Generation of allo-restricted peptide-specific T cells using RNA-pulsed dendritic cells
A three phase experimental procedure

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Abbreviations: APC, antigen-presenting cells; DC, dendritic cell; DLI, donor leukocyte infusion; HSCT, hematopoietic stem cell transplantation; mDC, mature DC; MFI, mean fluorescence intensity; MHC, major histocompatibility complex; TAA, tumor-associated antigen; TCR, T cell receptor; tg-TCR transgenic, TCR; TIL, tumor-infiltrating lymphocytes; PBMC, blood mononuclear cells; pMHC, peptide-MHC

The rapidly expanding field of adoptive T cell transfer for immunotherapy of cancer has revealed the important need to have sources of T cells that can effectively recognize and eliminate different types of malignant cells. The power of adoptive cell therapy to induce complete tumor remission lasting over many years has been clearly demonstrated in patients undergoing allogeneic hematopoietic stem cell transplantation (HSCT). In this setting, genetic differences between the donor and the patient enable strong T cell responses directed against minor histocompatibility determinants to eradicate malignant leukemia clones of the patient. Complete remissions often occur only after several months up to one year following donor leukocyte infusion (DLI), due to the need for specific T cells to be activated and expanded to adequate numbers in vivo. This time lag places limits on the efficacy of DLI in the treatment of acute leukemia after HSCT, since the growth of residual malignant cells often outpaces the expansion capacity of specific immune cells. Furthermore, this therapy option is not available for many patients who cannot undergo HSCT.

For immunotherapy of solid tumors, the adoptive transfer of tumor-infiltrating lymphocytes (TIL) has provided substantial clinical benefit for numerous patients, particularly those with melanoma. However the failure to obtain adequate numbers of TIL with good function hinders the treatment of many patients. Many TIL recognize self-MHC molecules that present peptides derived from self-proteins. These T cells are usually of low avidity because negative selection of high-avidity T cells recognizing self-proteins occurs in the thymus to prevent autoimmunity; however high-avidity T cells are needed to effectively eradicate tumor cells. Furthermore, the isolation and large-scale expansion of tumor-associated antigen (TAA)-specific T cells for each patient is expensive and time consuming. Thus, a new therapeutic strategy aims to rapidly imbue activated patient T cells with tumor specificity and high functional avidity by using transgenic expression of selected TCR sequences (TCR gene therapy).

Clinical studies have utilized several patient-derived transgenic TCR (tg-TCR) in phase I trials of advanced melanoma. When compared with adoptive transfer of heterogenous TIL, these initial trials suggested that clinical efficacy may require use of mini-repertoires of lymphocytes expressing different tg-TCR. Unfortunately, tg-TCR therapy was often accompanied by substantial toxicity due to cross-recognition of normal tissues, particularly when a TCR of higher avidity was used. Animal studies have also revealed that adoptive transfer of high-avidity tg-TCR specific for p53 leads to recognition of stem cells, causing hematological cell loss and death of recipient animals. Therefore, it is of critical importance to select tg-TCR for clinical application that will provide the best gain in clinical efficacy with the lowest
toxicity for self-tissues. This requires careful selection of the TAA that should serve as targets for tumor recognition, as well as careful selection of the corresponding therapeutic tg-TCR. While a number of TAA candidates have been identified over the past decades that may prove to be suitable target molecules, including tissue-restricted proteins and cancer-testis antigens, there is still a critical need to have access to more TCR that can be utilized for TCR gene therapy. In the end, rigorous selection of both the target TAA and the corresponding tg-TCR sequence will be needed to achieve optimal discrimination between tumor cells and normal tissues. However, after these parameters have been established, a tg-TCR can be widely used for large numbers of patients whose tumors express the corresponding tg-TCR ligand.

The problem of acquiring only low-avidity T cells as sources of tg-TCR, due to negative selection, can be bypassed because this selection process is limited to self-MHC molecules that are expressed in the thymus. Therefore, it is possible to isolate T cells of high avidity that recognize peptides derived from any TAA if they are presented by allogeneic MHC molecules. Previous work showed that transgenic lymphocytes that express allo-restricted peptide-specific tg-TCR can effectively eliminate tumor cells in pre-clinical models. The most common approach to obtain allo-restricted T cells uses peptide-pulsed T2 cells as antigen-presenting cells (APC). However, many T cell clones generated in this manner fail to kill tumor cells. As an alternative, activated B cells have been used following coating with allogeneic peptide-MHC (pMHC) monomers. This method takes advantage of the professional APC capacity of activated B cells, but it requires the development of many different pMHC-monomers if it is to be applied for numerous TAA. Allo-restricted peptide-specific T cells can also be obtained using peptide-loaded APC or tumor cells to stimulate T cells derived from HLA partial-mismatched individuals if suitable donor pairs can be identified. Here a major drawback is the rarity of individuals that differ by mismatched HLA allotypes for peptide presentation. Therefore all of these approaches have major limitations for generating allo-restricted TAA-specific T cells with high functional avidity.

