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METHODS: ORIGINAL ARTICLE

Mass Spectrometry Reveals Changes in MHC I Antigen Presentation After Lentivector Expression of a Gene Regulation System

Roland Vogel¹, Reem Al-Daccak², Oliver Drews³, Jessy Alonzeau¹, Gabor Mester³, Dominique Charron², Stefan Stevanovic³ and Jacques Mallet¹

The rapamycin-inducible gene regulation system was designed to minimize immune reactions in man and may thus be suited for gene therapy. We assessed whether this system indeed induces no immune responses. The protein components of the regulation system were produced in the human cell lines HEK 293T, D407, and HER 911 following lentiviral transfer of the corresponding genes. Stable cell lines were established, and the peptides presented by major histocompatibility complex class I (MHC I) molecules on transduced and wild-type (wt) cells were compared by differential mass spectrometry. In all cell lines examined, expression of the transgenes resulted in prominent changes in the repertoire of MHC I-presented self-peptides. No MHC I ligands originating from the transgenic proteins were detected. *In vitro* analysis of immunogenicity revealed that transduced D407 cells displayed slightly higher capacity than wt controls to promote proliferation of cytotoxic T cells. These results indicate that therapeutic manipulations within the genome of target cells may affect pathways involved in the processing of peptide antigens and their presentation by MHC I. This makes the genomic modifications visible to the immune system which may recognize these events and respond. Ultimately, the findings call attention to a possible immune risk.

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Subject Category: Methods section

Introduction

The use of gene transfer as a therapeutic tool requires a regulatory system allowing control of the expression of the therapeutic gene by the administration of a small inducer molecule. The treatment could then be adapted to the needs of the patients and, should complications arise, the therapy could be stopped or interrupted. Several regulatory systems have been developed.¹ Tetracycline-dependent regulatory systems² perform very well in transgenic animals.³ However, the transactivators used include peptide sequences from *Escherichia coli* and *Herpes simplex* and may therefore provoke immune reactions in patients. Immune responses against the transactivator of a recent version of the tetracycline-dependent regulatory system were observed after expression in the muscles of nonhuman primates.⁴ Moreover, an immunodominant HLA-A*0201 epitope was detected in the reverse tetracycline-dependent transactivator. This epitope caused cytolytic responses and compromised transgene expression under the control of the tetracycline-on system.⁵ As a consequence, adverse immunity may interfere, in this case, with gene transfer protocols and prevent gene therapy achieving its aims.

Recently, two novel regulatory systems have been developed which are induced by nonimmunosuppressive derivatives of rapamycin. The first system interferes with transcription and exploits the inducer-dependent interaction between Frap

kinase and the Frap kinase binding protein for the reversible *in situ* assembly of a functional transcription factor which activates transcription from a minimal promoter.⁶ The second system interferes with the secretory pathway and is adapted to controlling the production of secreted therapeutic factors. It exploits the ability of the inducer to control, in the endoplasmic reticulum, aggregation of a mutated Frap kinase binding protein fusion protein harboring the secreted polypeptide.⁷ Rapamycin-inducible transcription allows very tightly regulated expression of transgenes.⁸ Inducer-dependent secretion may be used in combination with inducible transcription for optimized control of the production of therapeutic factors such as the glia derived neurotrophic factor (GDNF).⁹

The key advantage of rapamycin-inducible systems is that they involve fusion proteins of human origin and consequently immune reactions in humans are minimized. They are therefore expected to be safe. However, the protein components include short peptide sequences that link the various peptide domains. These sequences may themselves constitute novel epitopes or may affect the proteasome cleavage pattern thereby generating novel peptide antigens from the fusion proteins. Any such novel antigens may be presented by the major histocompatibility complex (MHC). This possibility can readily be assessed by application of algorithms for the prediction of proteasome cleavage¹⁰ and MHC class I (MHC I) ligand motifs.¹¹ MHC I ligands may also emerge after

