules that are composed of a single alpha chain supported by β 2-microglobulin as a scaffolding molecule, MHC class II molecules are made up of two chains, α and β . While the peptide-binding groove in the MHC class I molecule is formed by the participation of α 1 and α 2 domains, the α 1 and β 1 domains from their corresponding chains within the MHC class II molecules take part in the formation of grooves to which the peptides can anchor. The peptides thus displayed by MHC class I and MHC class II molecules are recognized by CD8 or CD4 T cells, respectively, and CD8 and CD4 molecules act as co-receptors by binding to the corresponding MHC molecules [18]. Nonetheless, it should be noted that the affinity of T cell recognition of MHC-displayed peptides is weak ($KD \sim 0.1\text{--}500 \mu\text{M}$), and engagement of multiple TCRs with as many multiple MHC-peptide complexes favors stability of their interactions, leading to enhanced avidity [19, 20].

The creation of tetrameric reagents requires expression of soluble MHC molecules that represent the extracellular portions in each class of MHC molecule [7, 8]. The presence of a single α chain may offer some advantage in allowing MHC class I molecules to be expressed more easily than MHC class II molecules in which both α and β chains must stay together so peptides can be displayed in the MHC clefts. Several strategies have been successfully employed for the expression of MHC class II molecules for various alleles in both mice and humans [8, 21–24]. Expression of soluble MHC class II molecules requires

the creation of two constructs, one each for α and β chains. A BirA site for biotinylation is introduced into one of the two chains, usually the α chain (Fig 1).

To express soluble molecules, various mammalian and non-mammalian expression systems have been used [9, 25–27]. However, it is to be noted that creation of MHC constructs depends on the intended use of the soluble molecules. Three major approaches are commonly used. (1) *Derivation of peptide-tethered soluble MHC molecules*. In this approach, the nucleotide sequence for the desired peptide is covalently tethered to the N-terminus of the β chain such that the peptide-containing soluble MHC molecules are expressed directly [8, 23]. Two advantages have been noted with this approach: (i) the peptide-bound MHC molecules are readily available for downstream applications, so the loss/instability of peptides is not an issue [8, 27]; and (ii) the covalently assembled peptides may provide stability for MHC monomers during protein expression [8, 9]. The major disadvantage is the need to generate constructs individually for each peptide of interest. (2) *Derivation of soluble, empty MHC molecules*. Here, α and β chains without peptide sequences are expressed to obtain empty molecules, and the desired peptides are then loaded exogenously to generate peptide-bound MHC complexes. Technically, this approach is versatile and desirable, but the stability of peptides within the groove can be an issue [28–30]. (3) *Derivation of soluble MHC molecules containing class II-associated invariant chain peptide (CLIP)*. This approach is similar to approach 1 as described above, but instead of tethering the specific peptide of interest to the N-terminus of the β chain, the CLIP-conjugated sequence with a site for thrombin cleavage is inserted covalently to the N-terminus of the β chain. As a result, upon protein expression, the CLIP is cleaved by thrombin to obtain empty MHC molecules, into which peptides of interest can be loaded exogenously by a peptide-exchange reaction similar to that employed in approach 2 [21, 22, 31].

In all of these approaches, however, because it is possible for α and β chains to separate during expression, it is a common practice to insert leucine-zipper dimerization motifs from the transcription factors Fos (3.98 KDa) and Jun (3.92 KDa) to the α and β chains, respectively, of MHC class II molecules [23, 32, 33]. By virtue of high-affinity binding between Fos and Jun proteins (KD 5.4×10^{-8} M; [34]), the α and β chains are expected to stay together as single monomers during protein expression, as well as in the downstream steps of protein purification, multimerization and staining by flow cytometry. We commonly employ approaches 1 and 3 in our routine tetramer and dextramer preparation.