To overcome these obstacles we developed a procedure to isolate allo-restricted peptide-specific T cells as sources of high avidity tg-TCR which uses in vitro transcribed RNA (ivt-RNA)-transfected dendritic cells (DC) as APC. This method utilizes the important capacity of mature DC (mDC) to prime naïve lymphocytes. In the experimental procedure, T cells are co-cultured with autologous mDC that have been loaded with ivt-RNA which encodes a selected TAA, combined with ivt-RNA encoding a selected HLA allele that is not carried by the donor. Both proteins are expressed by the mDC and thereby new allo-pMHC ligands are presented at the cell surface where they function to activate allo-restricted peptide-specific responses in autologous T cells.

Results

T cells are generated using a three phase experimental procedure. The generation of allo-restricted peptide-specific T cells using mDC pulsed with ivt-RNA encompasses an experimental procedure that includes three distinct phases (Fig. 1). The first phase involves the generation of mDC from peripheral blood monocytes. This phase is spread over a time period of eight days, beginning on day 0 with respect to initiation of the primary (1°) mDC-T cell co-cultures that are established on day 0. A similar procedure must also be performed to prepare fresh mDC for use in the secondary (2°) co-cultures of mDC and 1° T cells. Therefore, a new set-up is performed in an identical manner, whereby the monocytes are isolated from fresh peripheral blood samples on day 1, in order to have mDC available for harvest and use when the 2° co-cultures are established on day 7.

The second phase of the experimental procedure involves the co-culture of CD8-enriched T cells with the ivt-RNA-pulsed mDC. The 1° co-cultures are established on day 0 and use mDC that are freshly loaded with ivt-RNA via electroporation. After ivt-RNA-loading, the mDC are cultured for 10 h in DC medium supplemented with 100 ng/ml GM-CSF and 20 ng/ml IL-4. After 10 h in DC medium, the mDC are harvested, counted and added to the pre-seeded T cells. The co-culture is performed in T cell medium. CD8+ T cells are generated from freshly drawn peripheral blood samples obtained on day 0. After Ficoll density centrifugation the lymphocytes undergo CD8-enrichment through depletion of contaminating lymphocyte populations with the use of a commercial kit. After washing and counting, mDC and CD8+ T cells are co-cultured at a ratio of 1:10. We aim to stimulate 2x10⁶ T cells in the 1° co-cultures. This number can vary dependent upon the numbers of viable mDC that are available after loading of ivt-RNA. The 1° co-cultures are continued for seven days, at which time the T cells are harvested, counted and replated with freshly prepared ivt-RNA-pulsed mDC, established from freshly isolated monocytes on day 1. The primed T cells are recovered on day 7 from the 1° co-cultures, washed, counted and resuspended in fresh culture medium. A ratio of 1:10 for mDC to T cells is also used in the 2° co-cultures. The numbers of primed T cells that can be replated in 2° co-cultures is dependent upon the numbers of viable mDC that are available for restimulation.

The third phase of the experimental procedure involves an enrichment step based on MHC-multimer sorting to obtain T cells that express TCR which recognize a pMHC ligand that is newly expressed on the mDC following transfection with both ivt-RNA. The T cells recovered from the 2° co-cultures are incubated with PE-labeled MHC-multimer and APC-labeled antibody specific for CD8. The double-positive cells are gated and sorted using either MoFlo or FACS Aria instruments. The multimer+ T cells are then immediately cloned in limiting dilution cultures. The remaining T cells are cultured in bulk to establish corresponding T cell lines. Both the T cell clones and the T cell lines are restimulated non-specifically every 14 d with anti-CD3 antibody and feeder cells.

Allogeneic MHC molecules are provided to mDC by ivt-RNA. In order to stimulate allogeneic T cells, some of which are peptide-specific, the mDC must express allogeneic MHC molecules at the cell surface after transfer of ivt-RNA encoding an HLA allotype. Since mDC display high levels of endogenous MHC class I, the surface expression of transgenic allogeneic MHC must be monitored using HLA allotype-specific monoclonal antibodies to analyze cells.
by flow cytometry. For example, expression of transgenic HLA-A2 molecules can be studied on mDC prepared from HLA-A2+ donors using a primary HLA-A2-specific BB7.2 monoclonal antibody, followed by a secondary PE-conjugated antibody. As an example HLA-A2 surface expression was already detected on the mDC surface at 1.5 h following electroporation of 25 μg ivt-RNA encoding the HLA-A*0201 allele (Fig. 2A). In this case, HLA-A2 expression reached a peak at 10 h, measured as mean fluorescent intensity (MFI), and then decreased over time. Nevertheless, around 50% of the mDC were still positive for HLA-A2 at 120 h. The greatest percentage of HLA-A2+ mDC (63%) was also seen at 10 h.

Various forms of ivt-RNA can be used to transfect mDC for allogeneic MHC expression. We analyzed four sources of ivt-RNA that contained α-globin or β-globin domains in the 3’ region, introduced to increase mRNA stability. The 3’ untranslated region of both the human α-globin and β-globin-gene was linked to the HLA-A2 construct in different ways leading to α-globin stabilized and β-globin stabilized HLA-A2 ivt-RNA. These four α-globin-stabilized ivt-RNA were compared with a standard ivt-RNA that was generated according to published procedures using commercially-available kits and reagents.15,16 This ivt-RNA carried a poly-A tail but did not contain any extra globin domain. When mDC were loaded with the different sources of RNA and analyzed by flow cytometry, a similar kinetics of HLA-A2 surface expression was observed, but use of ivt-RNA carrying a globin domain to improve mRNA stability did not increase the levels of surface HLA-A2. In fact, substantially higher levels of HLA-A2 were seen in mDC expressing the standard ivt-RNA (Fig. 2B). The differences in levels of HLA-A2 surface expression imbued by the various RNA also impacted on the capacity of the mDC to stimulate cytokine secretion by a T cell clone (JB-4) that recognizes HLA-A2 as an alloantigen.17 JB-4 cells released a far greater amount of IFNγ after stimulation with standard ivt-RNA (Fig. 2C).