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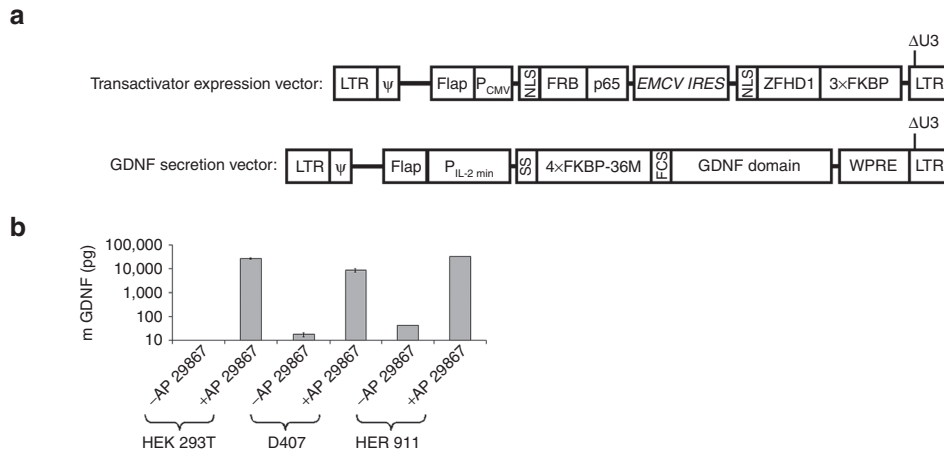


Figure 1 Rapamycin-inducible production and secretion of GDNF. (a) Lentiviral constructs: The transactivator expression vector (TEV) mediates constitutive expression of the fusion proteins NLS-FRB-p65 and NLS-ZFHD1-3xFKBP allowing inducible transcription from an engineered minimal interleukin 2 promoter ($P_{IL-2 min}$). $P_{IL-2 min}$ is used in the glia derived neurotrophic factor (GDNF) expression vector to produce the fusion protein SS-4xFKBP-36M-FCS-GDNF providing rapamycin-regulated secretion of GDNF.⁹ LTR, Ψ, and Flap sequences are derived from HIV-1 (the long terminal repeats, the packaging sequence, and the central Flap element, respectively). P_{CMV} is the CMV promoter, WPRE the Woodchuck hepatitis virus responsive element, and EMCV IRES an internal ribosome entry sequence from the encephalomyocarditis virus. Under the conditions used for cell amplification (*i.e.*, in the absence of inducer) only the fusion proteins NLS-FRB-p65 and NLS-ZFHD1-3xFKBP were produced from TEV. **(b)** Cell clones derived from HEK 293T, D407, and HER 911 cells displaying inducible production of GDNF. Cells (5×10^4) were cultivated in the presence and absence of 10 nmol/l of the rapamycin derivative AP 21967. Values (\pm SE, $n = 3$) indicate the amounts of GDNF in the medium after 2 days of culture.

production of the fusion proteins due to the changes in phenotype which may result from transactivatory effects on transcription and/or competitive effects on translation and protein degradation. Moreover, the transgenic protein components of the regulatory systems may directly interfere with pathways involved in antigen processing and thereby modulate the presentation of peptide antigens.

The immunogenic potential of rapamycin-inducible transcription is of particular importance because of the diverse possible clinical applications of small molecule-inducible gene regulation. We addressed this issue in transduced human cell lines using a mass spectrometry protocol allowing differential analysis of MHC I peptide ligands after stable isotope labeling.¹² We compared the presentation of MHC I peptides by cells expressing the protein components required for rapamycin-regulated transcription with the antigen presentation on the respective wild-type (wt) control cells. Production of the transgenic proteins was associated with major changes in the presentation of antigens by MHC I in all cell lines analyzed. Allogeneic *in vitro* immunogenicity assays provide first evidence that these changes may affect immune tolerance towards transduced cells, though in an individual manner.

Results

The human cell lines HEK 293T, D407,¹³ and HER 911¹⁴ were treated with lentiviral vectors mediating rapamycin-inducible production and secretion of GDNF as described previously⁹ (Figure 1a). Selected transduced clones displaying rapamycin-inducible production and secretion of GDNF (Figure 1b), and the corresponding wt cell lines were then characterized for their HLA genotype and their expression of immune relevant molecules. HEK 293T cells were *HLA-A*02/02*, *HLA-B*07/07*,

*HLA-C*07/07*, and *HLA-DRB1*15/15*, *HLA-DQB1*06/06*; D407 cells were *HLA-A*68:02/68:02*, *HLA-B*15:03/15:03* (B72), *HLA-C*12*, and *HLA-DRB1*01:02/01:02*, *HLA-DQB1*05/05*; and HER 911 cells were *HLA-A*02/24:03*, *HLA-B*37/51*, *HLA-C*06/15*, and *HLA-DRB1*10:01/13:01*, *HLA-DQB1*05/06*. Immune phenotyping revealed that all cell lines expressed MHC I, and low levels of MHC II molecules, but were negative for the costimulatory molecules CD80 (B7.1) and CD86 (B7.2) (Supplementary Figure S1).