The purified soluble MHC class II monomers are first biotinylated using biotin ligase (BirA) (Fig 1). While the peptide-tethered biotinylated proteins are directly multimerized using streptavidin (SA) conjugated with fluorochromes, such as phycoerythrin (PE) or allophycocyanin (APC), the CLIP-tethered proteins are first subjected to thrombin cleavage to release the CLIP, and the empty molecules are then assembled with peptides in the peptide-exchange reaction, followed by multimerization with SA-fluorochromes [21–23]. Although the latter approach is versatile in that we can generate reagents for multiple antigens by loading the peptides exogenously into empty MHC molecules, we noted a few limitations. First, peptides that are readily soluble in water or PBS can be reliably used to prepare the reagents, and addition of dimethyl sulfoxide to enhance the solubility of peptides

did not yield better results. Second, the sensitivity of tetramers and dextramers generated for some peptides was lower than that could be achieved with the reagents generated from the use of covalently tethered MHC molecules (unpublished observations).

To prepare multimeric MHC-peptide complexes using SA-fluorochrome conjugates, it is important to note that each SA can assemble up to four biotinylated MHC molecules. Hence, tetramers typically are expected to possess up to four peptide-bound biotinylated MHC molecules. In contrast to tetramers, the dextramers offer an advantage of assembling more peptide-bound MHC molecules – up to 20 per dextran molecule [15, 35]. Structurally, dextramers contain dextran backbones, which are polymers of glucose molecules attached through 1–6 and 1–3 linkages [35]. Each dextran molecule carries multiple moieties of SA to which biotinylated peptide-tethered MHC molecules can be assembled [35]. Therefore, dextramers can yield MHC-peptide complexes of large molecular weight, allowing them to engage multiple TCRs – more than could be achieved with tetramers.

To prepare the dextramers, we use dextran molecules (kind gift from Immudex, Denmark) at a working concentration of 3.2×10^{-8} M. Conjugation of dextran molecules with MHC-peptide molecules depends on the number of SA-fluorochrome-assembled moieties available in the dextran backbone. For example, a dextran-PE backbone containing 6.6 SA moieties has a molar ratio of 19.8 ($6.6 \times$ a valency of 3). Therefore, each mole of dextran-PE can be assembled with 19.8 moles of MHC class II monomers. We usually assemble 21 to 22 μg , equivalent to 3.17×10^{-10} moles of MHC class II monomers (~ 68 KDa) with 1.6×10^{-11} moles of dextran-PE [15]. On a reaction basis, we routinely use PE-dextramers containing 6.3×10^{-12} moles, equivalent to 0.45 μg of MHC protein ($\sim 3.8 \times 10^{12}$ molecules), to stain 1×10^6 cells [15]. Likewise, one mole of dextran-APC with a molar ratio of ~ 8.7 is used to assemble 8.7 moles of MHC class II monomers. To prepare APC-dextramers, we use ~ 9.5 to 10 μg of MHC class II protein, equivalent to 1.39×10^{-10} moles, to assemble 1.6×10^{-11} moles of dextran-APC; and to stain 1×10^6 cells, 2.78×10^{-12} moles (0.2 μg ; $\sim 1.67 \times 10^{12}$ MHC monomers) are used. Overall, the amount of MHC protein needed to prepare dextramer-APC is half the amount needed to prepare dextramer-PE, but the staining intensities obtained by the two reagents are comparable [15]. For routine use, the reaction conditions for each dextramer reagent are to be optimized. We found that staining at room temperature for a minimum of 30 minutes in growth medium (RPMI with 2.5% fetal bovine serum and interleukin-2, pH 7.63) was optimum for various MHC class II alleles assembled with a wide range of self- and foreign antigens [11, 14, 15].

Are MHC class II dextramers better reagents than MHC class II tetramers?

During the past 10 to 15 years, we have been using MHC class II tetramers in various autoimmune disease models in mice, primarily experimental autoimmune encephalomyelitis (EAE) induced with myelin proteolipid protein (PLP) 139–151 and myelin oligodendrocyte glycoprotein (MOG) 35–55; amoebic encephalitis; experimental autoimmune myocarditis (EAM) induced with cardiac myosin heavy chain (Myhc)- α 334–352; and experimental autoimmune uveoretinitis induced with interphotoreceptor retinoid-binding protein 201–216 [11, 15, 23, 33, 36–42]. Although the tetramer reagents were helpful, their use was largely limited to enumerating the frequencies of antigen-specific CD4 T cells in cultures activated