Co-expression of ivt-RNA encoding MHC and antigen creates new peptide-MHC ligands. New pMHC ligands of defined specificity can be created on mDC by introducing two different species of ivt-RNA, one encoding an allogeneic HLA and the second encoding
a new antigen, for example a defined TAA. To demonstrate that both proteins, encoded by the two co-transfected ivt-RNA species, are expressed in the mDC after electroporation, we used ivt-RNA encoding HLA-A2 combined with ivt-RNA encoding tyrosinase, which is expressed in many melanomas, as a model TAA. As shown above, HLA-A2 expression was easily monitored by surface staining of mDC. Tyrosinase expression was monitored by intracellular staining using tyrosinase-specific antibody since it is not expressed as a membrane protein.

In this analysis, two concentrations (24 µg and 48 µg) of ivt-RNA encoding HLA-A2 and tyrosinase were introduced simultaneously by electroporation into mDC prepared from an HLA-A2 donor (Fig. 3A). The intracellular expression of tyrosinase was analyzed after 3 h and surface expression of HLA-A2

Figure 2. Kinetics of HLA-A2 expression and stimulatory capacity of ivt-RNA-transfected mDC from an HLA-A2 donor. (A) HLA-A2 mDC are electroporated with 25 µg HLA-A2 ivt-RNA prepared by the standard method. Surface staining of HLA-A2 is performed at different time points (1.5 h, 3 h, 6 h, 10 h, 24 h, 48 h and 120 h) after transfection and analyzed by flow cytometry. Stained samples are represented by filled gray curves and corresponding controls by empty curves. Percent positive cells, mean fluorescence intensity (MFI) and x-fold expression are shown in the upper right corner of each histogram. The x-fold expression is calculated by dividing the MFI of the positive sample with the MFI of the control. (B) For comparison, mDC of an HLA-A2 donor are transfected with 25 µg of 5 different species of HLA-A2 ivt-RNA. The quality of the standard poly-A tail (pAAA)-stabilized HLA-A2 ivt-RNA is compared with α-globin-stabilized (1, 2, 3) and β-globin-stabilized HLA-A2 ivt-RNA. The expression at different time points is depicted in a bar histogram as MFI. (C) mDC transfected with the various HLA-A2 ivt-RNA species are used 24 h after electroporation for co-incubation with the HLA-A2 allo-reactive CTL clone JB4 to assess their stimulatory capacity. IFNγ is quantified in culture supernatants by ELISA and presented as pg/ml. Mean values and mean deviations are derived from duplicate measurements.
was assessed at 6 h, based on preliminary studies of each individual protein (data not shown). The percentages of HLA-A2+ mDC increased from around 30% using 24 μg ivt-RNA (Fig. 3A, panels 1 and 2) to around 60% using 48 μg ivt-RNA (Fig. 3A, panels 3 and 4). Only a small but non-significant increase in the percentages of mDC expressing intracellular tyrosinase protein was observed when the concentration of tyrosinase ivt-RNA was doubled (Fig. 3A, panels 1 and 3 vs. 2 and 4). A paradoxical decrease in HLA-A2 expression was seen when the concentration of tyrosinase ivt-RNA was increased, leading to a decrease in MFI from 101 when 24 μg of tyrosinase ivt-RNA (Fig. 3A, panel 3) was used to a MFI of only 53 when 48 μg of tyrosinase ivt-RNA was co-expressed in the mDC (Fig. 3A, panel 4). This suggested that the total amount of ivt-RNA expressed in the mDC might be

![Figure 3. Co-expression of two different ivt-RNA species encoding HLA-A2 and tyrosinase. mDC are electroporated with four different ivt-RNA combinations: (1) 24 μg HLA-A2 and 24 μg tyrosinase ivt-RNA, (2) 24 μg HLA-A2 and 48 μg tyrosinase ivt-RNA, (3) 48 μg HLA-A2 and 24 μg tyrosinase ivt-RNA and (4) 48 μg HLA-A2 and 48 μg tyrosinase ivt-RNA are mixed and co-transfected into mDC of an HLA-A2 donor. (A) Surface staining of HLA-A2 is performed at 6 h and intracellular staining of tyrosinase at 3 h after transfection and analyzed by flow cytometry. HLA-A2 expression is represented by filled curves, corresponding tyrosinase expression of the same sample is shown beneath by filled curves. Mock-control mDC are included in the respective histograms by open curves. Percent positive cells, MFI and x-fold expression are shown in the upper right corner of each histogram. (B) Stimulatory capacity of the mDC loaded with different concentrations of HLA-A2 and tyrosinase ivt-RNA is analyzed by induction of IFNγ secretion of specific T cell clones. Columns represent the amount of IFNγ (pg/ml) secreted by a tyrosinase-independent HLA-A2 allo-reactive CTL (JB4) and an HLA-A2-restricted tyrosinase peptide-specific CTL (Tyr-F8) after co-incubation with ivt-RNA-pulsed DC, 24 h after electroporation. IFNγ is quantified in culture supernatants by ELISA. Mean values and mean deviations are derived from duplicate measurements. IFNγ secretion of T cells alone and in co-culture with non-transfected mock mDC serve as negative controls. Maximal IFNγ release of the CTL is observed in co-cultures with the tumor cell line Mel-93.04A12 as the positive control.](www.landesbioscience.com/OncoImmunology/133)
inhibitory. This observation was similar to effects we observed when multiple sources of ivt-RNA that encode different TAA were introduced into mDC.