To focus on the effect of the constitutive expression of the transgenic fusion proteins mediating regulation of transcription, both transduced and wt cells of each cell line were amplified in the absence of inducers of the production and secretion of GDNF using parallel cultures maintained under identical rigorously controlled conditions. MHC I molecules were isolated from transduced and wt cells, and the peptides presented were extracted. To distinguish glutamine and lysine residues in mass spectrometry and to ensure specificity of the subsequent N-terminal labeling step, the ϵ -amino groups of lysine residues were modified by guanidination. A source-dependent difference in neutral mass of $m = 4.03$ was obtained by N-terminal nicotinylation using a ²H and ¹H nicotinoyloxy-succinimide reagent, maintaining the same physico-chemical properties for identical peptide sequences. Aliquots of labeled peptides from both cell populations were combined and analyzed by liquid chromatography linked to electrospray ionization mass spectrometry (LC-ESI-MS). The spectra included double signals (with peak constituents displaying a difference in neutral mass of $m = 4.03$) and single signals (Figure 2a,b) representing peptides presented in both and only one of the two specimens, respectively.¹² Single and double signals were counted for each comparative analysis (Table 1). Single signals revealing the appearance

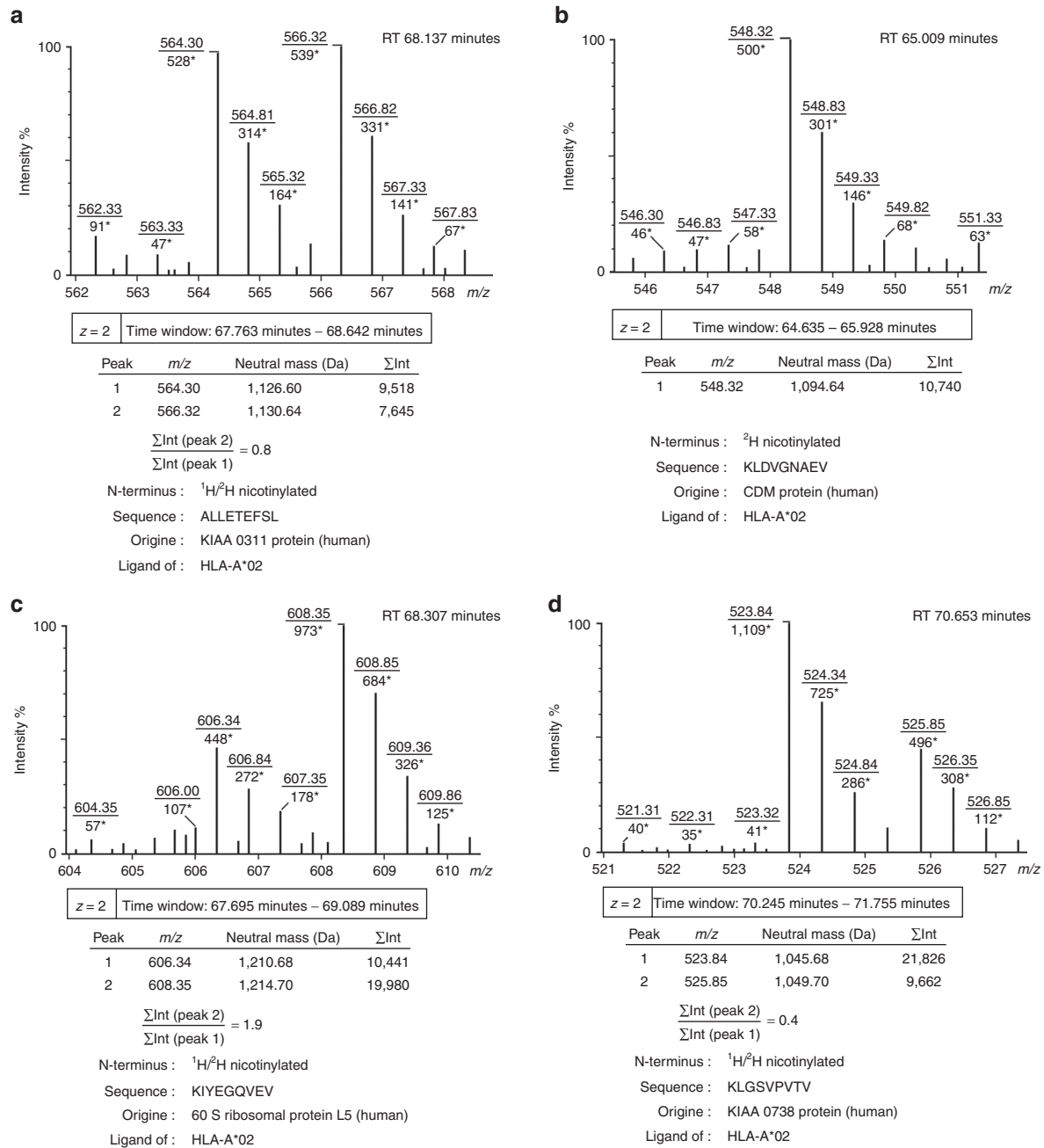