in vitro, and the direct *ex vivo* analysis of antigen-sensitized cells was a persistent problem. Enrichment procedures involving the use of antibodies to fluorochromes like PE to enrich the tetramer-PE-bound CD4 T cells by magnetic separation followed by extrapolating their frequencies to the total number of T cells have been commonly employed in the field [16, 22, 43, 44]. In these settings, the use of enzymes like protein tyrosine kinase inhibitors (PKIs) (e.g., Dasatinib, Bristol-Myers Squibb) prior to tetramer staining and enrichment procedures appears to enhance the sensitivity of tetramers [45, 46]. These methods can be labor intensive, and time consuming. In addition, technically, there also exists a possibility of non-antigen-specific cells to be present as contaminants in the enriched populations necessitating the use of multiple surface markers to eliminate them [16]. Likewise, peripheral blood mononuclear cells (PBMCs) from humans are usually stimulated with antigen-pulsed autologous antigen-presenting cells (APCs) for up to ~7 days prior to enrichment procedures [47]. Although these analyses may yield valuable information, absolute enumeration of the frequencies of antigen-specific T cells can be prone for errors leading to inaccurate estimations of their repertoire sizes. In some instances, frequencies of antigen-specific T cells can be detected only marginally higher than the background staining with the control tetramers because of difficulties in differentiating background staining from the specific tetramer staining [48]. To overcome these limitations with tetramers, we made efforts to create dextramers utilizing the dextran reagents conjugated with various fluorochromes (kind gift from Immudex, Denmark) for IA^s/PLP 139–151, IA^b/MOG 35–55, and IA^k/Myhc- α 334–352 [15]. We found that dextramers, but not tetramers, were helpful in enumerating the frequencies of antigen-specific CD4 T cells *ex vivo* in all the above models [15]. In addition, we noted that the detection sensitivity of dextramers could be increased by at least 4- to 5-fold compared with tetramers. Similarly, the specificity of staining obtained with dextramers was better than that could be achieved with tetramers, as background staining with the control dextramers was negligible [15]. One other group recently reported similar observations by generating the dextramers for human MHC class II allele, HLA-DR1 [49]. We think that dextramers consisting of multiple MHC-peptide complexes may more easily engage with multiple TCRs, leading to enhanced stability of dextramer-T cell interactions. As such, the dextramer-bound T cells are less likely to be detached during various steps in the staining reactions.

It is a common belief that tetramers bind poorly to low-affinity TCR-bearing T cells, because their binding depends on the activation status of T cells, and tetramers therefore are unlikely to bind the resting cells even if they are antigen-responsive [33, 50–52]. We and others have previously reported that the tetramer⁺ cells were found almost exclusively within the activated (CD4^{high}/CD25⁺) population [33, 47, 50, 51]. We had proposed that the activation status of cells may result in the reorientation or change in the configuration of the TCR and cell surface molecules leading to increase in the avidity of TCR-MHC binding [33]. To overcome this limitation, we had previously shown that antigen-responsive, rested cells can be made to bind with tetramers by first exposing the cells to neuraminidase (NASE), which essentially promotes T cell activation by an unknown mechanism [15, 23, 33]. In addition, removal of sialic acid residues from the surface of T cells by NASE-treatment may increase TCR affinity as shown with an insulin-insulin receptor system [53]. We verified that the dextramers can detect rested cells without the need for NASE-

treatment, thus overcoming the activation dependency problem noted with tetramers. Based on these successes, we expanded the utility of dextramers to detect antigen-specific CD4 T cells in target organs, such as brains in EAE mice and hearts in EAM mice, by flow cytometry [11, 15]. More importantly, we showed for the first time that MHC class II dextramers can be used to detect antigen-specific CD4 T cells *in situ* directly, both in the brain and heart in EAE and EAM models, respectively, by confocal microscopy without having to amplify the fluorochrome signals using secondary antibodies, which had been the case with MHC class I tetramers [12, 13, 54, 55]. This modality of *in situ* staining considerably reduced the duration of the process to less than a day (~12 hours), as compared to up to 3 days with MHC class I tetramers [54, 55]. Furthermore, the amount of reagent needed per *in situ* reaction is 2.5 µg/ml, which is significantly less than required with tetramers (20 µg/ml) [15, 56].