The response of a T cell clone (Tyr-F8) that is specific for a new pMHC ligand comprised of tyrosinase peptide derived from tyrosinase protein, demonstrated their independence from tyrosinase expression when co-expression was analyzed using independent sources of ivt-RNA to enable good co-expression of these protein. Clone Tyr-F8 responded to all four mDC populations, regardless of HLA-A2 and 24 ivt-RNA, demonstrating their independence from tyrosinase expression when co-expression was analyzed using independent sources of ivt-RNA to enable good co-expression of these protein.

When co-expression was analyzed using independent sources of ivt-RNA, the lower concentrations of ivt-RNA depicted in panels 1 and 2 (Fig. 3A) were generally found to stimulate weaker responses of Tyr-F8 cells than mDCs loaded with higher concentrations of ivt-RNA, as depicted in Figure 3A, panels 3 and 4. This revealed that both the percentages of HLA-A2+ mDC as well as the level of HLA-A2 expression were important. The small variations in tyrosinase protein expression seemed not to be relevant (Table 1).

Co-expression of other ivt-RNA combinations allows HLA and TAA expression on mDC. To establish that other combinations of ivt-RNA also led to strong co-expression of MHC and TAA in mDC, we analyzed melan-A and survivin as TAA, together with HLA-A2. Again, transgenic expression of HLA-A2 was monitored by surface staining and TAA protein expression was detected by intracellular staining. The amounts of ivt-RNA introduced into the mDC were also varied. With both combinations, it was also possible to detect high percentages of HLA-A2+ mDC and TAA+ mDC with substantial levels of protein expression (Fig. 4). These results demonstrated that this approach provides a robust method to efficiently generate mDC that co-express new MHC and TAA. Furthermore, the same method could be easily used to introduce ivt-RNA only encoding a TAA into mDC that naturally expressed the desired HLA restriction element due to presence of the endogenous class I allele, as shown in Table 1.

| HLA-A2 tyrosinase | Exp. | 1  | 2  | 3  | Mean |
|-------------------|------|----|----|----|------|
| Tyr-F8            | 24 µg| 24 µg| 0.0 | 1.5 | 4.1  | 1.9  |
| mDC* mock         | 24 µg| 48 µg| 3.6 | 2.2 | 4.0  | 3.3  |
| mDC 1             | 48 µg| 24 µg| 14.0| 3.6 | 9.9  | 9.2  |
| mDC 2             | 48 µg| 48 µg| 8.6 | 7.9 | 7.5  | 8.0  |
| mDC 3             | 14.0 | 14.7| 12.1| 13.6|
| mDC 4             | 7.4  | 17.5| 7.8 | 10.9|
| tumor cells       | 100.0| 100.0| 100.0| 100.0|

Table 1. Stimulatory capacity of mDC loaded with different amounts of ivt-RNA.
Figure 4. For figure legend, see page 134.
Induction of allo-restricted peptide-specific T cell lines using ivt-RNA-pulsed mDC. The preceding studies established the parameters for creating mDC that can be used as stimulating cells for the induction of allo-restricted peptide-specific T cells using CD8-enriched T cells that are prepared from the same donor as the mDC. An experiment demonstrating the application of these ivt-RNA-pulsed DC for T cell stimulation is shown in Figure 5. Here two comparisons are included: mDC were prepared from HLA-A2⁺ donors who carried an endogenous HLA-A*0201 allele and thereby received only tyrosinase-encoding ivt-RNA while the other mDC were prepared from HLA-A2⁻ donors that were pulsed with ivt-RNA encoding both tyrosinase and HLA-A2.

These DC were used to directly prime enriched, autologous CD8⁺ T cells, using two rounds of in vitro stimulation with freshly prepared mDC. Prior to activation and after the second stimulation, the frequency of CD8⁺ T cells bearing TCR specific for HLA-A2-tyrosinase₃₆₉–₃₇₇-peptide ligand was measured using anti-CD8 antibody and a tyrosinase₃₆₉–₃₇₇/HLA-A*0201-multimer.