Figure 2 Typical signals obtained by LC-ESI-MS analysis of combined aliquots of ^2H and ^1H nicotinylated MHC I peptides from transduced and wild-type cells, respectively. (a) Double signal revealing presentation of approximately equal quantities of the peptide antigen on transduced and wild-type cells. (b) Single signal representing a peptide antigen only presented on transduced cells as assessed by analysis of corresponding LC-ESI-MS/MS data. (c) Double signal revealing upregulation of the peptide antigen on transduced cells. (d) Double signal revealing downregulation of the peptide antigen on transduced cells. Within the spectra, m/z values are underlined, and intensity values are marked with asterisks. RT is the chromatographic retention time, and $\sum \text{Int}$ are intensity values integrated over the given time window. Sequence information was obtained by analysis of corresponding LC-ESI-MS/MS data. All examples were taken from the comparative analysis of transduced and wild-type HEK 293T cells.

or disappearance of particular MHC I peptides in association with the expression of the transgenic proteins accounted for 40% of all signals for HEK 293T, 21% for D407, and 43% for HER 911 cells. In addition, integration of the peak intensities over time revealed differences in intensity between the peak constituents of double signals (Figure 2a,c,d). These variations indicated two- to threefold up- and downregulation

Table 1 Differential analysis of MHC I peptides on transduced and wild-type cells: Quantification of total signals, single signals, and double signals

Cell line examined	Single signals	Double signals	Total signals
HEK 293T	95	144	239
D407	82	313	395
HER 911	99	130	229

Table 2 MHC I peptide ligands presented at the surface of HEK 293T, D407, and HER 911 cells after expression of the protein components required for rapamycin-regulated transcription