By extending the utility of dextramers to phenotype antigen-specific CD4 T cells in relation to cytokine-producing cells, we noted the unexpected finding that only a fraction of dextramer⁺ CD4 T cells are capable of producing Th cytokines, while the majority of cytokine-producing cells lie within the dextramer⁻ CD4 T cell populations (unpublished preliminary observations). Using IA^s/PLP 139–151 and IA^b/MOG 35–55 dextramers in the EAE system and IA^k/Myhc-α 334–352 dextramers in the EAM system, we are now phenotyping the cytokine-producing dextramer⁺ and cytokine-producing dextramer⁻ CD4 T cells corresponding to Th1, Th2 and Th17 subsets of cytokines to determine whether common cytokine signatures exist in different autoimmune diseases. More recently, we tested whether dextramers prepared by using naked (fluorochrome-unconjugated) dextran backbones (kind gift from Immudex) can be used as APCs to stimulate T cells in bioassays. An amount of 0.07 µg was found sufficient to induce robust proliferative responses in T cell hybridomas specific to PLP 139–151 and Myhc-α 334–352 (unpublished observations). These reagents may be useful in studying T cell-specific responses when the contribution of APCs must be avoided. One such potential application is the evaluation of reactive oxygen species (ROS) in T cells, whose detection within T cells has been debated, and the argument made that the T cells derive ROS as a contaminant from APCs.

Can MHC class II dextramers solve all problems with MHC class II tetramers? If not, what improvements can be made?

As described above, we have shown the utility of MHC class II dextramers in several experimental systems, finding the dextramers to be superior to their counterparts in terms of sensitivity, specificity and applications. If the dextramers can be so much more versatile than tetramers, a critical question is whether they can be used as reliable reagents to detect CD4 T cells with varied TCR affinity. Unfortunately, the answer is no. We performed an experiment to sort the dextramer⁺ and dextramer⁻ CD4 T cells from PLP 139–151-sensitized primary T cell cultures by flow cytometry, expecting that the dextramer⁻ cells would neither respond to antigen nor bind PLP dextramers upon restimulation. Contrary to our expectations, however, the dextramer⁻ cells did respond to PLP 139–151 and also stained with PLP dextramers [15]. We sorted CD4 T cells sensitized with PLP 139–151 that are negative for PLP 139–151 dextramers, and later stimulated them with PLP 139–151.

Approximately, 4% of cells from these cultures were found positive for PLP 139–151 dextramers, and expectedly, under similar conditions, the PLP 139–151 tetramer⁻ counterparts revealed more number of cells (7.2%) to be positive for PLP 139–151 dextramers [15]. These data suggest that a proportion of antigen-responsive T cells, possibly very low-affinity TCR-bearing cells, may not be detected by dextramers. Thus, the need for improving the detection sensitivity of dextramers still continues. Reports indicate that pre-exposure of cells to PKIs enhance the staining intensity of MHC class I tetramers/dextramers [49, 57]. Mechanistically, PKIs appear to inhibit downregulation of TCRs, including CD8 co-receptors, leading to their persistence on the surface of T cells [49, 57]. It is possible that similar strategies may work for MHC class II dextramers, although participation of CD4 co-receptor is not shown to be critical for binding with MHC class II tetramers [8, 58].