Figure 5. De novo priming of CD8⁺ T cells with RNA-pulsed DC. mDC derived from an HLA-A2⁺ donor transfected with 24 μg tyrosinase ivt-RNA and mDC derived from an HLA-A2⁻ donor transfected with 48 μg HLA-A2 and 24 μg tyrosinase ivt-RNA are used for a de novo T cell priming. (A) A2-tyr multimer staining (x-axis) of CD8-enriched T cell populations are determined for cells obtained from an HLA-A2⁺ donor and an HLA-A2⁻ donor before initiation of primary cultures with ivt-RNA-loaded autologous mDC and one day prior to cell sorting (i.e., 6 d after the 2° stimulation). Staining for CD8 is shown on the y-axis. The percentage of multimer⁺CD8⁺ T cells is displayed in the upper right quadrant, showing an increase after mDC stimulation. (B) Sorting scheme for two T cell cultures derived from an HLA-A2⁺ (left diagram) and an HLA-A2⁻ donor (right diagram). A2-tyr multimer staining is shown on the x-axis and CD8 staining on the y-axis. The boxed gates represent the multimer⁺CD8⁺ T cell populations that are selected for fluorescence-activated cell sorting. Dead cells are excluded by propidium iodide staining (data not shown). MFI of multimer staining is given for the gated cells in the lower right corner and the percentage of gated cells of total T cells is indicated beneath the gates. (C) Before initiation of primary cultures, CD8-enriched T cells are stained simultaneously for A2-tyr multimer, CD8 and either CD45RA or CD45RO. Histogram analysis for CD45RA⁺ and CD45RO⁺ cells is performed on gated multimer⁺CD8⁺ cells of both donors. (D) Sorted T cells are expanded as bulk lines and further re-analyzed after 7–22 d of culture via A2-tyr multimer (x-axis) and CD8 (y-axis) staining. T cell lines of two HLA-A2⁺ and two HLA-A2⁻ donors are depicted. Percentages of multimer⁺CD8⁺ T cells are displayed in upper right quadrant and MFI for this population is presented in lower right corner. T cells are labeled for 45 min with the multimer and either fixed immediately with 1% paraformaldehyde (0 h) to determine MFI of multimer staining (i.e., Intensity) or 2 h after washing of cells to remove unbound multimers with addition of HLA-A2-specific antibody to prevent rebinding of dissociated multimers to assess loss of multimer binding (i.e., Off-rate). Specificity is assessed with control HLA multimers, utilizing peptides from the pp65 protein of cytomegalovirus bound either to HLA-B7 or HLA-A2 molecules (i.e., Specificity).
 Frequencies of multimer-CD8+ T cells were approximately equal (0.46% and 0.49%) in enriched CD8+ T cells before DC stimulation (Fig. 5A). Multimer-CD8+ cells that were detected prior to DC stimulation revealed a phenotype of naïve T cells since they were positive for CD45RA and negative for CD45RO surface markers19 (Fig. 5C). The multimer cell frequencies increased to 0.65% of HLA-A2- and 1.39% of HLA-A2+ primed T cells, based on total T cells at the end of the 2° co-culture period (Fig. 5A). One day later, cultures were harvested and labeled with anti-CD8-antibody and A2-tyr multimer to perform fluorescence-activated cell sorting. Multimer+ cells were gated as shown and isolated (Fig. 5B). Distinct populations of multimer+ T cells were more difficult to identify in cultures from HLA-A2- donors but were easy to delineate in cultures from HLA-A2+ donors. The double-positive cells from the HLA-A2- donor shown here had a relatively low mean fluorescence intensity (MFI) of multimer binding (MFI = 3,963). In contrast, substantially higher multimer staining intensity (MFI = 17,110) was seen on gated cells of the HLA-A2+ donor. Sorted cells were cloned in limiting dilution cultures and the remaining cells were expanded as bulk cultures using antigen-independent stimulation.20

Bulk cultures of multimer-sorted T cells were re-analyzed after 7–22 d of in vitro expansion. Results are presented from four different donors, primed and sorted according to the full experimental procedure depicted in Figure 1. As expected, the majority of cells were CD8+ in all four T cell lines (Fig. 5D). MFI of multimer binding was used as a first estimate of structural TCR-pMHC binding affinity since higher multimer staining intensity has been shown to indicate a stronger interaction of TCR with their ligands.20 Multimer+ cells were present in the cultures of HLA-A2+ donors at frequencies of 11.2% and 8.3%, respectively. These T cells had low-intermediate intensities of multimer binding (MFI = 414 and 227). Apparently, substantial numbers of non-specific T cells were gated and sorted. In contrast, 93.1% and 78.2% of sorted cells of the two HLA-A2- donors bound HLA-multimers with higher intensities (MFI = 1,176 and 1,433). We also measured loss of HLA-multimer binding over time (i.e. HLA-multimer off-rate) as a second parameter of structural TCR-pMHC binding affinity. A slower off-rate indicates that the TCR-ligand interactions are more stable and thereby of higher structural affinity.21 After initial incubation with HLA-multimer and washing, the T cells were incubated for 2 h at 4°C in the absence of multimer and the presence of antibody specific for HLA-A2 molecules to prevent cellular re-association of released multimers. The rapid loss of multimer binding in T cells of HLA-A2+ donors indicated that the self-MHC-restricted T cells were of lower TCR affinity. After 2 h only 6.2% and 1.7% of the T cells were multimer+, accounting for losses of 65% and 79%, respectively. Multimer binding of T cells from HLA-A2+ donors was more stable over time, indicating a higher structural TCR binding affinity for the same pMHC ligands. After 2 h, 68.3% and 60.3% of the cells were multimer+, accounting for losses of only 27% and 23%, respectively. Specificity of multimer binding was confirmed using two control multimers, containing peptides derived from cytomegalovirus protein pp65. Both the HLA-B7-pp65 and HLA-A2-pp65 multimers showed negligible binding to the four T cell lines (0.1–1.0%). These results revealed that bulk T cell cultures containing highly enriched numbers of multimer+ T cells could be readily generated using the detailed experimental procedure. After expansion, the T cell lines could be further enriched by multimer sorting or they can serve as a source of T cell clones that can be isolated by single cell sorting or limiting dilution cultures. We have found that it is equally feasible to establish limiting dilution cultures directly from the multimer-sorted T cells on day 14 of the experimental procedure.22