Peptide sequence	Origine	NCBI accession number	Residues	Ligand of HLA allele
a) HEK 293T cells				
HPHSHDRIF	ICMT protein	Q7Z750	100–108	B*07:02
RLGADVCAV	Phosphomevalonate kinase	Q15126	31–39	A*02:01
RVGCASAPPL	cDNA	DA120873	11–20	A*02:01
APAPPKAEA	WUGSC:H_DJ0855D21.2 protein	O95014	43–51	B*07:02
RPQPGRENF	Transgelin-2	P37802	43–51	B*07:02
AARPATSTL	Translation initiation factor eIF-4 γ	A44453	914–922	B*07:02
APRAPSQVV	FUN 14 domain containing protein	NP_076423	5–13	B*07:02
APTGSQKTL	ATP-dependent RNA helicase DDX51	Q8N8A6	256–264	B*07:02
SLDQPTQTV	EIF3S8 protein	Q9BW98	246–254	A*02:01
NPSENRSLL	Angiomotin p130 isoform	AY987378	27–35	B*07:02
FVSCLGIV	IL23R protein	Q8IW84	192–199	A*02:01
KLDVGNAEV	CDM protein	S44279	167–175	A*02:01
GLATDVQTV	Proteasome b3 subunit	NP_002786	55–63	A*02:01
ILPPWPPTVL	cDNA	DW454284	28–37	A*02:01
b) D407 cells				
HSLGFEQLSL	Vesicular acetylcholine transporter	NP_003046	444–452	B*15:03 (B72)
HARPDYQIF	Putative protein hCG2038847	EAX06302	95–103	B*15:03 (B72)
ETFHDIAQV	Sequence 53 from patent WO 0166748	CAC88622	137–145	A*68:02
STDRVMTV	cDNA	BI907081	199–206	A*68:02
SRSEYAMM	cDNA	BG945501	10–17	B*15:03 (B72)
ISTPVIRTF	Oxidative stress ass. SRC activator	NP_055427	989–997	B*15:03 (B72)
AATQIRLAL	Ribosomal protein S6 kinase like protein	NP_113652	53–61	B*15:03 (B72)
EVFDKTYQF	C6orf 153 protein	AAH06293	132–140	B*15:03 (B72)/A*68:02
MALLCGLGQV	Transcriptional corepressor Cor1 1	NP_001026977	1–10	A*68:02
MASVLEQLNV	KIAA 0664 protein	BC004266	242–251	A*68:02
ETSPVLQKL	Ribonuclease III	AAF80558	1099–1107	A*68:02
TTFVGIVPLA	Transmembrane protein 132B	NP_443139	416–425	A*68:02
c) HER 911 cells				
DVANKIGII	Ribosomal protein L23a	AAH58041	148–156	B*51:01 ¹⁵
YVLFVARV	Sequence 2053 from patent EP1270724	CAD70044	1–8	A*02:01
FYQDSVGVV	MEST protein	Q5EB52	58–66	A*24:03
RFQSSAVMAL	Histone H3	AAH69079	84–93	A*24:03
LFLGNLAFI	Olfactory receptor OR 15-5	Q6IEY3	61–69	A*24:03
LLPPDLASPL	Probable DNA repair protein (XRCC9)	AAC07981	328–338	A*02:01
MRYVASYL	Acidic ribosomal protein p2	AAH05354	1–8	C*06:02
KYPHAAHIH	cDNA	BM920823	380–388	A*24:03

All sequences are from human origin.

of various MHC I peptides and provided evidence of more subtle changes in the presentation of MHC I antigens associated with the expression of the protein components mediating rapamycin-inducible transcription.

To obtain information about the sequences of peptides presented at the cellular surface by MHC I, aliquots of labeled peptides from transduced and wt cell populations were analyzed individually by liquid chromatography linked to electrospray ionization tandem mass spectrometry (LC-ESI-MS/MS). The results were first screened for peak constituents of double signals on the basis of chromatographic retention time and total mass data from LC-ESI-MS analysis. As could be expected from the labeling protocol (see Materials and

Methods section), the peak constituents that corresponded to the lower masses within double signals were from wt cells, whereas peak constituents corresponding to the higher masses were from transduced cells. For a significant number of double signals, MS/MS fragment spectra of both peak constituents were analyzed and gave identical peptide sequences. This confirms that the double signals represent peptides presented at the surface of both transduced and wt cells.

We next identified MHC I peptides that appeared or disappeared after expression of the transgenic proteins. Information on chromatographic retention time and total mass was again used to screen the LC-ESI-MS/MS data for the single signals detected by mass spectrometry in combined aliquots

Table 3 MHC I peptide ligands disappearing from the surface of HEK 293T, D407, and HER 911 cells after expression of the protein components required for rapamycin-regulated transcription

Peptide sequence	Origin	NCBI accession number	Residues	Ligand of HLA allele
a) HEK 293T cells				
RMLPHAPGV	Histone deacetylase 1	NP_004955	371–379	A*02:01
GPRTAALGLL	Reticulocalbin 2	EAW99216	58–67	B*07:02
TLVHYLAGI	cDNA	BF796701	143–151	A*02:01
RLWPKIQGL	FLJ 20500 protein	Q9NX09	184–192	A*02:01
LLLQPPAFL	Cell surface receptor FDFACT2 precursor	CAC19193	11–19	A*02:01
RLDELGGVYL	Ribophorin II precursor	B26168	190–199	A*02:01
ILTDITKGV	Elongation factor 2	P13639	660–668	A*02:01
KIADFGWSV	Aurora kinase C isoform 3	Q6AZY8	147–155	A*02:01
AMSSKFFLV	cDNA	DA555895	161–169	A*02:01
LLLPRLPLL	Sequence 78 from patent EP 1104808	AX969275	13–21	A*02:01
SLAQYLINV	hn RNP protein E2	S42471	345–353	A*02:01
b) D407 cells				
WAHQGQRY	cDNA	AW401591	35–42	B*15:03 (B72)
GSHSMRYF	MHC I antigen	AAM95704	9–16	B*15:03 (B72)
TASPVKAV	Hypothetical protein FLJ 22028	Q9H6P1	42–50	A*68:02
ETAAFIERL	Hypothetical protein	CAH10658	58–66	A*68:02
c) HER 911 cells				
SVHKGFAFV	Heterogen. nucl. ribonucleoprotein C	NP_001070910.1	47–55	A*02:01
DAHIYLNHI	Thymidylate synthase	Q8WYK4	171–177	B*51:01 ¹⁶
DAFRVNI	Assembly protein 50	G02088	30–37	B*51:01
VYPGDPLRF	tRNA splicing endonuclease 34 homolog	AAH00944	246–254	A*24:03