In summary, we have analyzed the importance of the utility of MHC class II dextramers, especially for detecting autoreactive CD4 T cells in relation to tetramers. While tetramers and other similar reagents (pentamers and protein-A conjugated multimers) are being made available through several academic (NIH Tetramer Core Facility, Atlanta, GA) and non-academic sources (Proimmune Inc, Beckman Coulter Inc, and MBL Inc, and TCMetrix Inc), dextramers are currently available only commercially (Immudex, Denmark). By creating dextramers for various autoantigens, we showed them to be superior to tetramers with respect to staining specificity, detection sensitivity, and various applications, but a fraction of antigen-specific CD4 T cells may still be undetected by dextramers [11–15, 59]. Nonetheless, specifically for *ex vivo* analysis, the dextramer reagents may offer an advantage by enumerating the frequencies of antigen-specific T cells directly, thereby avoiding the need to adopt other laborious procedures like enrichment of tetramer-bound T cells by magnetic separation [22, 43]. Finally, it should be noted that MHC class II tetramers are increasingly being used in clinical research in various areas, including, for example, studies of infectious diseases (Lyme arthritis, pulmonary tuberculosis, cytomegalovirus infection, and chronic hepatitis), immune-mediated diseases (type I diabetes and multiple sclerosis), and allergic diseases (atopic dermatitis and rye grass-associated allergy), and for monitoring T cell responses to vaccines (melanoma, influenza, and anthrax) [9, 17, 21, 47, 60–70]. In all of these evaluations, the frequencies of antigen-specific T cells have been reported to range from 0.006% to 3.5% in different body fluids, proving that tetramers are helpful in clinical research. But the real challenge is to be able to reliably detect the T cells *ex vivo* in a short period of time without having to stimulate the PBMCs, which is often the case with the human samples [21, 63]. Such limitations can possibly be overcome with the availability of MHC class II dextramers, since their detection sensitivity is higher than that of tetramers [15, 35]. To this end, we are now in the process of creating dextramers for use in patients with multiple sclerosis. We believe that potential exists for the use of MHC class II dextramers in both basic and clinical research as the reagents become available.

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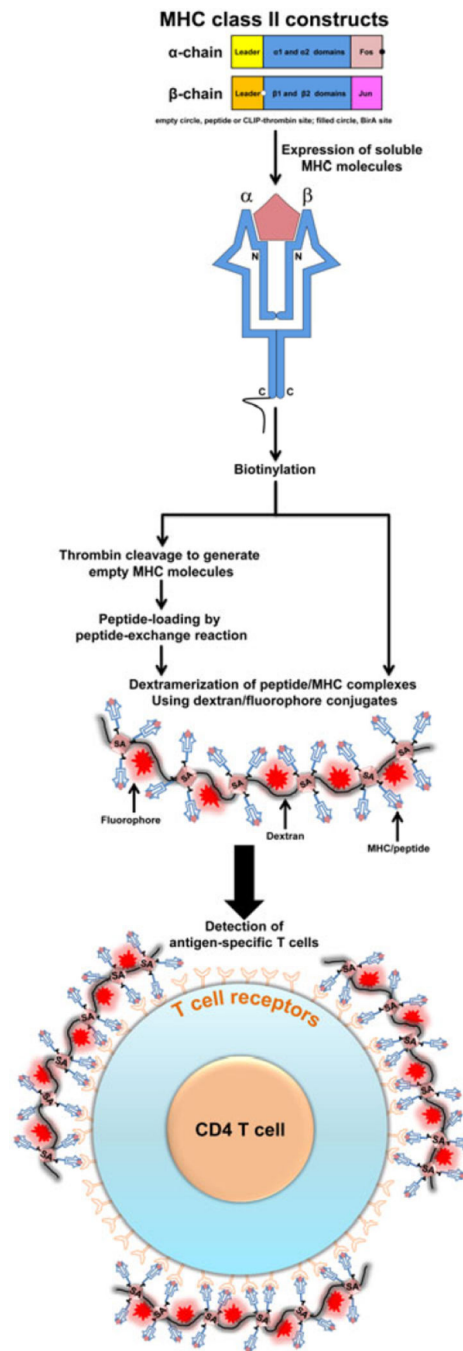


Figure 1. Derivation of MHC class II dextramers

To produce soluble MHC class II monomers, two constructs, each representing MHC class II α and MHC class II β chains, are first created. Generally, we insert the nucleotide sequence for specific peptides of interest or thrombin-cleavage site-tagged/CLIP into the N-terminal end of the β chain construct. Likewise, Fos-and Jun-sequences are attached to the C-terminal ends of α and β constructs, respectively, to facilitate stability of α and β chains during protein expression, whereas the BirA site for biotinylation is introduced in the α -construct. The constructs are used to express soluble MHC molecules using Baculovirus in