Discussion

The experimental procedure described here provides a highly flexible method to generate allo-restricted peptide-specific T cell lines and clones. We have applied this experimental approach to generate HLA-A2-alo-restricted T cell clones specific for tyrosinase, survivin,21 melan-A (unpublished observations), hyaluronan-mediated motility receptor (HMMR), Wilms’ tumor-1 (WT-1), NY-ESO among others. Two specificities of T cells arise in the priming cultures using mDC expressing new allogeneic MHC molecules and TAA. Since the responding cells and DC are derived from HLA-A2- donors, one fraction of T cell clones recognizes the transgenic HLA-A2 as an alloantigen, irrespective of specific TAA-derived peptide. The second fraction recognizes a TAA-derived peptide in association with allogeneic HLA-A2 molecules. These two types of response must be separated at the level of individual clones. We observed that about 50% of the multimer-sorted cells arising in the allogeneic co-cultures recognized HLA-A2 as an alloantigen, independent of TAA-derived peptide. Fortunately, after multimer sorting and cloning at the end of the 2° co-culture period around 30% of the T cell clones were allo-restricted and peptide-specific. As expected, none of the T cell clones derived from HLA-A2+ donors showed HLA-A2 alloreactivity. Thus, this experimental procedure provides a robust method for efficiently obtaining a variety of different T cell clones for further study.

The preparation of mDC as stimulating cells is a particularly critical component of the experimental procedure. The use of different combinations of ivt-RNA encoding MHC and TAA showed that the method is reliable for obtaining good co-expression of both proteins. In essence it can be used for any MHC allele and TAA for which cDNA are available to generate ivt-RNA. However it is important to track the expression of both the MHC and the TAA in mDC to assure adequate co-expression of the proteins. In the first instance, this can be done using flow
cytometry to detect expression of the new proteins. Ideally, detection of a new pMHC ligand through recognition of a specific T cell clone is useful to demonstrate that the functions of antigen processing and presentation are intact in the mDC. This may not always be feasible due to lack of specific T cell clones.

Because the entire TAA appears as an intracellular protein, it can be processed and presented by the natural antigen processing machinery of the mDC, thereby creating multiple new epitopes at the cell surface, often of unknown sequence. For known pMHC ligands, specific T cells can be sorted using multimers. For T cells of unknown peptide specificity, other isolation procedures can be employed, such as cytokine capture or CD137 expression (unpublished observations) after APC stimulation. Thereby, this method opens the possibility to tap T cells with high functional avidities for many new TAA specificities.

The major advantage of this experimental procedure is that responding CD8+ T cells and mDC can be used from any healthy donor who does not carry the HLA-allotype that is selected as the molecule for allo-restriction. This eliminates the need for extensive searches of HLA-typed donors to identify partial HLA-mismatched pairs that differ by only one HLA-allotype. Because an extensive bank of cDNA for different HLA alleles is available, or they can be quickly isolated from any donor of choice, studies of TAA presentation by HLA molecules other than HLA-A2 are now much easier to perform. Furthermore, it is possible to extend the studies to TAA presentation by MHC class II molecules using the same experimental procedure, with the modification that the TAA-encoding ivt-RNA should be modified to target the protein to the class II presentation pathway.

Materials and Methods

Culture of cell lines and effector T cell clones. The human melanoma cell line Mel-93.04A12 (HLA-A2+, tyrosinase+ melan-A+; gift of P. Schrier, Department of Immunohematology, Leiden University Hospital, The Netherlands) is cultured in RPMI 1640 medium (Invitrogen, 318725) supplemented with 12% fetal bovine serum (FBS) (Invitrogen, 10091148), 2 mM L-glutamine (Invitrogen, 25030-024) and 1 mM sodium-pyruvate (Invitrogen, 11360039) and non-essential amino acids (Invitrogen, 11140035).

For pre-testing the stimulatory capacity of ivt-RNA-pulsed mDC, the HLA-A2 allo-specific CTL J417 and the HLA-A*0201-restricted tyrosinase360–377 peptide-specific CTL TyrF8 (gift of P. Schrier, Department of Clinical Oncology, Leiden University Hospital, The Netherlands) are cultured as described. Generation of DC. Blood samples from healthy donors are collected after informed consent, in accordance with the Declaration of Helsinki and approval of the Institutional Review Board of the University Hospital of the Ludwig-Maximilians-University, Munich, Germany. Peripheral blood mononuclear cells (PBMC) are isolated by Ficoll density gradient centrifugation. PBMC are resuspended in 50 μl VLE (very low endotoxin) RPMI 1640 medium (Biochrom, FG1415) supplemented with 1.5% human serum (DC medium) at 7.5x10⁶ cells per 7 cm² culture flask (NUNC, 178905) and incubated at 37°C and 5% CO₂ for 1 h. Non-adherent cells are carefully removed by washing. Adherent monocytes are cultured in medium containing 100 ng/ml GM-CSF (Leukine® by Berlex, NDC50419-050-30) and 20 ng/ml interleukin-4 (R&D Systems, 104-IL-050-CF) and fed with the same medium on days 3 and 6. On day 6 of culture, the immature DC are differentiated into mDC by addition of medium containing 10 ng/ml IL-1β (R&D Systems, 201-LB-025-CF), 15 ng/ml IL-6 (R&D Systems, 206-IL-050-CF), 10 ng/ml TNFα (R&D Systems, 210-TA-050-CF) and 1 μg/ml PGE₂ (Sigma-Aldrich, p5640-10MG) for 2 d.

Production of HLA-A2, tyrosinase, melan-A and survivin ivt-RNA. The following linearized plasmids were used as in vitro transcription templates to produce single-species ivt-RNA with the aid of the mMESSAGE mMACHINE T7 kit and Poly(A) tailing kit (both Ambion, AM1344 and AM1350), according to the manufacturer’s instructions: pCDM8-HLA-A2 plasmid with HLA-A*0201 cDNA (gift of E. Weiß, Department Biology II, Ludwig-Maximilian-University, Munich, Germany), pZeoSV2+ huTyr with tyrosinase cDNA (gift of I. Dreher, Institute of Molecular Virology, Helmholtz Zentrum München, Munich, Germany), pcDNA1/Amp/Aa1 with melan-A cDNA (gift of T. Wölfel, Third Department of Medicine, Hematology and Oncology, Johannes-Gutenberg University of Mainz, Mainz, Germany) and pGEM4Z/survivin/A64 plasmid (cloned by S. Milosevic, Institute of Molecular Immunology, Helmholtz Zentrum München).

Also different modified mRNA molecules (RNActive®, CureVac) were used that encode HLA-A2 molecules. Four different HLA-A2 mRNA are compared, here whereby the 3’ untranslated region of either the human α-globin or β-globin gene is linked to the HLA-A2 construct in different ways, leading to α-globin or β-globin stabilized HLA-A2 mRNA.

Electroporation of DC. On day 8, mDC are harvested 48 h after addition of maturation cytokines. Electroporation is performed as previously described. In brief, 2 to 3x10⁶ cells are resuspended in approximately 200–300 μl OptiMEM I medium (Invitrogen, 31985047), placed in a 0.4-cm electroporation cuvette (Bio-Rad, 5000835) and incubated for 3 min on ice. Electroporation is performed with the Gene Pulser Xcell (Bio-Rad) using the exponential protocol at 250 V and 150 μF. RNA species are transfected using different amounts of individual RNA, between 24–50 μg per electroporation.

FACS analysis. HLA-A2 molecules are stained on the surface of DC with the Biocytin (PE) goat anti-mouse IgG (Jackson ImmunoResearch, 115-116-146) for 45 min on ice followed by a secondary antibody conjugated with phycoerythrin (PE). The intracellular protein expression in ivt-RNA-transfected mDC is detected using the following primary antibodies: tyrosinase-specific primary monoclonal antibody (clone T311; diluted 1:10; Novocastra Laboratories Ltd., 600741), melan-A-specific primary monoclonal antibody (clone A103; diluted 1:20; DakoCytomation, M7196) and survivin-specific primary monoclonal antibody (clone 91630; diluted 1:2.5; R&D Systems, MAB886) followed by staining with Cy5-conjugated secondary antibody (rat anti-mouse IgG; Jackson ImmunoResearch, 415-176-166). The intracellular staining is performed as follows. The mDC samples are first fixed in FACS buffer containing 1% paraformaldehyde (Sigma,
mDC serve as negative controls and show almost no IFN-γ release assay. Stimulatory capacity of RNA-pulsed mDC is determined by the mean fluorescence of untransfected mDC. mDC are treated in the same manner and used as negative controls. Data acquisition and analysis are done on a FACSCalibur (BD Biosciences) using CellQuest Pro software or FloJo software (Tree Star). X-fold expression is calculated by dividing the mean fluorescence of ivt-RNA-transfected mDC by the mean fluorescence of untransfected mDC.

To determine the phenotype of unprimed multimer CD8+ T cells, cells are double-stained with PE-labeled HLA-A2 tyrosinase-369-377 (YMCDTQMSV) (A2-tyr) multimer,26 CD8-specific antibody (clone RPA-T8, BD PharMingen, 555369) and simultaneously with either CD45RO- (clone UCHL1, Immunotech, IM1247) or CD45RA-specific (clone H1100, BD PharMingen, 550855) antibody. Histogram analysis for CD45RO and CD45RA staining is performed on gated multimer+CD8+ T cells. Data acquisition and analysis are made with an LSRII instrument (BD Biosciences) using FloJo software (Tree Star).

**IFNγ release assay.** Stimulatory capacity of RNA-pulsed mDC is analyzed 24 h after electroporation. JB4 or Tyr-F8 CTL suspensions (2 × 10^6 cells in 100 μl) are added to the RNA-loaded mDC (4 × 10^5 cells in 100 μl) in round-bottom 96-well plates. T cells without stimulator cells and with mock-transfected mDC serve as negative controls and show almost no IFNγ secretion. T cells are co-cultured with the HLA-A2+ tyrosinase+, melan-A+ tumor cell line Mel-93.04A12, as positive control. Culture supernatants are harvested after 24 h co-culture and assessed by a standard ELISA using the OptEIA™ Human IFNγ Set (BD Biosciences, 555142). Data represent mean values with corresponding mean deviations calculated from duplicate determinations.

**De novo priming of T cells with ivt-RNA-pulsed DC.** mDC are transfected with pre-selected optimal ivt-RNA concentrations via electroporation as described above. mDC derived from HLA-A2+ donors are loaded with 24 μg tyrosinase ivt-RNA and mDC prepared from HLA-A2- donors are co-transfected with 24 μg tyrosinase and 48 μg HLA-A2 ivt-RNA. Untouched autologous CD8+ T lymphocytes are enriched from PBMC via negative selection using a commercial kit, according to the manufacturer’s instructions (CD8+ T Cell Isolation Kit, Miltenyi Biotec, 130-094-156) on the same day. Co-cultures are initiated 10 h after electroporation in 24-well plates (TPP, Z707805-64EA). 1 × 10^5 ivt-RNA-pulsed mDC are added to 1 × 10^6 CD8+ T cells, yielding a mDC:T cell ratio of 1:10 in T cell medium consisting of RPMI 1640 (Invitrogen, 318725), supplemented with 10% heat inactivated human serum, 4 mM L-glutamine (Invitrogen, 25030-024), 12.5 mM HEPES (Invitrogen, -15630049), 50 μM β-mercaptoethanol (Sigma, -63700-50ML-F) and 100 U/ml penicillin/streptomycin (Invitrogen, -15070063). On day 0, 5 ng/ml IL-7 (Promokine, -C-61712) is added. Addition of IL-2 (Prol leukin® S, Novartis) is delayed for 2 d to decrease proliferation of non-specific CD8+ T cells,19 then it is added every 3rd day. The 2nd stimulation of the primed T cells is made seven days later using freshly generated ivt-RNA-pulsed mDC, prepared as above.

**HLA-multimer staining and sorting.** Prior to priming and six days after the 2nd stimulation of CD8-enriched T cells with ivt-RNA-pulsed mDC, HLA-A2-restricted tyrosinase-specific T cells are detected by staining with PE-labeled HLA-A2 tyrosinase-369-377 (YMCDTQMSV) (A2-tyr) multimer,26 CD8-specific antibody (clone RPA-T8, BD PharMingen) and propidium iodide (PI; 2 μg/ml). Up to 1 × 10^6 cells are incubated in 50 μl volume for 25 min with 4 μg PE-labeled multimer on ice in the dark. For sorting, multiple samples of up to 5 × 10^6 cells are incubated with 12 μg multimer in 100 μl PBS + 0.5% human serum. APC-labeled, CD8-specific antibody is then added for an additional 25 min. After staining, an aliquot of cells is washed and fixed for analysis by flow cytometry using a FACSCalibur (BD Biosciences) and the remaining cells are pooled and sorted on a FACS Aria instrument (BD Biosciences) as described.27

To observe HLA-multimer dissociation (off-rate), cells are washed after multimer binding and resuspended in FACS buffer containing saturating amounts of monoclonal HLA-A2-specific BB7.2 antibody to capture detached multimers and prevent rebinding to T cells. After 2 h, samples are fixed and analyzed by flow cytometry.21 PE-labeled HLA-B7/pp65417–427 (TPRVGGGAM) peptide/human β2m multimers serve as the HLA control and HLA-A*0201/pp65495–503 (NLVPVMVAT) peptide/human β2m multimers serve as a peptide-specificity control. Peptides for these multimers are derived from the pp65 protein of human cytomegalovirus.

**Culture of peptide-specific T cell lines.** Multimer-sorted T cells are expanded as bulk T cell lines in 96-well round-bottom plates (TPP, 500077) in 200 μl/well T cell medium (600 cells per well). 50 IU/ml IL-2 is supplemented every three days with 5 ng/ml IL-7 (Promokine, C-61712) and 10 ng/ml IL-15 (Peprotech Inc., AF-200-15) every seven days. Every 14 d T cell lines are restimulated non-specifically with anti-CD3 antibody (0.1 μg/ml) (OKT-3; gift of E. Kremmer, Institute of Molecular Immunology, Helmholtz Zentrum München) and provided with 1 × 10^5 feeder cells per 96-well (TPP, 92097), consisting of irradiated (50 Gy) PBMC derived from a pool of five unrelated donors. Proliferating T cells are transferred into 24-well plates (TPP, Z707805-64EA) and cultured in 1.5 ml T cell medium plus cytokines. 1 × 10^6 allogeneic irradiated PBMC are added per well as feeder cells in 24-well plates.

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Declaration of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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