All sequences are from human origin.

of labeled peptides from transduced and wt cells. The single signals were detected exclusively in either the transduced or the wt cell samples. Signal intensities were sufficiently robust to allow sequencing of a significant number of peptides, and the masses of the respective b1-fragments (representing the ²H or ¹H nicotinylated N-terminal amino acid residues) provided controls confirming the presence of the respective peptides in the transduced or wt cell samples. MHC I peptides that appeared or disappeared after expression of the transgenic proteins are listed in **Tables 2** and **3**, respectively. The sequences correspond to structural motifs required for binding to MHC I molecules. Epitope prediction using the Net-MHCpan algorithm¹¹ indicated, in HEK 293T cells, the presence of peptides binding to MHC I molecules derived from the HLA alleles A*02:01 and B*07:02. In D407 cells, peptides were observed that bind to MHC I molecules from the HLA alleles A*68:02 and B*15:03. In HER 911 cells, peptides were found that bind to MHC I derived from the HLA alleles A*02:01, A*24:03, B*51:01, and C*06:02. Two of the peptides – DVANKIGII originating from the ribosomal protein L23a and DAHIYLNHI originating from thymidylate synthase – have previously been described as ligands of MHC I derived from the HLA allele B*51:01.^{15,16} The peptides identified were derived from a wide variety of cellular proteins, *i.e.*, self-proteins. No HLA peptide antigens originating from the expressed transgenic proteins were detected by the analytical method used.

In one-way allogeneic mixed lymphocyte reactions, we then investigated the capacity of transduced and wt HEK 293T,

HER 911, and D407 cells to elicit an allogeneic response from CD3⁺CD4⁻ T cells which mainly comprise cytotoxic CD8⁺ T cells. The expression of CD25 was used as a marker of activation and CFSE labeling was used to assess proliferation. Transduced D407 cells activated more strongly (1.7-fold) CD3⁺CD4⁻ T cells than wt D407 cells (**Figure 3a**), and this significantly increased activation was followed by a significantly increased proliferation of CD3⁺CD4⁻ T cells (**Figure 3b**). No significant differences compared with the respective wt controls were observed for HEK 293T and HER 911 cells suggesting that immunogenicity is limited to individual cell clones.

Discussion

The regulatory system allowing rapamycin-inducible transcription is promising for clinical applications in part, because it is composed of fusion proteins comprising polypeptide sequences that all originate from human proteins thereby minimizing the risk of immune reactions in patients. However, the sequences at the junctions between the various domains could create sites that are potentially immunogenic. Therefore, we assessed the potential immunogenicity associated with rapamycin-inducible transcription.

We assessed changes in the presentation of MHC I peptides associated with the production of the fusion proteins allowing rapamycin-inducible expression of transgenes and observed variations in the presentation of peptide antigens. A significant number of peptides appeared and disappeared,