Sf9 insect cells. After biotinylation, the MHC monomers obtained from the constructs containing covalently tethered, specific peptide sequences are assembled with dextran molecules to obtain dextramers directly. In contrast, the biotinylated proteins generated from the CLIP-tethered constructs are first treated with thrombin to release the CLIP peptide, and the resulting empty MHC monomers are then used to load peptides of interest exogenously by peptide-exchange reaction. For dextramerization, the biotinylated MHC monomers generated from any of the above two approaches are mixed with dextran-SA-fluorochrome at molar ratios ranging from ~9:1 or 20:1, and the dextramers are then used to stain T cells to analyze their frequencies by flow cytometry.

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Table 1

List of MHC class II tetramers and dextramers, and their use for the determination of antigen-specific T cell responses in mice and humans

MHC class II alleles/epitopes	Application
(a) Tetramers	
<u>Mice</u>	
<u>IA^b</u>	
IRBP ₂₇₁₋₂₉₀ and IRBP ₇₇₁₋₇₉₀	Thymic selection of T cells [71]
LCMV GP ₆₆₋₇₇ , GP ₁₂₆₋₁₄₀ , GP ₆₋₂₀ and CLIP	Detection of LCMV-specific T cells in infected mice [72]
MOG ₃₅₋₅₅ , MOG ₃₈₋₄₉ and hCLIP ₁₀₃₋₁₁₇	Detection of MOG-specific T cells in EAE mice [40, 73]
OVA ₃₂₃₋₃₃₉ , CLIP and LCMV GP ₆₁	Detection of OVA-specific T cells in immunized mice [74]
FliC ₄₂₇₋₄₄₁ , 2W1S, OVA ₃₂₃₋₃₃₉	Determination of the frequencies of antigen-specific T cells in naïve repertoires [43]
HY	Detection of antigen-specific T cells in naïve and immunized mice [75]
<u>IA^d</u>	
OVA ₃₂₃₋₃₃₉ and influenza HA ₁₂₆₋₁₃₈	Peptide/MHC display and antigen-specificity studies [8, 76, 77]
<u>IA^{g7}</u>	
p8G p6- α 62, p8G p11- α 72, p8E p6- α 62, p8E p11- α 72, insulin, CHGA and HEL	Detection of insulin-specific T cells in NOD mice [78]
GAD ₅₂₄₋₅₄₃ , GAD ₁₀₄₀₋₇₉ , GAD ₁₁₃₆₋₁₇ , p206, p524, GAD ₂₀₆₋₂₂₀ and GAD ₄₇₁₋₄₉₀	Detection of GAD-specific T cells in NOD mice [22, 47, 79]
<u>IA^k</u>	
Myhc- α ₃₃₄₋₃₅₂ and RNase ₄₃₋₅₆	Detection of Myhc- α -specific T cells in EAM mice [31]
IRBP ₂₀₁₋₂₁₆ , and EHC ₄₄₋₅₉	Detection of IRBP, and IRBP- and EHC-cross-reactive T cells in EAU mice [36]
<u>IA^s</u>	
PLP ₁₃₉₋₁₅₁ and TMEV ₇₀₋₈₆	Detection of PLP-specific T cells in EAE mice [33]
PLP ₁₃₉₋₁₅₁ and ACA ₈₃₋₉₅	Detection of PLP- and ACA-cross-reactive T cells in EAE mice [14]
<u>IE^k</u>	
MCC ₈₈₋₁₀₃	First report to demonstrate that MHC class II tetramers detect antigen-specific CD4 T cells [78]
U1-70 ₁₃₁₋₁₅₀ , U1-70 ₁₃₁₋₁₅₀ , P(140) and MCC ₈₈₋₁₀₃	TCR-signaling studies, and antigen-specific T cell responses [58, 80, 81]
<u>Humans</u>	
<u>DP4</u>	
MAGE-3	Detection of MAGE-3-specific T cells in vaccinated patients [70]
<u>DQ2</u>	
α I- and α II-peptides of gliadin	Detection of gliadin-specific T cells in patients with celiac disease [82]

