Identification of MAGE-3 Epitopes Presented by HLA-DR Molecules to CD4⁺ T Lymphocytes

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Summary
MAGE-type genes are expressed by many tumors of different histological types and not by normal cells, except for male germline cells, which do not express major histocompatibility complex (MHC) molecules. Therefore, the antigens encoded by MAGE-type genes are strictly tumor specific and common to many tumors. We describe here the identification of the first MAGE-encoded epitopes presented by histocompatibility leukocyte antigen (HLA) class II molecules to CD4⁺ T lymphocytes. Monocyte-derived dendritic cells were loaded with a MAGE-3 recombinant protein and used to stimulate autologous CD4⁺ T cells. We isolated CD4⁺ T cell clones that recognized two different MAGE-3 epitopes, MAGE-3114–127 and MAGE-3121–134, both presented by the HLA-DR13 molecule, which is expressed in 20% of Caucasians. The second epitope is also encoded by MAGE-1, -2, and -6. Our procedure should be applicable to other proteins for the identification of new tumor-specific antigens presented by HLA class II molecules. The knowledge of such antigens will be useful for evaluation of the immune response of cancer patients immunized with proteins or with recombinant viruses carrying entire genes coding for tumor antigens. The use of antigenic peptides presented by class II in addition to peptides presented by class I may also improve the efficacy of therapeutic antitumor vaccination.

Key words: human • invariant chain • peptide • tumor • histocompatibility leukocyte antigen class II

From cultures of blood lymphocytes or tumor-infiltrating lymphocytes of cancer patients stimulated with autologous tumor cells, it is possible to isolate CTLs that show specificity for tumor cells (1, 2). These CTLs have been used as tools to isolate genes that code for tumor antigens, such as those of the MAGE gene family (3, 4). The MAGE antigens are of particular interest for cancer immunotherapy because of their strict tumoral specificity and because they are shared by many tumors. MAGE genes are activated in tumors of many different histological types. They are silent in normal cells, except in testicular germ cells, which do not express MHC class I molecules and are therefore incapable of presenting antigens to CTLs (5, 6). Other families of genes with the same pattern of expression have been identified (7, 8). In addition to the genes identified with the approach involving antitumor CTLs, several other MAGE-type genes, i.e., genes expressed only in tumors and in male germline cells, have been identified by purely genetic approaches or by the use of antibodies present in the sera of cancer patients (9–11). As a result, there is now a large supply of sequences potentially coding for tumor-specific shared antigens.

Among the MAGE genes, MAGE-3 is one of the most frequently expressed in tumors. For instance, it is expressed in 76% of metastatic melanomas (12). Five antigenic peptides presented by class I molecules have been identified in the MAGE-3 protein. They are presented to CTLs by HLA-A1, -A2 (two epitopes), -A24, and -B44 (12–17). Some of these epitopes were identified as targets of CTL clones obtained by stimulation of T lymphocytes with au-
tologous tumor cells. Others were identified by “reverse immunology” approaches, based on the knowledge of gene sequences and in vitro stimulation of lymphocytes of non-cancerous individuals with candidate peptides considered likely to bind to a given HLA. MAGE-3 codes for a protein of 315 amino acids, and it is very likely that many other peptides derived from MAGE-3 bind to various HLA molecules and hence could be additional target antigens for an antitumor CTL response.

So far, the identification of tumor antigens has focused mainly on antigens recognized by CTLs, almost all of which are CD8+, even though the importance of CD4+ T cells in antitumor response has been demonstrated in numerous animal models (18–21). Only a few examples of antigens recognized by CD4+ T cells on human tumors have been described. CD4+ T cells that were raised in vitro against a peptide centered on the fusion region of bcr-abl were found to recognize HLA-DR4 leukemic blasts expressing bcr-abl (22). The same approach had been previously used to identify an epitope presented by HLA-DQ7 molecules in a mutated region of K-ras (23). Antigenic peptides encoded by the melanocyte differentiation proteins tyrosinase and gp100 have also been identified (24–26).

No antigen presented by a class II molecule has yet been reported for a MAGE-type gene. In an attempt to characterize such an antigen for MAGE-3, we used monocytederived dendritic cells loaded with a MAGE-3 recombinant protein to stimulate autologous CD4+ T cells. We report here the identification of MAGE-3 epitopes presented by HLA-DR molecules.

Materials and Methods

Cell Lines, Media, and Reagents. The EBV-transformed B (EBV-B) cell lines and tumor cell line M22-MEL.43 were cultured in IMDM (GIBCO BRL) supplemented with 10% FCS (GIBCO BRL), 0.24 mM l-asparagine, 0.55 mM l-arginine, 1.5 mM l-glutamine (AAG), 100 U/ml penicillin, and 100 µg/ml streptomycin. The PhoenixAMPHO cell line (provided by Dr. Nolan, Stanford University, Stanford, CA) is a high-titer amphotropic retrovirus–producing cell line that was generated by stable transfection of 293T cells with a Moloney amphotropic envelope gene driven by a CMV promoter and coselected with the diphtheria toxin resistance gene (pHED-7). This producer cell line is helper virus-free. The PhoenixAMPHO packaging cell line (provided by Dr. Nolan, Stanford University, Stanford, CA) is a high-titer amphotropic retrovirus–producing cell line that was generated by stable transfection of 293T cells with a Moloney amphotropic envelope gene driven by a CMV promoter and coselected with the diphtheria toxin resistance gene (pHED-7). This producer cell line is helper virus-free. The PhoenixAMPHO packaging cell line (provided by Dr. Nolan, Stanford University, Stanford, CA) is a high-titer amphotropic retrovirus–producing cell line that was generated by stable transfection of 293T cells with a Moloney amphotropic envelope gene driven by a CMV promoter and coselected with the diphtheria toxin resistance gene (pHED-7). This producer cell line is helper virus-free.