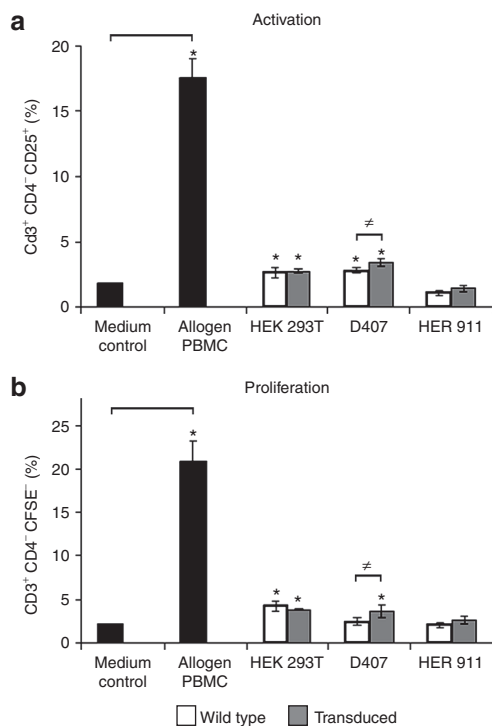


Figure 3 Allogeneic immune responses elicited by wild-type and transduced cells. CSFE-labeled peripheral blood mononuclear cells (PBMCs) were incubated with medium alone, or cocultured with control allogeneic PBMCs, wild-type or transduced cells. (a) Activation of CD3⁺CD4⁺ T cells as determined by the expression of CD25. (b) Proliferation of CD3⁺CD4⁺ T cells as evaluated by determining the percentages of dividing T cells. Results are expressed as mean values \pm SD for three independent experiments each performed in triplicate at different times of culture ($*P < 0.05$). Note: Control allogeneic PBMCs triggered large responses both in terms of activation and proliferation, as expected (positive control).

and the amounts of some peptides were affected. Surprisingly, all these peptides were derived from self-proteins. We detected no peptide antigens originating from the two transgenic proteins allowing inducible transcription of a therapeutic transgene. Using an *in vitro* immunogenicity assay, we demonstrated that the expression of these transgenic proteins and the associated changes in the repertoire of MHC I-presented peptide antigens could affect the immunogenicity of the cells.

The mass spectrometry protocol used was developed for direct and quantitative comparison of MHC-presented peptides in pairs of samples.¹² In contrast to approaches addressing changes in the transcriptome and the proteome, this technique is able to reveal events after protein degradation and the binding of peptides to distinct MHC I molecules. Interestingly, there is only a weak correlation between mRNA copy number and the density of corresponding MHC ligands¹⁷ indicating that the presentation of peptide antigens has to be investigated directly to obtain pertinent information about the immunological consequences of the expression of transgenes. No peptides originating from the transgenic proteins were detected. The detection limit of the method is about 100 fmol, so peptides present at 6–10 copies per cell could be detected; nevertheless, we have to consider the possibility that peptides

from the two transgenic proteins are presented by MHC I molecules in amounts too low to be detected. Also, peptides derived from the degradation of the transgenic proteins may have low affinity for the MHC I molecules tested. Moreover, we cannot exclude the possibility that these peptides are presented by MHC II despite its low abundance in the cell lines used. In contrast to the absence of MHC ligands of transgenic origin, numerous changes in the presentation of peptides of self-origin were apparent, although, most probably, we only detected the largest effects because more subtle changes may have escaped detection for the reasons discussed above. Abundant MHC ligands such as those detected by the analytical protocol can repeatedly be found in batches from cultured cells harvested at different time points^{15,18–22} which underlined evidence for the observed impact of transgene expression on the presentation of peptide antigens by MHC I.

The presentation of self-peptides may result from the phenomenon of insertional mutagenesis, commonly associated with integrative vectors. Sequences randomly inserted into the genome can modify the expression of genes in the proximity of the integration sites. Indeed, such insertions may be deleterious and even lead to the activation of oncogenes. In a clinical gene therapy trial for X-linked severe combined immune deficiency, 3 of 13 patients developed leukemia 30 months after treatment as a consequence of oncogene activation by the integrating vector genome.²³ Activation of oncogenes is accompanied by substantial changes at the proteome level²⁴ which may ultimately provoke appearance and disappearance of peptide antigens of self-origin. In the context of infection with HIV, di Marzo Veronese *et al.* showed that infected cells from HIV-1 positive subjects overexpressed vinculin and that three peptides originating from this self-protein are presented by MHC I at the surface of these infected cells, leading to the activation of specific cytotoxic T-lymphocytes.²⁵ Kane *et al.* recently reported that lentiviral vectors may provoke somatic cell reprogramming as a result of dysregulated host gene expression in the vicinity of integration sites.²⁶