MHC class II alleles/epitopes	Application
<u>DR*1101</u>	
MAGE-A3 _{191–205} , MAGE-A3 _{281–295} and MAGE-A3 _{286–300}	Epitope-binding studies [83]
<u>DR0301</u>	
IGRP peptides and Flu MP _{169–181}	Tetramer-guided epitope mapping [84]
<u>DR0401</u>	
IGRP peptides and Flu HA _{306–318}	Tetramer-guided epitope mapping [84]
<u>DRB1</u>	
Bet v 1 _{141–155} , Der p 1 _{16–30} , Der p 1 _{171–185} and Phl p 1 _{194–208}	T cell responses in allergy [85]
<u>DRB1*0101</u>	
HIV Gag p24 _{294–313} , Gag p24 _{264–283} , TT _{830–843} , influenza HA _{307–319} and CMV pp65 _{108–127}	Detection of HIV-, tetanus toxoid-, influenza-, or CMV-specific T cells by magnetic bead enrichment [86]
HA _{307–319} and HA _{306–318}	Detection of influenza-specific T cells [87, 88]
Fel d 1	Detection of Fel d 1-specific T cells in atopic dermatitis patients [64]
<u>DRB1*0401</u>	
HA _{306–318} and GAD	Detection of influenza-specific T cells in vaccine-recipients [88]
MOG _{97–109}	Detection of MOG-specific T cells [89]
OspA _{164–175}	Detection of OspA-specific T cells in Lyme arthritis patients [62]
Human CII, HCgp39 and Osp-A	Detection of cartilage antigen-specific T cells in rheumatoid arthritis patients [90]
HCV 1248, HCV 1579 and HCV 1770	Detection of HCV-specific T cells in viral hepatitis [21, 63]
MART 151–173	Detection of MART-1-specific T cells in vaccine-recipients [69]
Lol P1	Detection of allergen-specific T cells in allergic individuals [65]
<u>DRB1*0402</u>	
Dsg3	Detection of Dsg3-specific T cells in pemphigus vulgaris [91]
<u>DRB1*0404</u>	
PA _{112–127}	Detection of antigen-specific T cells in recipients of Anthrax vaccine [67]
HIV Gag p24 _{164–183} and HCV NS3 _{1248–1261}	Detection of HIV- or HCV-specific T cells by magnetic bead enrichment [86]
<u>DRB1*1302</u>	
PA _{373–393} and PA _{381–392}	Detection of antigen-specific T cells in recipients of Anthrax vaccine [67]
<u>DR3</u>	
MTB 1–13 and CMV pp65 _{510–522}	Detection of MTB- and CMV-specific T cells in patients/vaccine-recipients [92]
<u>DR4</u>	

MHC class II alleles/epitopes	Application
HA _{p306-318} and NY-ESO-1 _{p119-130}	Detection of influenza- or NY-ESO-1-specific T cells by immunoscope [25]
Ag85B and ESAT-6	Detection of MTB-specific T cells in patients with pulmonary tuberculosis [61]
<u>DR401</u>	
GAD65 ₅₅₅₋₅₆₇ and HSV-p61	Detection of GAD-specific T cells in type 1 diabetes [47]
<u>DR404</u>	
GAD65 ₅₅₅₋₅₆₇ and HSV-p61	Detection of GAD-specific T cells in type 1 diabetes [47]
<hr/>	
(b) Dextramers	
<hr/>	
<u>Mice</u>	
<u>IA^b</u>	
MOG ₃₅₋₅₅	Detection of MOG-specific T cells in EAE mice [59]
<u>IA^k</u>	
Myhc- α ₃₃₄₋₃₅₂ and RNase ₄₃₋₅₆	Detection of Myhc- α -specific T cells in EAM mice and CVB3-infected mice [11, 15]
<u>IA^s</u>	
PLP ₁₃₉₋₁₅₁ and TMEV ₇₀₋₈₆	Detection of PLP-specific T cells <i>ex vivo</i> , <i>in vitro</i> and <i>in situ</i> in EAE mice [12, 13, 15]
PLP ₁₃₉₋₁₅₁ and ACA ₈₃₋₉₅	Detection of PLP- and ACA-cross-reactive T cells in ACA infected mice [14]
<u>Humans</u>	
<u>HLA-DR1</u>	
Influenza A HA ₃₀₇₋₃₁₉ and HIV-1 p24 Gag ₂₉₉₋₃₁₂	Detection of influenza-specific T cells [49]
<u>HLA-DP2</u>	
Mim2, Mim2 and Be, and PLXNA4 and Be	Detection of Be-specific T cells [93, 94]

* This list is intended to capture broad applications of tetramer and dextramer reagents in basic and clinical research, and by no means is a comprehensive list. As such, some reagents may not have been listed, and the authors wish to acknowledge this limitation.

IRBP, interphotoreceptor retinoid-binding protein; LCMV, Lymphocytic choriomeningitis virus; GP, glycoprotein; CLIP, Class II-associated invariant chain peptide; MOG, myelin oligodendrocyte glycoprotein; EAE, experimental autoimmune encephalomyelitis; OVA, ovalbumin; HA, hemagglutinin; FLiC, flagellar filament structural protein; HY, Y chromosome-encoded transplantation antigens; NOD, non-obese diabetic; GAD, glutamic acid decarboxylase; CHGA, chromogranin A; HEL, hen egg lysozyme; Myhc- α , cardiac myosin heavy chain- α ; RNase, Ribonuclease; EAM, experimental autoimmune myocarditis; EHC, *Ehrlichia canis*; EAU, experimental autoimmune uveitis; PLP, myelin proteolipid protein; TMEV, Theiler's murine encephalomyelitis virus; ACA, *Acanthamoeba castellanii*; MCC, moth cytochrome C; MAGE, melanoma-associated antigen; IRGP, islet-specific glucose-6-phosphatase catalytic subunit-related protein; MP, Matrix protein; Bet v 1, *Betula verrucosa*; Der p 1, *Dermatophagoides pteronyssinus*; Phl p 1, pollen allergen; HIV, Human immunodeficiency virus; Gag, group-specific antigen; TT, Tetanus toxoid; CMV, Cytomegalovirus; OspA, *Borrelia burgdorferi* outer surface protein-A; CII, type 2 collagen; HCV, Hepatitis C virus; MART, from Melan-Ae/melanoma antigen recognized by T cells; Lol P1, rye grass allergen; HCgp, Human cartilage glycoprotein; Dsg3, desmoglein3; PA, protective antigen of *Bacillus anthracis*; NS, nonstructural protein; MTB, *Mycobacterium tuberculosis*; NY-ESO-1, New York esophageal squamous cell carcinoma; Ag85B, Antigen 85; ESAT-6, Early Secretory Antigenic Target; CVB3, coxsackie virus B3; Mim2, mimotope-2; Be, beryllium

Table 2

Advantages of using MHC class II dextramers over tetramers for detecting antigen-specific, autoreactive CD4 T cells.

Parameters	Tetramers	Dextramers
a. Generation of reagents		
• Creation of MHC constructs, expression of soluble MHC monomers and biotinylation	Steps are similar for both the reagents	
• Multimerization	Require fluorophore/streptavidin-conjugated molecules	Require fluorophore/streptavidin-conjugated dextran molecules
b. Staining characteristics based on flow cytometry		
• Number of MHC monomers per reaction	12.6×10^{-12} moles	2.78×10^{-12} moles
• Specificity	Yes	Yes, but enhanced
• Sensitivity	Low	High
• Duration	Variable (usually 2 to 3 hours in our hands)	Relatively short (0.5 to 2 hours)
c. Applications		
• Detection of T cells <i>ex vivo</i>	May or may not work	Reliable and consistent
• Detection of T cells activated <i>in vitro</i>	Useful	Useful
• Detection of T cells <i>in situ</i>	May work	Reliable and consistent
• Cytokine-phenotyping by flow cytometry	Yes	Yes, but with enhanced sensitivity
• Bioassay	Yes	Yes