Human recombinant IL-2 was purchased from Eurocetus, IL-7 from Genzyme, GM-CSF from Schering Plough, and TNF-α from R&D Systems. Human recombinant IL-4, IL-6, and IL-12 were produced in our laboratory.

MAGE-3 Protein. The recombinant MAGE-3 protein was produced by SmithKline Beecham Pharmaceuticals. The full-length MAGE-3 sequence, preceded by the amino acid (aa) sequence MHHHHHTHGG, was engineered into a vector bearing the pM1B1 replicon and the PL short promoter. This vector was used to transform Esherichia coli strain AR58 (27). Bacteria were grown in LB medium containing 50 µg/ml kanamycin at 30°C. After heat induction, the MAGE-3 expression product became detectable as a 46-kD band when assayed by Western blot (SDS-PAGE 12.5%; revealed by a rabbit polyclonal antiserum to MAGE-3). Protein purification was carried out at room temperature, and involved the following steps: cell lysis and centrifugation, repeated washing of the centrifugation pellet followed by solubilization of the pellet containing MAGE-3, immobilized metal ion affinity chromatography, anion exchange chromatography, concentration, and dialysis. The resulting MAGE-3 protein was >95% pure, as assessed by Coomassie blue staining.

Construction of pMFG-II:MAGE-3. The plasmid lippSV51L, containing a cDNA encoding the human invariant chain (II), was provided by Dr. J. Pieters (Basel Institute for Immunology, Basel, Switzerland). The MFG plasmid was provided by Dr. O. Danos (Somatix Therapy Corp., Alameda, CA). The MFG retroviral vector is derived from the Moloney murine leukemia virus, which lacks a drug resistance marker and does not express any other potential antigenic protein except for the inserted cDNA (28). The cDNA encoding the N-terminal domain (i.e., the cytoplasmic tail and the transmembrane domain) of the human (hu)-li polypeptide (residues 1–80) was amplified by PCR using lippSV51L as the template. The following primers were used: hu-li sense, 5'-TTTGTAGATGGATGACCCGGCG-3', and hu-li antisense, 5'-TTTGGATCTCGGAGCTCATGCAGG-3' (the recognition sites for Ncol and BamHI are in italics). The PCR product was cloned into pCR2.1 and sequenced. The Ncol-BamHI amplification product was cloned into pMFG and open reading with the enzymes Ncol and BamHI, resulting in pMFG-II. A BglII recognition site, replacing the ATG codon and in frame with the BamHI site at the 3' end of the truncated li-cDNA, was introduced at the 5' end of the MAGE-3 cDNA by PCR using the following primers: BglII sense, 5'-TTTGGATCTCGGAGCTCATGCAGG-3', and BglII antisense, 5'-CCCGAGATCTTCATCCGGTTC-3' (the recognition sites for BglII are in italics). The PCR product (BglII, MAGE-3:BglII) was cloned into pCR2.1 and sequenced. The recombinant plasmid, pMFG-II, was reopened with BamHI, and the BglII:MAGE-3:BglII amplification product was ligated to the compatible ends. Recombinant plasmids containing the MAGE-3 cDNA in frame and in the right orientation were identified by restriction fragment analysis.

Production of High-titer Ii.MAGE-3-encoding Recombinant Retrovirus. The MAGE-3-encoding retroviral vector plasmid MFG-II:MAGE-3 was introduced into the PhoenixAMPHO packaging cells by transfection. The transfection procedure is a modification of the calcium phosphate–mediated transfection protocol of Graham and Van der Eb (29). 24 h before transfection, PhoenixAMPHO cells were plated in cell growth medium in a 75-cm2 tissue culture flask (Falcon; Becton Dickinson Labware). After adding the cells, the flask was gently shaken forward and backward to distribute cells evenly in the flask bottom. The cells were incubated at 37°C and 5% CO2. At the time of transfection, when the cells had reached a confluence of 70–80%, the medium was removed and replaced by 14 ml fresh PhoenixAMPHO cell growth medium containing 25 mM chlorouridine (Sigma). A transfection cocktail was prepared in a 50-ml tube by adding 40 µg retroviral vector plasmid DNA to water and diluting to 1,575 µl final volume. To this DNA solution 225 µl of 2 M CaCl2 was added. Then, 1,800 µl of 2× HEBS (50 mM Hepes, 10 mM KCl, 12 mM
were washed and added at 10^4 per round-bottomed microwell to 1 ml. The CD4^+ T lymphocytes were transferred to 4 ml of infection cocktail containing 50% viral supernatant, the retroviral supernatant was harvested after 20 h and the chloroquine-containing medium was changed for fresh PhoenixAMPHO cell growth medium. Approximately 24 h before the harvest of the retroviral supernatant, the PhoenixAMPHO medium was removed and gently replaced by 9 ml of IMDM containing only 2.5% FCS. The retroviral supernatant was harvested 48 h after transfection by removing the medium from the cells and filtering through a 0.45-μm filter to remove cell debris. After harvest and filtration, the virus-containing medium was kept on ice, aliquoted in appropriate volumes in 15-ml polypropylene tubes, and stored at −80°C.

Retroviral Transduction of Cell Lines. Target cells were resuspended in 60-mm tissue culture plates (Falcon) at a density of 10^6 cells in 4 ml of infection cocktail containing 50% viral supernatant in growth medium and 6 μg/ml of protamine sulfate. The plates were centrifuged for 2 h at 32°C and 1,200 rpm, followed by another 2 h of incubation in a humidified incubator at 37°C. Cells were then transferred to 4 ml of growth medium. This transduction cycle was carried out immediately after plating the cells and was repeated at 24 and 48 h.

Dendritic Cells. Blood cells were collected asuffy coat preparations from hemochromatosis patients. PBMCs were isolated by Lymphoprep (Nycomed Pharma) density gradient centrifugation. To generate autologous dendritic cells, PBMCs were depleted from T lymphocytes by rosetting with sheep erythrocytes (Bio Mérieux) treated with 2-aminoethylisothiouronium (Sigma). The lymphocyte-depleted PBMCs were treated with 2-aminoethylisothiouronium (Sigma). The lymphocyte-depleted PBMCs were transferred to 2 medium supplemented with IL-4 (100 U/ml) and GM-CSF (100 ng/ml). On day 5, the nonadherent cell population was used as a source of enriched dendritic cells.

Mixed Lymphocyte/Dendritic Cell Culture. Autologous dendritic cells (5 × 10^6/ml) were incubated at 37°C, 5% CO_2 for 18–20 h in complete medium supplemented with IL-4 (100 U/ml), GM-CSF (100 ng/ml), and TNF-α (1 ng/ml) in the presence of the recombinant MAGE-3 protein (20 μg/ml). Cells were washed and added at 10^6 per round-bottomed microwell to 10^2 CD4^+ lymphocytes in 200 μl IMDM supplemented with AAG and 10% human serum (hereafter referred to as complete IMDM) in the presence of IL-6 (1,000 U/ml) and IL-12 (10 ng/ml). The CD4^+ lymphocytes were restimulated on days 7, 14, and 21 with autologous dendritic cells freshly loaded with the MAGE-3 protein and were grown in complete IMDM supplemented with IL-2 (10 U/ml) and IL-7 (5 ng/ml). Due to the limited supply of dendritic cells for melanoma patient 7002, we used only 6 × 10^3 dendritic cells instead of 10^4 and restimulations were performed on days 10 and 20 instead of 7, 14, and 21. The microcultures containing proliferating CD4^+ T cells were assessed on days 35–37 for their capacity to produce TNF and/or IFN-γ when stimulated with autologous EBV-B cells loaded with protein MAGE-3. Autologous EBV-B cells were incubated for 18–20 h in the presence of 20 μg/ml of protein MAGE-3, or OVA (Sigma) as a negative control. Protein-pulsed EBV-B cells were washed and distributed at 5,000 cells per round-bottomed microwell together with 2,500 CD4^+ T lymphocytes in 150 μl of complete IMDM supplemented with IL-2 (25 U/ml). After 20 h, the supernatant was collected and its TNF content was determined by testing its cytotoxic effect on WEHI-164 clone 13 cells (30) in a MTT colorimetric assay (31, 32). IFN-γ released in the supernatant was measured by ELISA using reagents from Medgenix Diagnostics-Biosource. Inhibition with mAbs W6/32 (anti–HLA class I) or 2B6 (anti–HLA-DR) was performed by addition of a 1:20 dilution of ascites during the experiment.

CD4^+ T Cell Clones. The microcultures that recognized cells loaded with the MAGE-3 protein were cloned by limiting dilution, using as stimulating cells either the autologous EBV-B cell line loaded with the MAGE-3 protein or the autologous EBV-B cell line transduced with a retrovirus encoding II.MAGE-3. Allogeneic EBV-B cells (LG2-EBV) were added as feeder cells. CD4^+ T cell clones were grown in complete IMDM supplemented with IL-2 (50 U/ml), IL-7 (5 ng/ml), and 0.5 μg/ml purified PHA (HA 16; Murex Diagnostics). The clones were supplemented with fresh culture medium once a week and passaged with feeder cells (1.5 × 10^6 allogeneic PBLs plus 5 × 10^6 LG2-EBV per well in a 24-well plate) at 1–2-wk intervals. Occasionally, clones were restimulated with 2 × 10^5 autologous EBV-B cells transduced with II.MAGE-3 as stimulating cells and 10^6 LG2-EBV. Established CD4^+ T cell clones were then tested for TNF and/or IFN-γ production after stimulation with autologous EBV-B cells pulsed with protein MAGE-3 and MAGE-3-derived peptides. Recognition A says with Peptides. Peptides were synthesized on solid phase using Fmoc for transient N H_2-tert-aminomethyl protection and were characterized using mass spectrometry. All peptides were >80% pure, as indicated by analytical HPLC. Lyophilized peptides were dissolved in DM SO and used at a concentration of 5 μg/ml. EBV-B cells were incubated for 2 h at 37°C in the presence of the different peptides, the indicated concentrations representing their concentrations during the incubation step. They were distributed at 5,000 cells per round-bottomed microwell together with 2,500 CD4^+ T lymphocytes in 150 μl of complete IMDM supplemented with IL-2 (25 U/ml). Supernatants were harvested after 20 h and assessed for TNF and/or IFN-γ production. The results represent the average of duplicate or triplicate cultures.
dritic cells were then washed, and 2,500 CD4+ lymphocytes were added in 150 μl of complete IMDM supplemented with IL-2 (25 U/ml). Supernatants were harvested after 20 h and assessed for IFN-γ production.

Results

Blood monocytes were cultured in medium supplemented with GM-CSF and IL-4 to favor their differentiation into dendritic cells (Fig. 1). Autologous plasma was used to avoid loading the dendritic cells with bovine or allogeneic proteins. After 5 d, the dendritic cells were incubated overnight with 20 μg/ml of a MAGE-3 protein produced in E. coli and with TNF-α to induce their maturation. 96 microcultures were set up with 10^6 autologous responder CD4+ T cells and 10^4 dendritic cells loaded with protein MAGE-3 as stimulator cells. IL-6 and IL-12 were added during the first week to activate the T cells. The responder T cells received three additional weekly restimulations with dendritic cells pulsed with protein MAGE-3 in the presence of IL-2 and IL-7. After a resting period of 2 wk, the responder cells of each microculture were tested on day 35 for TNF or, more often, for IFN-γ production after stimulation for 20 h with autologous EBV-B cells loaded with protein MAGE-3.

CD4+ T Cell Clones Directed against a MAGE-3 Antigen. CD4+ T cells of hemochromatosis patient LB 1555, stimulated with dendritic cells loaded with protein MAGE-3, were tested for their ability to produce TNF upon stimulation with EBV-B cells loaded with either protein MAGE-3 or OVA. It was possible to measure the production of TNF by the T cells because, contrary to most EBV-B cell lines, that of patient LB 1555 did not produce TNF. The microculture (B6 in Fig. 2) that produced the highest level of TNF after stimulation with protein MAGE-3 was cloned by limiting dilution using autologous EBV-B cells loaded with protein MAGE-3 as stimulator cells. These stimulator cells were used for the cloning step because of the limited supply of autologous dendritic cells. A positive CD4+ clone was obtained, referred to hereafter as clone 37. It recognized autologous EBV-B cells loaded with protein MAGE-3. Even though the protein, which had been produced in bacteria, was >95% pure, we could not at this stage exclude the possibility that clone 37 recognized a bacterial contaminant. Therefore, we prepared a set of peptides of 16 aa, which overlapped by 12 and covered the entire MAGE-3 protein sequence. Each of these peptides was incubated with autologous EBV-B cells and tested for recognition by clone 37. We identified two overlapping peptides, MAGE-3111-126 and MAGE-3115-130 (see below), that stimulated production of both TNF (data not shown) and IFN-γ (Fig. 3 A). This demonstrated unambiguously that clone 37 recognized a MAGE-3 antigen and not a contaminant.

A CD4+ clone was obtained from another microculture which was also very well stimulated by cells loaded with protein MAGE-3 (F3 in Fig. 2). However, here the screening with the set of overlapping MAGE-3 peptides was negative. We found that the antigen recognized by this CD4+ clone was present in an HPLC fraction other than the one containing the MAGE-3 protein (data not shown). Our tentative conclusion is that this clone recognizes a bacterial contaminant.

Additional CD4+ clones were obtained from five independent microcultures, set up with cells from another hemochromatosis patient. Their ability to recognize cells loaded with protein MAGE-3 appeared to decrease upon further purification of the protein by HPLC. Two of these clones were also tested for their response to the set of MAGE-3 peptides, with negative results, suggesting further that they recognized a contaminant in the MAGE-3 batch. Therefore, it appears that bacterial contaminants in this MAGE-3 protein batch were more likely to activate CD4+ T cells than was MAGE-3 itself, so that CD4+ clones specific for MAGE-3 constituted only a minor fraction of the clones obtained with this procedure.

Figure 1. Overview of the procedure used to obtain anti-MAGE-3 CD4+ T cell clones.

770 MAGE-3 Epitopes Presented by HLA Class II Molecules
Clone 37 recognizes Peptide AELVHFLLLKYRAR on HLA-DR13. CD4+ clone 37 recognized two peptides which overlapped by 12 aa, namely RKVAELVHFLLLKYRA (aa 111–126) and ELVHFLLLKYRAREPV (aa 115–130) (Fig. 3 A). The recognition by clone 37 of cells loaded with protein MAGE-3 was abolished by an anti–HLA-DR antibody (data not shown). Patient LB 1555 was serologically typed DR3, DR13, and DR52. To identify the presenting HLA-DR molecule, we tested additional EBV-B cell lines expressing DR3, DR13, or DR52. All and only those expressing DR13 were able to present the MAGE-3111–126 and MAGE-3115–130 peptides to clone 37 (Table I). 31 different HLA-DR13 alleles have been described to date (33). LB 1555 expresses the DRB1*1302 allele, and the other DR13 cell lines tested express DRB1*1301 or DRB1*1302, the latter two representing 80% of the HLA-DR13 alleles.

Unlike the peptides presented by class I molecules, those presented by class II usually vary in length and tolerate extensions at both the NH2 and COOH termini because they are not fixed by their ends in the groove (34). We tested a large number of MAGE-3 peptides of different lengths (data not shown) and concluded that the shortest peptide well recognized by clone 37 is AELVHFLLLKYRAR (aa 114–127; Fig. 3 B). Half-maximum value of stimulation was obtained by incubating the stimulator cells with 10 nM of this peptide. This compares favorably with the results obtained with other epitopes recognized on HLA class II molecules by CD4+ T cells: a minimum of 30 nM of a bcr-abl peptide was needed to induce a significant proliferation of an anti–bcr-abl CD4+ cell line, whereas 330 nM of peptide p21-ras was required for anti-p21-ras CD4+ clones (22, 23).

### Table I. HLA-DR13 Cells Present the Two Peptides MAGE-3111–126 and MAGE-3115–130 to CD4+ Clone 37

| EBV-B cell line | Serological specificity | IFN-γ production by clone 37 stimulated with peptide |
|-----------------|------------------------|----------------------------------------------------|
| LB 1555         | DR3⁺ DR52⁺             | >4,000     >4,000                                 |
| LB 1118         | DR3⁺ DR52⁺             | 3,761      3,909                                 |
| LB 1622         | DR3⁺ DR52⁺             | 1,731      1,349                                 |
| OMW             | DR3⁻ DR52⁺             | >4,000     >4,000                                 |
| MZ2             | DR3⁻ DR52⁻             | 3,429      3,096                                 |
| BM16            | DR3⁻ DR52⁺             | 99         120                                   |
| BOB             | DR3⁻ DR52⁺             | 128        101                                   |
| BOLETH          | DR3⁻ DR52⁺             | 77         67                                    |

Autologous LB 1555 EBV or allogeneic EBV-B cells were incubated for 2 h with 5 μg/ml of peptides MAGE-3111–126 and MAGE-3115–130, and washed. Clone 37 was then incubated for 20 h with 5,000 peptide-pulsed EBV-B cells IFN-γ released in the supernatant was measured by ELISA.

**Figure 2.** Production of TNF by CD4+ T cells stimulated with autologous EBV-B cells incubated with protein MAGE-3. EBV-B cells from hemochromatosis patient LB 1555 were incubated with 20 μg/ml of OVA or MAGE-3 protein for 20 h. 5,000 cells were distributed in microsolves to which an aliquot of each of the 96 CD4 microcultures was added (~3,000 effector cells). TNF production was estimated after overnight culture, by the toxicity of the supernatants for the TNF-sensitive WEHI 164-13 cells. The dots represent the average result obtained with two aliquots of each CD4 microculture.

**Figure 3.** MAGE-3 peptides recognized by clone 37. (A) Stimulation of clone 37 by two overlapping MAGE-3 peptides. Autologous EBV-B cells were incubated for 2 h with different concentrations of the peptides. Clone 37 (2,500 cells) was then cocultured with 5,000 peptide-pulsed cells for 20 h. IFN-γ production in the supernatant was measured by ELISA. (B) Stimulation by the shortest peptide recognized by clone 37. Conditions were as in A.
Dendritic Cells Incubated with Cell Lysates Present the MAGE-3 Antigen. We tested dendritic cells, incubated with decreasing concentrations of MAGE-3 protein, for their ability to stimulate clone 37 (Fig. 4A). Half-maximum production of IFN-γ was obtained when $10^4$ dendritic cells were preincubated in a volume of 100 μl with 30 ng of the protein, which is approximately the amount present in $2 \times 10^4$ MZ2-MEL.43 tumor cells (Godelaine, D., personal communication). This suggested that a dendritic cell, having endocytosed debris of just one or a few tumor cells expressing MAGE-3, would be capable of stimulating anti-MAGE-3 CD4+ T cells. To test further the ability of dendritic cells to process debris of cells expressing MAGE-3, 293-EBNA cells transiently transfected with the MAGE-3 gene were lysed and incubated for 24 h with HLA-DR13 dendritic cells. These cells stimulated clone 37 (Fig. 4B). The amount of IFN-γ released by clone 37 increased with the amount of plasmid used for the transfection.

Presentation of the Antigen by Cells Expressing the Ii MAGE-3 Construct. A melanoma cell line which expresses HLA-DR13 and MAGE-3 was unable to stimulate the release of IFN-γ by clone 37, unless it was pulsed with the MAGE-3115–130 Peptide (Fig. 5A). This suggests that the MAGE-3 protein synthesized endogenously does not reach the class II presentation pathway. It has been reported that signals within the Ii could be used to target endogenously synthesized protein to the class II antigen-processing compartments. The expression of several constructs, which encoded fusion proteins containing part of the mouse Ii followed by fragments of the OVA gene, generated OVA peptides recognized by murine CD4+ T cells on MHC class II molecules. The highest level of antigen presentation was obtained with a construct containing the sequence coding for the first 80 amino acids of the Ii (35). Accordingly, MZ2-MEL.43 cells were transduced with a retroviral construct encoding the first 80 amino acids of the human Ii fused with MAGE-3 (retro-ⅱ.MAGE-3). MZ2-MEL.43 cells transduced with retro-ⅱ.MAGE-3 stimulated a high production of IFN-γ by clone 37, indicating that the Ii.MAGE-3 protein was processed through the class II presentation pathway (Fig. 5A). HLA-DR13 EBV-B cells transduced with the same construct also stimulated clone 37 (Fig. 5B). In contrast, HLA-DR13 EBV-B cells transduced with retro-MAGE-3 alone were unable to stimulate clone 37.

Two Other CD4+ Clones Recognized a Neighboring MAGE-3 Epitope on HLA-DR13. CD4+ T cells isolated from the blood of melanoma patient 7002, who had been immunized with the MAGE-3 protein, were stimulated with autologous dendritic cells loaded with MAGE-3 protein. After three restimulations, one microculture produced IFN-γ upon stimulation with autologous EBV-B cells loaded with the protein. CD4+ clone 2 was isolated, which produced IFN-γ upon stimulation with cells that were either loaded with MAGE-3 protein or transduced with Ii.MAGE-3, the latter proving that the clone was directed against MAGE-3 and not against a contaminant of the protein batch (Fig. 6A).

Figure 4. Presentation of the MAGE-3 antigen by dendritic cells incubated with purified protein or cell lysates. (A) HLA-DR13 dendritic cells were cultured for 24 h with different concentrations of MAGE-3 protein. The cells were washed, then incubated with 2,500 cells per well of clone 37. IFN-γ production was measured after 20 h by ELISA. The results shown represent the average of triplicate cultures. (B) 293-EBNA cells (5 × 10^6 cells per well) were transfected with different doses of pCEP4-MAGE-3 mixed with Lipofectamine®. 24 h after transfection, the transfected cells were lysed by freeze–thawing. HLA-DR13 dendritic cells (10^5 cells per well) were cultured with lysates at the equivalent of 5 293-EBNA cells per dendritic cell for 24 h. The experiment was pursued as in A.

Figure 5. Recognition by clone 37 of cells expressing an Ii MAGE-3 construct. (A) MZ2-MEL.43 melanoma cells express both HLA-DR13 and MAGE-3. Cells were pulsed for 2 h with 5 μg/ml of peptide MAGE-3115–130 and washed, or transduced with a retroviral construct encoding Ii.MAGE-3. (B) MZ2-EBV cells express HLA-DR13. As a positive control for stimulating cells, we incubated them for 20 h with 20 μg/ml of protein MAGE-3. MZ2-EBV cells were transduced with a retroviral construct encoding MAGE-3 alone or Ii.MAGE-3. Tumor cells (10^6) and EBV-B cells (5 × 10^6) were distributed in microwells, and 2,500 cells of clone 37 were added. IFN-γ production was measured after 20 h of coculture by ELISA. The results shown represent the average of triplicate cultures.
When the set of overlapping MAGE-3 peptides were tested for recognition by clone 2, peptide ELVHFLLKYRAREPV (aa 115–130) scored positive (Fig. 6 B). This peptide was shown previously to be recognized by clone 37, isolated from hemochromatosis patient LB 1555. However, peptide FLLLKYRAREPVTKAE (aa 119–134) scored positive with clone 2 but was not recognized by clone 37 (Fig. 7 B). Moreover, peptide RKVAELVHFLLKYRA (aa 111–126), previously shown to be recognized by clone 37, was not recognized by clone 2. This indicated clearly that clones 2 and 37 were directed against two adjacent but distinct epitopes. Several EBV-B cell lines were tested for their ability to present peptide MAGE-3119–134 to clone 2. All and only those expressing DR13 were capable of presenting the peptide (Table II). Patient 7002 expresses the DRB1*1302 allele. We tested a large number of MAGE-3 peptides at different concentrations to identify

Table II. HLA-DR13 Cells Present Peptide MAGE-3119–134 to CD4+ Clones 2 and 22

| EBV-B cell line | Serological specificity | IFN-γ production after stimulation with peptide MAGE-3119–134 (pg/ml) |
|-----------------|-------------------------|---------------------------------------------------------------|
| 7002            | DR 1<sup>+</sup> DR 52<sup>+</sup> | >4,000                                                |
| LB 1158         | DR 1<sup>+</sup> DR 52<sup>+</sup> | 3,967                                                   |
| LB 1118         | DR 1<sup>+</sup> DR 52<sup>+</sup> | >4,000                                                |
| LB 1622         | DR 1<sup>+</sup> DR 52<sup>+</sup> | 1,250                                                   |
| O M W           | DR 1<sup>+</sup> DR 52<sup>+</sup> | 3,651                                                   |
| M Z 2           | DR 1<sup>+</sup> DR 52<sup>+</sup> | >4,000                                                |
|                  | DR 13 negative         | 2,030                                                   |
| L K T 3         | DR 1<sup>+</sup> DR 52<sup>+</sup> | 0                                                       |
| R SH            | DR 1<sup>+</sup> DR 52<sup>+</sup> | 0                                                       |
| B M 16          | DR 1<sup>+</sup> DR 52<sup>+</sup> | 7                                                       |

EBV-B cells were incubated for 2 h with 1 μg/ml of peptide MAGE-3119–134 and washed. Clones 2 and 22 were then incubated for 20 h with 5,000 peptide-pulsed cells. IFN-γ release in the supernatant was measured by ELISA. Among the DR 13<sup>+</sup> cells, some were DRB1*1301 and others were DRB1*1302.
the shortest peptide efficiently recognized by clone 2 (data not shown). This proved to be LLKYRAREPVTKAE (aa 121–134; Fig. 7).

Interestingly, peptide MAGE-3121–134 is also present in proteins MAGE-1, -2, and -6. Therefore, we loaded DR13 EBV-B cells with a MAGE-1 recombinant protein and used them to stimulate the CD4+ clones. Clone 2, but not clone 37, produced IFN-γ (data not shown).

From hemochromatosis patient LB 1158, yet another anti-MAGE-3 CD4+ clone was obtained, referred to hereafter as clone 22. Patient LB 1158 expresses the DRB1*1301 allele. Like clone 2, clone 22 was stimulated by DR13 cells presenting peptide MAGE-3119–134 (Table II) and did not recognize peptide MAGE-3111–126 (data not shown). It was also stimulated by DR13 cells loaded with the MAGE-1 protein (data not shown).

Discussion

The procedure described here seems efficient for the activation of anti-MAGE CD4+ precursors present in the blood of individuals with or without cancer, and for the resulting isolation of anti-MAGE CD4+ permanent T cell clones. We are quite confident that it will prove to be applicable to other proteins for the identification of new antigens presented by HLA class II molecules. But our approach is not without disadvantages: a large number of the CD4+ clones obtained were apparently directed against contaminants of the protein batch. Therefore, it will be preferable to use one source of antigen for stimulation of the T cells and another to test their specificity. This may be achieved using two batches of protein produced in different organisms, such as bacteria and insect cells infected with baculovirus constructs, or, as shown here, by the use of presenting cells expressing the protein of interest fused to a truncated II.

Antigens can be processed through two different pathways that lead to presentation on HLA class II molecules. The endogenous pathway channels some proteins synthesized in the cell towards endosomal compartments, where they are cleaved into peptides that then associate with class II molecules (36). In the exogenous pathway, the APC takes up protein by endocytosis. Early endosomes undergo a progressive transition to late endosomes and then to lysosomes. The late endosomes have a relatively low pH, contain lysosomal hydrolases, and are enriched in HLA-DM and MHC class II molecules, which are associated with II in nonameric complexes. Therefore, they are also called MHC class II compartments (MIIC). II is rapidly degraded but class II-associated II peptide (CLIP), a small part of it, remains bound within the groove of the MHC molecule and blocks access of other peptides until HLA-DM catalyzes its dissociation from class II molecules. This permits the subsequent loading of potentially antigenic peptides. The peptide–HLA class II complexes are then directed to the cell surface (37).

Melanocyte-specific proteins such as tyrosinase contain a lysosomal targeting sequence that enables them to follow the endogenous class II processing pathway (38). Therefore, these differentiation antigens can be recognized directly by CD4+ T cells on those melanomas that express class II molecules (39). Constitutive expression of class II occurs frequently in melanoma. In line with their cytotoxic and nuclear localizations, the MAGE proteins appear to lack the targeting sequences that would enable them to follow the endogenous class II pathway (5, 40, 41). This probably explains why the anti-MAGE-3 CD4+ clones obtained were unable to recognize an HLA-DR13 tumor cell line expressing MAGE-3.

The anti-MAGE-3 CD4+ clones could be stimulated by HLA-DR13 dendritic cells loaded with extracts of cells producing a large amount of MAGE-3 protein. However, they failed to recognize dendritic cells incubated with lysates of tumor cells expressing MAGE-3. This might be due to a lower quantity of protein MAGE-3 in these tumor cell extracts. However, this process may be more efficient in vivo, where loading of dendritic cells with apoptotic bodies from tumor cells might result in very efficient presentation of MAGE-3 epitopes (42). The notion that MAGE-3 antigens can be presented to CD4+ T cells in vivo is supported by the observation that some cancer patients produce anti-MAGE antibodies (11, 43, 44). This is probably due to destruction of some tumor cells followed by uptake of the debris by macrophages or dendritic cells.

In a recently completed clinical trial, 25 tumor-bearing HLA-A1 melanoma patients with advanced disease received 3 subcutaneous injections of a MAGE-3 peptide presented by HLA-A1 (45). Tumor regressions were observed in seven patients, and three of these were complete. No increase in anti-MAGE CTLs could be detected in the blood of these patients, including those whose tumor regressed. The regressions occurred very slowly, suggesting that they may have been caused by a weak immune response. A major limitation of such a class I peptide–based approach might be that the CTLs elicited by the peptide reach the tumor but fail to be restimulated properly at the tumor site so that CTL amplification does not occur. This could be due to the lack of help by tumor-specific CD4+ T cells. Vaccination strategies with MAGE-3 products may benefit from the induction of anti-MAGE-3 CD4+ T cells, despite the fact that tumor cells that express both class II molecules and MAGE-3 are unable to activate CD4+ T cells directly. Upon lysis of some tumor cells by a first CTL attack, tumor debris could be processed by tumor-infiltrating APCs. Activated tumor-specific CD4+ T cells located around the tumor could then be restimulated by these cells. These CD4+ T lymphocytes might then favor the activation and proliferation of the effector CD8+ T cells, provoke the maturation of dendritic cells via CD40-CD40L interactions, and mediate recruitment of additional immune cells at the tumor sites (46–48). The resulting amplification of the immune response might then lead rapidly to the complete destruction of the tumor mass. Immunization with MAGE-3 peptides presented by class I and class II molecules or with a purified protein may represent one possibility for inducing both MAGE-3–specific CD4+ and CD8+ T cells. Another possibility is the reinjection into pa-
tients of autologous APCs loaded with MAGE-3 peptides binding to both HLA class I and class II molecules. This will require the characterization of several epitopes to cope with HLA restriction. Alternatively, autologous dendritic cells could be infected with recombinant viruses encoding an II.MAGE-3 fusion protein. Expression of II.MAGE-3 in EBV-transformed B cells was shown to result not only in the presentation of class II epitopes, but also in a very efficient presentation of a class I epitope recognized by an anti-MAGE-3.A1 CTL (Corthals, J., and K. Thielemans, manuscript in preparation). Patients could also be immunized directly with recombinant viruses encoding II.MAGE-3.

A number of results obtained in mice support the importance of immunization against antigens presented by class II molecules. Direct injection of a recombinant vaccinia encoding a human papilloma virus E7 protein linked to the sorting signal of the lysosomal-associated membrane protein (LAMP-1) was very effective in inducing protective immunity against a challenge with an E7-+ class II+ tumor. In contrast, animals injected with a vaccinia coding for the normal E7 protein were not protected (49). In another mouse model, strong protection was achieved against normal E7 protein (53). In a recent study, mice infected with recombinant vaccinia expressing MAGE-3 peptides linked to human HLA class II molecules and injected with recombinant viruses encoding an Ii.MAGE-3 fusion protein. Expression of Ii.MAGE-3 in EBV-transformed B cells was shown to result not only in the presentation of class II epitopes, but also in a very efficient presentation of a class I epitope recognized by an anti-MAGE-3.A1 CTL (Corthals, J., and K. Thielemans, manuscript in preparation). Patients could also be immunized directly with recombinant viruses encoding II.MAGE-3.

A number of results obtained in mice support the importance of immunization against antigens presented by class II molecules. Direct injection of a recombinant vaccinia encoding a human papilloma virus E7 protein linked to the sorting signal of the lysosomal-associated membrane protein (LAMP-1) was very effective in inducing protective immunity against a challenge with an E7-+ class II+ tumor. In contrast, animals injected with a vaccinia coding for the normal E7 protein were not protected (49). In another mouse model, strong protection was achieved against highly aggressive tumor cells lacking MHC class II expression through a single vaccination with a tumor-specific T helper peptide encoded by the Moloney murine leukemia virus (21). In this model, the CD8+ CTLs were helped efficiently by peptide-primed tumor-specific CD4+ T cells.

A clinical trial was recently initiated with a MAGE-3 recombinant protein. In this trial as in others involving recombinant viruses harboring large MAGE sequences, it will be essential to have reliable monitoring of the anti-MAGE-3 CD4+ response. Our observations that CD4+ cells directed against a minor contaminant in the protein batch can easily be activated in vitro have implications for this monitoring. To avoid possible misinterpretation of in vitro assays and of delayed-type hypersensitivity (DTH) assays, a product produced in another organism or autologous cells expressing the antigen endogenously should be used for these assays. Another possibility, which narrows the analysis to certain epitopes, is the use of a set of relevant peptides that can be used either to select and amplify peptide-specific T cells in vitro or to label directly T cell receptors with soluble HLA tetramers presenting the relevant peptide (50, 51). Multimeric soluble MHC class II molecules, complexed with a peptide attached covalently, were recently shown to bind with appropriate specificity and affinity to mouse-specific T cells (52). This approach requires prior identification of the antigenic peptides, as described in this report. Considering that several MAGE genes share the sequence coding for one of the two MAGE-3.DR13 epitopes reported here, these peptides are relevant to 16% of Caucasian melanoma patients, as 20% of Caucasians express HLA-DR13 and ~80% of metastatic melanomas express MAGE-1, 2, -3, or -6.

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