Mechanisms other than those associated with genotoxicity may also contribute to the changes in the presentation of peptide antigens, including particular mechanisms involving the fusion proteins allowing rapamycin-inducible expression of transgenes. In our model, these transgenic proteins were strongly expressed under the control of a CMV promoter. This could lead to interference with the proteome by competing for the protein synthesis and degradation machinery. Furthermore, specific interference of these transgenic proteins might trigger changes in the presentation of self-peptides by MHC I. The transgenic proteins include domains from the human Frap kinase and the Frap kinase binding protein, which may interfere with the endogenous Frap kinase and Frap kinase binding protein in the regulation of autophagy.²⁷ Recently, it has emerged that autophagy is a pathway for the processing of antigens presented by MHC I.²⁸ Consistent with this possibility, we detected several peptide antigens originating from membrane proteins that could not have been produced by proteasome-mediated proteolysis but might have got access to MHC I by autophagy-mediated digestion of membrane components and mitochondria.²⁹ However, these peptides may also be produced by translation of mRNA using alternative translational reading frames.³⁰

Regardless of the molecular mechanism involved in the changes in the repertoire of MHC I-presented peptides, our coculture experiments revealed that transduction of cells with lentiviral vectors might affect the immunogenicity of individual transduced cells. In particular, transduced cells were able to activate and trigger proliferation of T cells. Cytotoxic T cells may arise, if the presentation of the respective MHC peptide complexes in the thymus is insufficient during the selection process leading to self-tolerance. In clinical settings, this phenomenon could lead to an immunological rejection of the transduced cells and thereby prevent a gene therapy protocol achieving its aims.

Any adverse immune reaction probably depends on a variety of factors including cell type, the site of vector integration and the HLA haplotype. Furthermore, the changes in the presentation of peptides by MHC I may only be recognized, if cytotoxic T cells bearing appropriate T cell receptors are available in the repertoire. We studied three cell lines, and only one, the transduced D407-derived cell clone, induced proliferation of CD4⁺ T cells (comprising cytotoxic CD8⁺ T and NKT cells). It is therefore likely that immune reactivity to transduced cells depends on the details of each individual case and context. Most probably, following administration of lentiviral vectors to patients, only a subset of the broad diversity of transduced cells could be eliminated by the immune system. Interestingly, in a recent trial of gene therapy for β -thalassemia, one particular clone of transduced cells came to predominate over time and provided therapeutic efficacy,³¹ suggesting that the majority of the transduced cells were probably successively eliminated by the immune system.

The effects on the presentation of peptide epitopes might not be restricted to MHC I. They might also be found in the repertoire of peptides presented by MHC II as transduced HER 911 cells (but not D407 or HEK 293T cells) revealed, compared with wt controls, significantly enhanced capacity to activate CD3⁺CD4⁺ allogeneic T cells, although no significant proliferation was observed (**Supplementary Figure S2**). It would therefore be of interest to assess the effect of the vectors on the presentation of MHC II peptides. These studies should also address whether activated CD4⁺ cells belong to effector or regulatory subtypes which induce adverse immunity and immune tolerance, respectively. Indeed, particular individual subsets of transduced cells might escape elimination by the immune system by activation of tolerance-mediating CD4⁺ regulatory T cells, a scenario frequently observed in studies monitoring the progression of tumors in patients.³²

Taken together, our findings provide the first evidence (i) that pathways involved in processing of peptide antigens and their presentation by MHC I render therapeutic manipulations within the genome of particular target cells visible to the immune system and (ii) that the immune system may recognize these events and respond.

Materials and methods

Cell culture, lentiviral gene transfer, and generation of stable cell lines. All cell lines were maintained at 37 °C under a water-saturated atmosphere of 5% CO₂/95% air. HER 911¹⁴ and HEK 293T cells were cultivated in Dulbecco's modified Eagle medium supplemented with 10% fetal calf serum, 20

U/ml penicillin G, and 20 μ g/ml streptomycin sulfate; D407 cells¹³ were cultivated in Dulbecco's modified Eagle medium containing 5% fetal calf serum, and antibiotics as above.

For gene transfer, 20,000 cells were incubated overnight with aliquots of the transactivator expression vector and the GDNF secretion vector⁹ (**Figure 1a**), each corresponding to 60 ng of capsid protein p24. Single viable cells were then seeded one in each well of 96-well culture dishes. After 3 weeks, wells were screened for cell clones which were tested for inducible production of GDNF using the GDNF Emax immunoassay system (Promega, Charbonnières, France). