T Cells Specific for a Mycobacterial Glycolipid Expand after Intravenous Bacillus Calmette–Guérin Vaccination

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Intradermal vaccination with Mycobacterium bovis bacillus Calmette–Guérin (BCG) protects infants from disseminated tuberculosis, and i.v. BCG protects nonhuman primates (NHP) against pulmonary and extrapulmonary tuberculosis. In humans and NHP, protection is thought to be mediated by T cells, which typically recognize bacterial peptide Ags bound to MHC proteins. However, during vertebrate evolution, T cells acquired the capacity to recognize lipid Ags bound to CD1a, CD1b, and CD1c proteins expressed on APCs. It is unknown whether BCG induces T cell immunity to mycobacterial lipids and whether CD1-restricted T cells are resident in the lung. In this study, we developed and validated Macaca mulatta (Mamu) CD1b and CD1c tetramers to probe ex vivo phenotypes and functions of T cells specific for glucose monomycocoyl (GMM), an immunodominant mycobacterial lipid Ag. We discovered that CD1b and CD1c present GMM to T cells in both humans and NHP. We show that GMM-specific T cells are expanded in rhesus macaque blood 4 wk after i.v. BCG, which has been shown to protect NHP with near-sterilizing efficacy upon M. tuberculosis challenge. After vaccination, these T cells are detected at high frequency within bronchoalveolar fluid and express CD69 and CD103, markers associated with resident memory T cells. Thus, our data expand the repertoire of T cells known to be induced by whole cell mycobacterial vaccines, such as BCG, and show that lipid Ag-specific T cells are resident in the lungs, where they may contribute to protective immunity.

M. tuberculosis bacillus Calmette–Guérin (BCG) is the oldest vaccine still in clinical use today, with over 3 billion doses administered since 1921 (1). In most countries, infants undergo intradermal vaccination with BCG at birth, which protects them from disseminated tuberculosis and death in early childhood (2). BCG provides variable efficacy against tuberculosis disease when provided later in life (3). The reasons for this discrepancy are not fully understood but may relate to qualitative differences in the adaptive immune response to BCG (1).

Studies in mice and nonhuman primates (NHP) show that T cells are critical for the protective efficacy of BCG. T cell–deficient mice fail to control M. tuberculosis to the same extent as wild-type mice after BCG (4). CD8 T cell depletion in NHP abrogates the protective effect of BCG (5). More recently, i.v. BCG was shown to provide superior protection to M. tuberculosis challenge in rhesus macaques when compared with the standard intradermal route (6, 7). Higher levels of Ag-specific T cells in the lung and bronchoalveolar lavage (BAL) fluid were induced by i.v. BCG, suggesting that they may be directly responsible for improved protection (6).

Canonicly, T cells are activated by foreign peptide Ags presented by autologous MHC molecules (8). However, during vertebrate evolution, T cells have also acquired the capacity to recognize bacterial lipid Ags via the CD1 lipid Ag presentation pathway (9). Humans express four CD1 Ag-presenting molecules (CD1a, CD1b, CD1c, and CD1d), which vary in the configuration of their binding grooves, patterns of cellular expression, and subcellular trafficking (10). The first structurally defined lipid Ag was mycolic acid, which is composed of long (C90) alkyl chains, is a ubiquitous component of the mycobacterial cell wall, and is presented by CD1b to T cells (11). Follow-up studies identified the structurally related Ags glucose monomycocoyl (GMM) and glycerol monomycocoyl (12, 13). Several other mycobacterial lipid Ags have been discovered as T cell Ags, whose recognition is mediated by CD1a, CD1b, or CD1c (group 1) in humans (10). Current evidence suggests that these lipid-specific T cells are expanded in clinical settings of mycobacterial exposure or infection.
Lipid-loaded CD1 tetramers have facilitated further ex vivo studies of the phenotypes of CD1-restricted T cells in humans (17–21). T cell recognition of GMM is exquisitely sensitive to the position of the glucose headgroup but relatively independent of lipid chain length, allowing both short-chain (C_{32}) and C_{30}-GMM to be used in tetramers (12, 17, 22). Other studies have revealed that germline-encoded residues within TRAV-2 are required for recognition of GMM and that TRBV4-1 may bias T cells toward recognition of CD1b and CD1c (17, 23–25). Because CD1 genes exhibit very few polymorphisms in humans, CD1 tetramers can be used without prior knowledge of genotype, enabling population-based studies (15, 26).

Further studies elucidating the contribution of CD1a-, CD1b-, and CD1c-restricted T cells to protective immunity during M. tuberculosis infection have been hampered by the lack of a suitable animal model. Mice have only one ortholog of human CD1d, so this model has provided a narrow window into the role of CD1d-restricted invariant NKT (iNKT) cells in tuberculosis (27, 28). Genomic studies suggest that CD1a, CD1b, and CD1c genes were deleted from Rodentia or a common ancestor (29). Humanized CD1 transgenic (hCD1Tg) mice express CD1a, CD1b, and CD1c in a pattern similar to that seen in humans and support the development of group 1 CD1-restricted T cells (30). Infection of hCD1Tg mice with BCG induces mycolic acid-specific T cells, which home to pulmonary granulomas and mediate partial protection after M. tuberculosis challenge in a CD1-restricted manner (31). CD1b was shown to be expressed in human granulomas as well as conserved TCR sequences expressed by CD1b-restricted T cells (32, 33). Whether BCG induces T cell responses to mycobacterial lipids in humans and NHP is currently not known.

Evolutionary genetic analysis suggests that mice are the exception among mammals, most of which have expanded CD1 gene families similar to humans (34). Specifically, a five-gene CD1 locus is highly conserved between humans and NHP, which stands in stark contrast to genes encoded within the MHC (35). Several studies have investigated the phenotypes and functions of iNKT cells in NHP (35–37). However, there have been only a few attempts to explore CD1-restricted T cells. One effort using a human lymphoblastoid cell line (T2) stably transfected with a GMM-specific human TCR in the presence of GMM (38). Another study explored the restriction of T cells line transfected with a GMM-specific human TCR in the presence of GMM (38). Yet another study attempted to explore CD1-restricted T cells. One effort using a human lymphoblastoid cell line (T2) stably transfected with a GMM-specific human TCR in the presence of GMM (38). These limited and conflicting data suggest that T cells specific for mycobacterial glycolipids may be induced by vaccination, but their restricting element, phenotypes, and tissue distribution have not been determined.

We generated Mamu CD1b and Mamu CD1c tetramers and validated their ability to detect GMM-specific T cells by sorting them from the blood of BCG-vaccinated animals and performing in vitro expansion and functional assays. We find that the response to GMM is restricted by CD1b and CD1c in NHP and, surprisingly, in humans as well. We incorporated CD1c-GMM tetramers into a multiparameter flow cytometry panel and show that GMM-specific T cells are expanded in the blood of NHP 4 wk after i.v. BCG with a predominant effector memory T (TEM) phenotype. These cells are also detectable in the lung as tissue-resident memory T (T_{EMR}) cells 4 wk after vaccination. These data establish the NHP as a natural model to study CD1-restricted T cells, which are resident in the lung and may contribute to protective immunity induced by whole cell mycobacterial vaccines.

Materials and Methods

Evolutionary genetic analysis of group 1 CD1 genes

CD1a, CD1b, and CD1c sequences were collected for 11 primate species with sequenced genomes (human, chimpanzee, bonobo, gorilla, gibbon, orangutan, rhesus macaque, crab-eating macaque, green monkey, olive baboon, and marmoset) using a custom bioinformatics pipeline adapted from Young and Trask (40). Briefly, the Basic Local Alignment Search Tool (BLAST) was used with human CD1a, CD1b, and CD1c protein sequences as queries to identify potentially homologous regions in each primate genome. Putative exons were defined based on a low e-value (<1 x 10^-8), contiguity in sequence orientation, and proximity within 20 kb. These putative loci were then analyzed using exonere to predict protein-to-genome mode with human group 1 CD1 sequences as queries to identify putative open reading frames. For each paralog, sequences were aligned using Clustal Ω (41), and tests of positive selection were performed using the codeml application in PAML (v4.8) with the negative selection tree (43), and if significant, sites with Bayes-empirical Bayes-posterior probabilities ≥0.8 were considered under positive selection. Positively selected sites were modeled onto available crystal structures (Protein Data Bank numbers 5J1A, 5L2J, and 30V6 for CD1a, -b, and -c, respectively) using PyMOL v2.

Preparation of lipid-loaded CD1 tetramers

C_{32}GMM isolated from Rhodococcus equi, synthetic phosphomycoloykette (PM), and diaoylated sulfoylglycolipid (AcSGL) were generously provided by the laboratory of D. B. Moody (20, 44, 45). Synthetic C_{30}GMM was prepared as we recently described (45). Stock GMM, PM, or AcSGL was solubilized in chloroform/methanol (2:1, v/v) at a single concentration of 1 ng/ml. Manu and Homo sapiens CD1b and CD1c biotinylated monomers were provided by the National Institutes of Health (NIH) Tetramer Core Facility (Emory University, Atlanta, GA). CDMalo-loaded CD1b tetramers were generated as previously described with the following modifications: C_{32}GMM was used as a 40-fold excess, and lipid loading was performed overnight at 37°C (15). GMMLoaded CD1c tetramers were generated as follows: a 100-fold excess of C_{32}GMM was aliquoted into a single well, glass vial and evaporated under a nitrogen stream to remove the solvent. One hundred microliters of PBS was added, and the vial was sonicated at 37°C for 15 min and then moved to the 37°C water bath. Following the addition of CD1c monomer, the solution was incubated at 37°C overnight. On the following day, streptavidin (Life Technologies, Carlsbad, CA; BioLegend, San Diego, CA; BD Biosciences, San Jose, CA) was added to tetramerize the CD1c monomers. The tetramer was filtered through a Spin X column (Sigma-Aldrich, St. Louis, MO) to remove large aggregates and stored at 4°C until use. CD1b and CD1c mock-loaded tetramers were generated in an analogous manner but without the addition of lipid Ag. M1R-5-(2-oxopyrrolidinemono)-6-ribitylaminoetracil (5-OP-RU) and CD1d-α-galactosylceramide (α-GalCer) tetramers were provided by the NIH Tetramer Core Facility.

Generation of monocyte-derived dendritic cells

PBMCs were isolated from human (Bloodworks, Seattle, WA) or NHP whole blood (Washington National Primate Research Center, Seattle, WA) by density centrifugation with Ficoll-Paque PLUS media (Cytiva, Marlborough, MA) and plated in tissue culture dishes at a density of 10 million cells/ml in R10 media (RPMI 1640 [Life Technologies] with 10% FBS [HyClone Laboratories, Logan, UT]), supplemented with recombinant human GM-CSF (final concentration of 100 ng/ml) (PeproTech, Cranbury, NJ) and incubated at 37°C for 3 h. After incubation, residual nonadherent cells were removed by washing gently with R10. R10 supplemented with rhGM-CSF (final 100 ng/ml) and rhIL-4 (final 20 ng/ml) (PeproTech) was added to the adherent fraction, and the cells were then incubated at 37°C for 3 d (46). Following the incubation period, adherent monocyte-derived dendritic cells were harvested and cryopreserved in liquid nitrogen. Cells were screened by flow cytometry for CD1 expression by staining with the following anti-human CD1 Abs: anti-CD1a PE (clone HE149), anti-CD1b FITC (clone SN13), anti-CD1c allophecocyanin (clone L161) (Bio-Legend), and anti-CD1d PE (clone 42.1) (BD Biosciences).

Generation of human and NHP T cell lines

Cryopreserved human PBMC were thawed and rested overnight and then mixed with γ-irradiated (5000 rad) allogeneic monocyte-derived dendritic
cells at a ratio of 5:1 and incubated in T cell media (R10 supplemented with 100 U/ml penicillin, 100 mg/ml streptomycin, 55 mM 2-ML-0.3% essential amino acids, 60 mM nonessential amino acids, 11 mM HEPES, and 800 mM L-glutamine [Life Technologies]) with 1 μg/ml of C3d-GMM and 50 U/ml recombinant human IL-2 (rhIL-2; Prometheus Laboratories through University of Washington Medical Center clinical pharmacy) for 12–14 d. T cell lines were screened with H. sapiens CD1b-GMM or H. sapiens CD1c-GMM tetramer. To further purify GMM-specific T cells present within T cell lines, cells were stained with PE-conjugated tetramers in FACS buffer without additional additives. An anti-streptavidin PE Ab (clone 3A20.2) (BioLegend) was also included to enhance the fluorescence intensity of tetramer-specific cells (47). After Ab staining, the cells were resuspended in culture media and kept on ice until sorting. Tetramer-positive T cells were sorted at the University of Washington’s Department of Immunology Flow Cytometry Core using a FACSaria II (BD Biosciences) cell sorter equipped with violet (407-nm), blue (488-nm), and red (641-nm) lasers.

Sorted T cells were expanded using a modified version of a previously published rapid expansion protocol (48). Briefly, T cells were mixed with 5 million γ-irradiated (8000 rad) lymphoblastoid cells (courtesy of D. Koelle, University of Washington) and 25 million γ-irradiated (3300 rad) PBMC provided by an anonymous blood donor to act as feeder cells in T25 tissue culture flasks (Costar, St. Louis, MO). Anti-CD3 (clone OKT3) was added at a final concentration of 30 ng/ml, and the mixture was incubated overnight at 37°C. On the following day, rhIL-2 was added to a final concentration of 50 U/ml. On day 4, the cells were washed twice in T cell media to remove the anti-CD3 Ab and resuspended in fresh media supplemented with rhIL-2 at 50 U/ml. Half the media was replaced every 3 d or split into new T25 flasks as determined by cell proliferation. After 13 d in culture, T cell lines were screened using H. sapiens CD1b-GMM or H. sapiens CD1c-GMM tetramer. T cell lines were cryopreserved until use.

To generate NHP T cell lines, cryopreserved PBMC were thawed and rested overnight and then mixed with γ-irradiated Mamu allologeneic monocyte-derived dendritic cells and C3d-GMM as described above. After 2 wk in culture, T cells were stained using Mamu CD1b-GMM or Mamu CD1c-GMM PE-Cy5 tetramer and tetramer-positive T cells were sorted. Sorted T cells were expanded in vitro using the rapid expansion method detailed above, except that irradiated Mamu PBMC and splenocytes were used as feeder cells. Repeated rounds of expansion were achieved by using γ-irradiated human PBMC and lymphoblastoid cells in the presence of rhIL-2 and/or PHA (Remel, San Diego, CA).

Flow cytometry
PBMC or T cell lines were thawed in RPMI 1640 (Life Technologies) supplemented with 10% FBS (Atlas Biologicals, Fort Collins, CO), and viable cells were enumerated by trypan blue (MilliporeSigma, Burlington, MA) exclusion. The PBMC or T cell lines were plated at a final density of 1 million cells per well and then washed twice with 1× PBS (Life Technologies). The cells were then stained with Aqua LIVE/DEAD (Life Technologies), the cells were centrifuged at 250 × g for 10 min and rested overnight at a density of 2 million cells/ml. On the following day, the cells were counted and plated at a density of 4 million cells per well in a 96-well U-bottom plate. The staining procedure was performed as above with the following reagents: H. sapiens CD1d–εGalCer PE Tetramer, Mamu MR1–5-OP-RU BV421 Tetramer (NIH Tetramer Core Facility, Atlanta, GA), Mamu CD1c-GMM BV650 tetramer, Mamu CD1c-GMM PE-Cy7 tetramer, Mamu CD1c-mock PerCP Cy5.5 tetramer (custom), anti-CD3 BV737 (clone SP3-42,–CD4 allophycocyanin (clone T208), and anti-CD8 BV785 (clone 3C10) (Mabtech) and incubating overnight at 4°C. After Ab staining, the cells were resuspended in culture media and kept on ice until sorting. Tetramer-positive T cells were sorted at the University of Washington’s Department of Immunology Flow Cytometry Core using a FACSaria II (BD Biosciences) cell sorter equipped with violet (407-nm), blue (488-nm), and red (641-nm) lasers.

For results described in Figs. 4 and 5, the rhesus macaque PBMC and BAL samples were thawed in warm media consisting of R10 and 2 μM Benzoxane (MilliporeSigma) sterile-filtered and centrifuged at 250 × g for 10 min. The supernatant was decanted, and the viable cells were enumerated using Guava Viacount Reagent (MilliporeSigma) and acquired on a Guava easyCyte Benchtop Flow Cytometer (MilliporeSigma) according to manufacturer’s instructions. The cells were centrifuged at 250 × g for 10 min and rested overnight at a density of 2 million cells/ml. On the following day, the cells were counted and plated at a density of 4 million cells per well in a 96-well U-bottom plate. The staining procedure was performed as above with the following reagents: H. sapiens CD1d–εGalCer PE Tetramer, Mamu MR1–5-OP-RU BV421 Tetramer (NIH Tetramer Core Facility, Atlanta, GA), Mamu CD1c-GMM BV650 tetramer, Mamu CD1c-GMM PE-Cy7 tetramer, Mamu CD1c-mock PerCP Cy5.5 tetramer (custom), anti-CD3 BV737 (clone SP3-42,–CD4 allophycocyanin (clone T208), and anti-CD8 BV785 (clone 3C10) (Mabtech) and incubating overnight at 4°C. After Ab staining, the cells were resuspended in culture media and kept on ice until sorting. Tetramer-positive T cells were sorted at the University of Washington’s Department of Immunology Flow Cytometry Core using a FACSaria II (BD Biosciences) cell sorter equipped with violet (407-nm), blue (488-nm), and red (641-nm) lasers.

Flow cytometry
PBMC or T cell lines were thawed in RPMI 1640 (Life Technologies) supplemented with 10% FBS (Atlas Biologicals, Fort Collins, CO), and viable cells were enumerated by trypan blue (MilliporeSigma, Burlington, MA) exclusion. The PBMC or T cell lines were plated at a final density of 1 million cells per well and then washed twice with 1× PBS (Life Technologies). The cells were then stained with Aqua LIVE/DEAD (Life Technologies) according to manufacturer’s instructions and incubated for 15 min at room temperature. LIVE/DEAD staining and all following steps were performed in the dark at room temperature. At the end of the incubation, the cells were washed twice in PBS and blocked by incubating the cells with human serum (Valley Biomedical, Winchester, VA) and FACS buffer (PBS supplemented with 0.2% BSA [Sigma-Aldrich]) mixed 1:1 for 15 min at 4°C. Anti-CCR7 Ab (BD Biosciences) was then prepared with 50 nM dasatinib (MilliporeSigma) diluted 1:1000 in PBS and incubated overnight at 4°C. After Ab staining, the cells were washed twice in PBS and incubated with 4 μg/ml of Anti-CCR7 Ab (clone 7-B6-1) (Mabtech) diluted to 1 μg/ml; PBS/0.5% FBS and incubated for 2 h at room temperature. The cells were then washed five times with PBS. One hundred microliters of ExtrAvidin–Alkaline Phosphatase (Sigma-Aldrich) diluted 1:1000 in PBS was then added, and the plates were incubated for 1 h at room temperature. The samples were washed five times and incubated with BCIP/NBT substrate (Sigma-Aldrich) prepared according to manufacturer’s instructions for 15 min in the dark at room temperature to develop the membrane. IFN-γ spots were measured using an ImmunoSpot S6 Core Analyzer (Cellular Technology, Cleveland, OH).

Detection of IFN-γ produced by Mamu T cells was performed similarly but with the following modifications: a MultiScreen PVDF Plate (MilliporeSigma) was wetted using 15 μl of 35% ethanol for no longer than 1 min. The plate was then washed with sterile water five times before coating the plate with 100 μl of PBS containing 15 μg/ml of anti-human IFN-γ Ab (clone MT126L) (Mabtech) and incubating overnight at 4°C.

Animals
Derivation of macaque T cell lines began with PBMC isolation from a female Chinese rhesus macaque 12 wk after intradermal vaccination with BCG. Data presented in Figs. 4 and 5 derived from 10 Indian-origin rhesus macaques who underwent i.v. vaccination with 3×10^7 CFU/ml of Danish strain 1331 BCG as part of a larger study that was recently published (6). Cryopreserved PBMCs derived from blood collected at baseline or 4 wk postvaccination as well as BAL collected 4 wk postvaccination were used for the experiments described in this study. This study was established and approved by the Animal Welfare Act and the Guide for the Care and Use of Laboratory Animals. Macaques were monitored for physical health, food consumption, body weight, temperature, complete blood counts, and serum chemistries.
Clinical cohort

Derivation of human T cell lines began with cryopreserved PBMC donated by a South African adolescent who was enrolled into a study that aimed to determine the incidence and prevalence of tuberculosis infection and disease (51). Twelve- to eighteen-year-old adolescents were enrolled at high schools in the Worcester region of the Western Cape of South Africa. Subjects were screened for the presence of latent tuberculosis by a tuberculin skin test and IFN-γ release assay QuantIFERON-TB Gold In-Tube (QIAGEN, Hilden, Germany) at study entry. PBMC were isolated from freshly collected heparinized blood via density centrifugation and cryopreserved. For this study, we chose a single donor with latent tuberculosis from whom sufficient PBMC were archived for the proposed experiments.

Ethics

Clinical study protocols were approved by the institutional review boards of the University of Cape Town and the University of Washington. Written informed consent was obtained from parents and/or legal guardians of the adolescents who participated. In addition, written informed assent was obtained from the adolescents.

Data analysis and statistics

Initial compensation, gating, and quality assessment of flow cytometry data were performed using FlowJo version 9.9.6 (FlowJo, TreeStar, Ashland, OR). Boolean gating and subsequent analysis was performed using the OpenCyto framework in the R programming environment (52, 53). Cell frequencies were compared using Wilcoxon signed-rank tests and results are reported as unadjusted p values. Two pre-BCG PBMC samples were excluded during analysis because of low viability and low CD3 cell counts.

Uniform Manifold Approximation and Projection (UMAP) was performed using the uwot package in R (L. McNees, J. Healy, and J. Melville, manuscript posted on arXiv, DOI: 1802.03426) with the following parameters: spread = 10 and min_dist = 0.15 for the Mamu CD1c–GMM tetramer data; spread = 16 and min_dist = 0.2 for the H. sapiens CD1d–α-GalCer tetramer data; and spread = 9 and min_dist = 0.02 for the Mamu MR1–5-OP-RU tetramer data. For the analysis of data generated using Mamu MR1 tetramer, cells were sampled evenly from each batch, as required for each experiment, and analyzed. However, for data generated using Mamu-CD1c and H. sapiens–CD1d tetramers, animals were unevenly sampled because of low cell counts. The following markers were used in the UMAP analysis: CD45RA, CD69, CD103, CCR7, CD38, γδ, CD4, 82, CD8, HLA-DR, CD3, Mamu CD1c–GMM PE-Cy7, Mamu CD1c–GMM BV560, H. sapiens CD1d–α-GalCer, and Mamu MR1–5-OP-RU. Fluorescence intensities of each marker were scaled within each batch prior to running UMAP. The raw data and analysis code used to generate the figures in this manuscript are available upon reasonable request.

Results

Evolutionary genetics of group 1 CD1 genes among primates

We recently showed that a five-gene CD1 locus is present among all primate clades and that CD1d shows evidence of purifying selection during primate evolution. This allowed us to use H. sapiens CD1d tetramer loaded with α-GalCer to study the molecular mechanisms of Ag recognition by iNKT cells in rhesus macaques (35). To further delineate the evolutionary history of simian CD1A, CD1B, and CD1C genes, CD1 homolog sequences were mined from 11 primate genomes that include great apes, Old World monkey, and New World monkeys, spanning ~45 million years of evolutionary time. In contrast to the very high conservation of CD1D, tests of molecular evolution across primates found significant signatures of diversifying (positive) selection in CD1A, CD1B, and CD1C (Table I). The signal is most robust in CD1A, which was also found to have experienced gene duplication independently in both olive baboon and marmosets (35). When the sites under positive selection were visualized on H. sapiens–derived crystal structures, most sites were localized within the lipid Ag binding pocket (Fig. 1A) (54–56). These data reveal that CD1A, CD1B, and CD1C have undergone diversifying selection during primate evolution, possibly leading to differences in the ability to bind and present lipid Ags to T cells.

GMM is presented by CD1b to T cells in macaques

We used the sequences for CD1b and CD1c obtained from cynomolgus macaques (M. fascicularis) and rhesus macaques (M. mulatta) to generate species-specific CD1b and CD1c tetramers (Supplemental Fig. 1). We used an established loading protocol to generate Mamu CD1b loaded with C32-GMM and tested its ability to stain a human GMM-specific T cell line (G02) (17). We found that Mamu CD1b-GMM successfully stained human T cells compared with mock-loaded control tetramer but at a lower mean fluorescence intensity compared with H. sapiens CD1b-GMM (Fig. 1B). The human GMM-specific T cell repertoire consists of high-avidity T cells expressing TRAV1-2 or low-avidity T cells expressing TRBV4-1 (57, 58). We considered the possibility that Mamu CD1b-GMM might be missing some of these specific subpopulations. However, we found that Mamu CD1b-GMM detected similar proportions of T cells expressing TRAV1-2 or TRBV4-1 as H. sapiens CD1b-GMM (Fig. 1C). Mamu CD1b-GMM tetramer stained 0.1% of T cells in PBMC of a BCG-vaccinated rhesus macaque compared with a mock-loaded control tetramer (Fig. 1D). These T cells were sorted and successfully expanded ~100-fold into a T cell line (bG52R). Because human PBMC were used as feeder cells, we verified the species origin of these T cells by staining with two anti-CD3e Abs, one that is cross-reactive with human and NHP and a second that is only reactive against human CD3e (Fig. 1E). We noted that nearly all the Mamu CD1b-GMM–specific T cells failed to stain with Abs targeting CD4 or CD8 coreceptors, which is in contrast to studies using H. sapiens CD1b-GMM that have revealed mostly CD4+ T cells in humans (Fig. 1F) (58). Finally, we noted that T cells that were identified using Mamu CD1b-GMM tetramer largely fail to costain with H. sapiens CD1b-GMM tetramer (Fig. 1G). Thus, GMM-specific T cells are present within the CD1b-restricted T cell repertoire of rhesus macaques but are largely “invisible” without a species-specific tetramer to identify them.

GMM is presented to mutually exclusive T cell populations by CD1b and CD1c in humans

Several studies have confirmed that GMM is presented to T cells by CD1b in humans (12, 17, 58, 59), so it was not surprising to find the same was true in rhesus macaques. However, a recent study examining rhesus macaques after repeated vaccination with BCG and liposomal GMM suggested that the dominant mechanism by which GMM is presented to T cells might be mediated by CD1c instead of CD1b (39). To investigate this possibility in humans, we incubated PBMC from a South African adolescent with latent tuberculosis infection with allogeneic dendritic cells that express all human CD1 isoforms and C32-GMM, allowing for selective expansion of Ag-specific T cells. Two weeks later, we noted that 0.4% of T cells stained with H. sapiens CD1b–C32-GMM tetramer but not with mock-loaded CD1b control tetramer (Fig. 2A). We also observed 0.1% of T cells stained with H. sapiens CD1c–C32-GMM tetramer but not with mock-loaded CD1c (Fig. 2A). These cells were independently sorted and successfully expanded ~100-fold into T cell lines (Fig. 2B). The CD1c-GMM–restricted T cell line (cG11) stained only with CD1c-GMM tetramer and not with CD1c mock-loaded or CD1b-GMM tetramer (Fig. 2B, 2C). The CD1b-GMM–restricted T cell line (bG11) stained only with CD1b-GMM tetramer and not with CD1b mock-loaded or CD1c-GMM tetramer (Fig. 2B, 2C). Both bG11 and cG11 also largely failed to stain with CD1b and CD1c tetramers loaded with irrelevant mycobacterial lipid Ags (Ac2SGL and PM, respectively) (Fig. 2D) (18, 20). We used K652 cells stably transfected with

i.v. BCG AND MYCOBACTERIAL GLYCOLIPID-SPECIFIC T CELLS IN NHP
Table I. Summary of tests of molecular evolution across primates in CD1a, CD1b, and CD1c paralogs

| CD1 Paralog | M8A (Neutral Model) | M8 (Positive Selection) | 2ΔlogL | p Value |
|-------------|----------------------|-------------------------|--------|---------|
| CD1A        | ρ0 = 0.4533, p = 0.0050, q = 67.7584 | ρ0 = 0.8848, p = 0.0245, q = 0.0209 | 58.4007 | 1.1 × 10⁻¹⁴ |
| CD1B        | ρ0 = 0.5473, p = 0.0050, q = 40.4342 | ρ0 = 0.6643, p = 0.0050, q = 2.6196 | 3.2376 | 0.036 |
| CD1C        | ρ0 = 0.4527, ρ1 = 1.0000, q = 10.9279 | ρ0 = 0.3357, ρ1 = 1.6799 | 16.4367 | 2.5 × 10⁻⁵ |

For each paralog, sequences were aligned using Clustal Ω, and tests of positive selection were performed using the codeml application in PAML (v4.8) with the primate species tree. Nested models of neutral evolution (M8a) and one allowing for positive selection (M8) were compared. In both models, the fraction of sites under purifying selection (ρ0) are described by evolutionary rates (nonsynonymous to synonymous substitution rates, or dN/dS) drawn from a β distribution with parameters p and q. Under the neutral M8a model, the remaining sites (ρ1 = 1 – ρ0) are described by an additional dN/dS term (ω) constrained to 1. Under the positive-selection M8 model, ω is a free parameter that can be > 1. The two models were compared by likelihood-ratio test with 1 df, and if significant, sites with Bayes-empirical Bayes-posterior probabilities ≥0.8 were considered under positive selection. CD1A and CD1C show stronger evidence of positive selection compared with CD1B during primate evolution.

Among CD1b-GMM-specific T cells within the bG11 T cell line, we noted that 8% expressed TRAV1-2, and 0.4% expressed TRBV4-1, whereas 2.5% expressed both TCR genes (Fig. 2G). Among CD1c-GMM-specific T cells within the cG11 T cell line, we noted that 99.9% stained with TRBV4-1 and found almost no evidence of TRAV1-2 expression (Fig. 2G). These data suggest that TRBV4-1 may be more important than TRAV1-2 for CD1c-mediated recognition of GMM (24, 25). The volume of the

FIGURE 1. GMM is presented by CD1b to T cells in macaques. (A) Evolutionary genetics of group 1 CD1 genes among primates. CD1a, CD1b, and CD1c sequences were collated, aligned, and compared as described in the Materials and Methods. Amino acid residues under positive selection were modeled onto H. sapiens CD1 crystal structures using PyMOL v2.4 and are noted in red. (Protein Data Bank numbers 5J1A, 5L2J, and 3OV6 for CD1a, CD1b, and CD1c, respectively). (B) H. sapiens CD1b-GMM and Mamu CD1b-GMM tetramers were used to identify GMM-specific T cells present within a human T cell line (G02) by staining with both GMM-loaded (BV421) and mock-loaded (FITC) CD1b tetramers. Mean fluorescence intensities (MFI) of populations identified by H. sapiens and Mamu CD1b-GMM tetramer present within G02 were identified by staining with either H. sapiens CD1b-GMM or Mamu CD1b-GMM tetramers, and TCR genes expressed by GMM-specific T cells were identified using Abs targeting TRAV1-2 and TRBV4-1. (D) PBMC from an Indian-origin rhesus macaque that was vaccinated intradermally with BCG were stained with Mamu CD1b-GMM and Mamu CD1b mock-loaded tetramers, sorted, and expanded in vitro as described in the Materials and Methods. The resulting T cell line (bG52R) was restained with Mamu CD1b-GMM and Mamu CD1b mock-loaded tetramer. (E) The species origin of bG52R was confirmed by staining with an anti-CD3ε Ab (clone UCHT1) that does not bind with Mamu CD3ε as well as an anti-CD3ε Ab (clone SP34-2) that binds CD3ε from both species. (F) T cells staining with Mamu CD1b-GMM tetramer present within bG52R were further examined for coreceptor expression using Abs targeting CD4 and CD8α. (G) bG52R was simultaneously stained with Mamu CD1b-GMM tetramer (BV421) and H. sapiens CD1b-GMM tetramer (allophycocyanin). Data are representative of at least two independent experiments. See also Supplemental Fig. 1 and Table I.
binding pocket of CD1b is nearly 50% larger than that of CD1c, so it is unclear whether the smaller CD1c cleft might accommodate C80-GMM (60). We tested the ability of cG11 or a CD1b-restricted T cell line (G10) to recognize long-chain (C80) or short-chain (C32) GMM when presented by CD1-transfected K562 cells. The CD1b-restricted T cell line was able to recognize both C80- and C32-GMM tetramer loading in vitro to generate the bG11 T cell line. T cells stained with H. sapiens CD1b-GMM tetramer were sorted and expanded in vitro to generate the cG11 T cell line. (C) bG11 stains with H. sapiens CD1b-GMM tetramer (PE) but fails to costain with H. sapiens CD1c-GMM tetramer (allophycocyanin). cG11 stains with H. sapiens CD1c-GMM tetramer but fails to costain with H. sapiens CD1b-GMM tetramer. (D) bG11 stains specifically with H. sapiens CD1b-GMM tetramer (allophycocyanin) and fails to costain with H. sapiens CD1c-GMM tetramer (PE) and fails to costain with H. sapiens CD1c (allophycocyanin) loaded with PM, another CD1c-presented mycobacterial lipid Ag. cG11 stains specifically with H. sapiens CD1c-GMM tetramer (PE) and fails to costain with H. sapiens CD1b (BV421) loaded with Ac2SGL, another CD1b-presented mycobacterial lipid Ag. (E) bG11 or (F) cG11 were coincubated with K562 cells that were stably transfected with H. sapiens CD1a, CD1b, CD1c, CD1d, or EV, and INF-γ production was measured by ELISPOT. (G) T cells within bG11 that stained with H. sapiens CD1b-GMM tetramer or T cells within cG11 that stained with H. sapiens CD1c-GMM tetramer were stained with Abs targeting TRAV1-2 and TRBV4-1 to investigate TCR gene usage. (H) A CD1b-GMM–restricted T cell line (G10) was coincubated with K562-CD1b–transfected APCs, and a CD1c-GMM–restricted T cell line (cG11) was coincubated with K562-CD1c–transfected APCs in the presence of titrating concentrations of short-chain (C32) or long-chain (C80) GMM. INF-γ production was measured by ELISPOT. Data are representative of at least three (A–C and G) or two (D–F and H) independent experiments.

**FIGURE 2.** GMM is presented to mutually exclusive T cell populations by CD1b and CD1c in humans. (A) PBMC from a healthy South African adolescent were incubated with allogeneic dendritic cells, GMM, and IL-2 for 2 wk. The resulting T cell line was stained with either H. sapiens CD1b-GMM (allophycocyanin) and H. sapiens CD1b mock-loaded (PE) tetramers or H. sapiens CD1c-GMM (allophycocyanin) and H. sapiens CD1c mock-loaded (PE) tetramers to identify GMM-specific T cells. (B) T cells staining with H. sapiens CD1b-GMM tetramer were sorted and expanded in vitro to generate the bG11 T cell line. T cells stained with H. sapiens CD1c-GMM tetramer were sorted and expanded in vitro to generate the cG11 T cell line. (C) bG11 stains with H. sapiens CD1b-GMM tetramer (PE) but fails to costain with H. sapiens CD1c-GMM tetramer (allophycocyanin). cG11 stains with H. sapiens CD1c-GMM tetramer but fails to costain with H. sapiens CD1b-GMM tetramer. (D) bG11 stains specifically with H. sapiens CD1b-GMM tetramer (allophycocyanin) and fails to costain with H. sapiens CD1c-GMM tetramer (PE) and fails to costain with H. sapiens CD1c (allophycocyanin) loaded with PM, another CD1c-presented mycobacterial lipid Ag. cG11 stains specifically with H. sapiens CD1c-GMM tetramer (PE) and fails to costain with H. sapiens CD1b (BV421) loaded with Ac2SGL, another CD1b-presented mycobacterial lipid Ag. (E) bG11 or (F) cG11 were coincubated with K562 cells that were stably transfected with H. sapiens CD1a, CD1b, CD1c, CD1d, or EV, and INF-γ production was measured by ELISPOT. (G) T cells within bG11 that stained with H. sapiens CD1b-GMM tetramer or T cells within cG11 that stained with H. sapiens CD1c-GMM tetramer were stained with Abs targeting TRAV1-2 and TRBV4-1 to investigate TCR gene usage. (H) A CD1b-GMM–restricted T cell line (G10) was coincubated with K562-CD1b–transfected APCs, and a CD1c-GMM–restricted T cell line (cG11) was coincubated with K562-CD1c–transfected APCs in the presence of titrating concentrations of short-chain (C32) or long-chain (C80) GMM. INF-γ production was measured by ELISPOT. Data are representative of at least three (A–C and G) or two (D–F and H) independent experiments.

GMM is presented by CD1c to T cells in macaques

Having established that GMM is presented to T cells by CD1b and CD1c in humans, we proceeded to investigate this paradigm in NHP. We generated Mamu CD1c–C32-GMM tetramers and stained PBMC obtained from a rhesus macaque 4 wk after intradermal BCG vaccination. We noted that Mamu CD1c-GMM tetramer stained 0.02% of T cells compared with Mamu CD1c-mock tetramer (Fig. 3A). These T cells were sorted and successfully expanded ~100-fold into a T cell line (cG52R) (Fig. 3B). We verified the species origin of these T cells by staining with two anti-CD3ε Abs before (Fig. 3C). Within cG52R, we noted the majority of T cells expressed CD4, in contrast to bG52R, which was predominantly CD4−CD8+ (Figs. 1F, 3D). Finally, we noted that T cells that were identified using Mamu CD1c-GMM tetramer largely failed to costain with H. sapiens CD1c-GMM tetramer, demonstrating species specificity of the T cell response (Fig. 3E). However, GMM-specific T cells within cG52R were partially activated by K562 cells transfected with H. sapiens CD1c in the presence of GMM, demonstrating some cross-reactivity using a functional assay (Fig. 3F). These data reveal that in rhesus macaques, as with humans, C32-GMM is presented by both CD1b and CD1c to two mutually exclusive populations of T cells.

Resident memory MAIT and iNKT cells are present in macaque lungs after i.v. BCG

Taking advantage of the donor-unrestricted nature of T cells that recognize these nonpolymorphic Ag-presenting molecules, we next...
sought to investigate whether mycobacterial vaccination might lead to the expansion of CD1- and MR1-restricted T cells and whether these T cells might be found among T cells resident in the lung (62, 63). To accomplish this, we made use of archived PBMC and BAL from our recent study, which demonstrated the superiority of i.v. BCG vaccination over standard routes in conferring protection from our recent study, which demonstrated the superiority of i.v. BCG vaccination over standard routes in conferring protection from M. tuberculosis challenge (6). We developed a multiparameter flow cytometry panel to measure memory (CD45RA and CCR7), activation (HLA-DR and CD38), tissue-residency (CD69 and CD103), and T cell (CD4, CD8, and γδ) phenotypes of GMM-specific T cells with additional tetramers to profile mucosal-associated invariant T (MAIT) cells and iNKT cells. Our experiments were designed to answer two statistically independent questions: 1) are donor-unrestricted T cells induced in PBMC after i.v. BCG, and 2) is the distribution of donor-unrestricted T cells different between PBMC and BAL at the peak of the response to i.v. BCG?

Cryopreserved samples from 10 animals were chosen for analysis consisting of PBMC at baseline, PBMC at 4 wk postvaccination, and BAL at 4 wk postvaccination. When comparing 4 wk after i.v. BCG vaccination to baseline, we found no change in the frequency of MAIT cells (p = 0.109) (Supplemental Fig. 2A, 2B) nor in the frequency of iNKT cells (p = 0.078) (Supplemental Fig. 2C, 2D) in the blood. Although we did see that MAIT cells were found at higher frequency in the BAL compared with PBMC (p = 0.008) (Supplemental Fig. 2B), this did not hold true for iNKT cells (p = 0.734) (Supplemental Fig. 2D). These data are consistent with our published data, which revealed a selective induction of MAIT cells in BAL after i.v. BCG when compared with both aerosol and intradermal administration (6). We then used UMAP to examine whether subpopulations of MAIT or iNKT cells were expanded after i.v. BCG (Fig. 4A, 4F, Supplemental Fig. 3). Indeed, we observed an increase in the frequency of MAIT cells in PBMC expressing a CD45RA+CCR7+TEM phenotype (p = 0.023) (Fig. 4A–C). We also observed an increase in the frequency of MAIT cells expressing TEM markers CD69 and CD103 in BAL compared with PBMC at the 4 wk time point (p = 0.004) (Fig. 4A, 4D, 4E). We observed similar expansions of iNKT cells expressing TEM and RM markers in blood (p = 0.078) and BAL (p = 0.008), respectively (Fig. 4F–J). Notably, T cells with a TEM phenotype were broadly expanded in the blood 4 wk after i.v. BCG (p = 0.023), and T cells expressing RM markers were broadly expanded in the BAL compared with the blood (p = 0.004) (Supplemental Fig. 2E, 2F). These data extend our published findings by revealing TEM phenotypes in the lung as well as expansions of MAIT and iNKT cell subsets expressing TEM phenotypes in the blood after i.v. BCG vaccination.

GMM-specific T cells are expanded after i.v. BCG

Finally, we examined the effect of i.v. BCG on CD1c-GMM–specific T cells (Fig. 5, Supplemental Fig. 3). Limiting cell numbers precluded us from simultaneously profiling CD1b-GMM–specific T cells in all samples. We prioritized the detection of CD1c-GMM–specific T cells based on the report by Morita et al. (39), which suggested that this was the dominant mechanism of GMM recognition by T cells in macaques. Specificity of staining and accurate quantification of low-frequency events were achieved by using two Mamu CD1c-C32-GMM–loaded tetramers in different fluorescence channels as well as a mock-loaded negative-control CD1c tetramer (Fig. 5A, Supplemental Fig. 4). At baseline, we noted that the majority of Mamu CD1c–GMM–specific T cells expressed either a CD4 or CD8 coreceptor, which was consistent with our in vitro results (Figs. 3D, 5B). We observed an increase in the frequency of T cells staining with Mamu CD1c-GMM tetramers in PBMC at 4 wk after BCG vaccination compared with baseline (p = 0.016) (Fig. 5C). As with MAIT and iNKT cells, this expansion was especially pronounced within the TEM subset (p = 0.008) (Fig. 5D–F). We noted a positive correlation between the frequency of CD1c-GMM–specific T cells and both iNKT (r = 0.85; p = 0.002) and MAIT cells (r = 0.63; p = 0.051) expressing a TEM phenotype in the blood (Fig. 5G). By contrast, we observed a lower total frequency of T cells staining with Mamu CD1c-GMM tetramers in BAL compared with PBMC at the 4-wk time point (p = 0.02) (Fig. 5H). However, this was contrasted by a statistically significant difference in the subset of Mamu CD1c-GMM–specific T cells expressing TEM markers.
These data reveal that i.v. BCG leads to clonal expansion and terminal differentiation of GMM-specific T cells in the blood. GMM-specific T cells are also detectable as TRM cells in the lung.

Discussion

Our data advance several aspects of CD1 biology and tuberculosis vaccinology. First, we developed and validated Mamu CD1b-GMM and Mamu CD1c-GMM tetramers, establishing the NHP as a natural animal model in which to study the phenotypes and functions of CD1-reactive T cells in health and disease. Second, we discovered that both CD1b and CD1c can present GMM to T cells in humans and NHP, providing a potential mechanism for how primates evolved the capacity to discriminate among Ags with identical head groups but various lipid chain lengths. Third, we discovered that i.v. BCG leads to the expansion of GMM-specific T cells in the blood, which are also detectable in BAL fluid, suggesting that lipid Ag-specific T cells may contribute to the protective immune response to *M. tuberculosis*.

We specifically chose to study GMM-specific T cells in the setting of i.v. BCG vaccination because this route has been shown to provide superior protection compared with intradermal or aerosol vaccination in rhesus macaques, which are exquisitely susceptible to *M. tuberculosis* infection and disease (6, 7, 64). In our prior work, most animals vaccinated with i.v. BCG showed evidence of sterilizing immunity and had substantial expansions of MAIT cells in the lung at the time of *M. tuberculosis* challenge (6). The natural question is whether particular Ags are being targeted by lung T cells. Much effort is sure to be spent on distinguishing among the ~4000 proteins expressed by *M. tuberculosis*. Our data suggest that T cells mediating protective immunity may also be targeting lipids that are shared between BCG and *M. tuberculosis*, such as MA and GMM (65).

Morita et al. (39) were the first to describe the presentation of GMM by CD1c rather than CD1b in rhesus macaques by using an unusual vaccination strategy. GMM was purified from cultured mycobacteria and CD1c dependence was demonstrated through the use of blocking Abs and transfected APCs. Our data confirm

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**FIGURE 4.** Resident memory MAIT and iNKT cells are present in macaque lungs after i.v. BCG. Multiparameter flow cytometry was used to profile T cells present in PBMC prior to i.v. BCG vaccination (pre-BCG) and in PBMC and BAL at 4 wk after BCG vaccination in 10 rhesus macaques. MAIT cells were identified using *Mamu* MR1–5-OP-RU loaded tetramer and iNKT cells were identified using *H. sapiens* CD1d–α-GalCer–loaded tetramer. (A) Expression of CD45RA, CCR7, CD69, and CD103 on MAIT cells is displayed after aggregating across all animals and performing dimensionality reduction with UMAP. Mean fluorescence intensities (MFI) were scaled to achieve a mean = 0 and SD = 1. (B) Representative staining of CD45RA and CCR7 among CD8+ MAIT cells identified in PBMC from one animal pre-BCG or 4 wk after i.v. BCG vaccination. (C) The frequency of CD4+ and CD8+ MAIT cells expressing a T EM phenotype was compared with pre-BCG and at 4 wk after i.v. BCG vaccination in all 10 animals. (D) Representative staining of CD69 and CD103 among CD8+ MAIT cells identified in PBMC and BAL at 4 wk after i.v. BCG vaccination. (E) The frequency of CD4+ and CD8+ MAIT cells expressing a T RM phenotype was compared in PBMC from one animal pre-BCG or 4 wk after i.v. BCG vaccination. (G) The frequency of CD4+ and CD8+ iNKT cells expressing a T EM phenotype was compared pre-BCG and at 4 wk after i.v. BCG vaccination. (H) Representative staining of CD69 and CD103 among CD8+ iNKT cells identified in PBMC and BAL at 4 wk after i.v. BCG vaccination. (I) The frequency of CD4+ and CD8+ iNKT cells expressing a T RM phenotype was compared pre-BCG and at 4 wk after i.v. BCG vaccination in all 10 animals. Results from line plots represent data from all animals. Statistical testing was performed using Wilcoxon signed-rank tests, and unadjusted *p* values are reported. The experiment was performed once. See also Supplemental Figs. 2 and 3.
and extend these findings through the use of synthetic GMM, newly validated Mamu CD1c tetramers, paired comparisons between blood and BAL, and context within i.v. BCG. In one representative animal, there is an apparent increase in the frequency of T cells staining with both GMM-loaded tetramers at 4 wk after i.v. BCG. (D) Expression of CD45RA, CCR7, CD69, and CD103 on T cells staining with Mamu CD1c-GMM tetramer is displayed after aggregating across all animals and performing dimensionality reduction with UMAP. Mean fluorescence intensities (MFI) were scaled to achieve a mean = 0 and SD = 1. (E) Representative staining of CD45RA and CCR7 among T cells staining with Mamu CD1c-GMM tetramer in one animal pre-BCG or 4 wk after i.v. BCG vaccination. (F) The frequency of Mamu CD1c-GMM staining T cells expressing a TEM phenotype was compared pre-BCG and at 4 wk after i.v. BCG vaccination in all animals. (G) The frequency of cells expressing a TEM phenotype as a percentage of total tetramer staining cells in the blood is plotted for CD1c-GMM versus iNKT cells, CD1c-GMM versus MAIT cells, or iNKT versus MAIT cells. Pearson correlation R and nominal p values are displayed for each comparison. (H) Frequencies of T cells staining with Mamu CD1c-GMM tetramer as a percentage of total CD3+ cells were compared between PBMC and BAL at 4 wk after i.v. BCG vaccination. (I) Representative staining of CD69 and CD103 among T cells staining with Mamu CD1c-GMM tetramer in PBMC and BAL at 4 wk after i.v. BCG vaccination. (J) The frequency of T cells staining with Mamu CD1c-GMM tetramer expressing a TEM phenotype was compared between PBMC and BAL at 4 wk after i.v. BCG vaccination. Statistical testing was performed using Wilcoxon signed-rank tests, and unadjusted p values are reported. The experiment was performed once. See also Supplemental Figs. 3 and 4.
detectable in the lungs 4 wk after i.v. BCG and express an activated phenotype. Because we have previously reported that residual BCG is detectable in BAL at this time point, it is possible that some of these cells may also represent activated T cells because of the persistence of mycobacterial Ag (6). Whether they persist as true resident memory T cells, express antibacterial functions, and contribute directly to reducing bacterial burdens in the lung is the subject of future work.

When we used H. sapiens CD1b-GMM and H. sapiens CD1c-GMM tetramer to isolate GMM-specific T cell lines, we found that they stained only with the tetramer used to isolate them. These data suggest that the TCR repertoires identified by these tetramers are mutually exclusive. CD1b-restricted T cells specific for GMM can express TRAV1-2 or TRBV4-1 as well as diverse TCRs (57, 70). CD1c-autoreactive T cells have been shown to preferentially express TRBV4-1 (25). A detailed molecular study of both GMM-specific and autoreactive CD1b-restricted T cells showed that germline-encoded residues within TRBV4-1 are essential for binding both CD1b and CD1c independent of the ligand (24). We find that nearly all T cells that stain with H. sapiens CD1c-GMM express either TRAV1-2 or TRBV4-1, but only a subset of T cells identified by H. sapiens CD1b-GMM express either TRAV1-2 or TRBV4-1. This is consistent with a diverse repertoire among H. sapiens CD1b-GMM T cells that may not hold for H. sapiens CD1c (70).

Validation of Mamu CD1b and CD1c tetramers and the discovery of tissue phenotypes of CD1c-GMM-specific T cells in rhesus macaques represents direct evidence for tissue-specific, polyclonal responses to CD1-presented lipid Ags. Studies in humans have advanced our understanding of the structure, function, and regulation of CD1 expression (60, 71). Access to peripheral blood and tissues have enabled the study of T cell phenotypes and regulation of CD1 expression (60, 71). Access to peripheral blood and tissues have enabled the study of T cell phenotypes and regulation of CD1 expression (60, 71). Access to peripheral blood and tissues have enabled the study of T cell phenotypes and regulation of CD1 expression (60, 71). Access to peripheral blood and tissues have enabled the study of T cell phenotypes and regulation of CD1 expression (60, 71). Access to peripheral blood and tissues have enabled the study of T cell phenotypes and regulation of CD1 expression (60, 71). Access to peripheral blood and tissues have enabled the study of T cell phenotypes and regulation of CD1 expression (60, 71).
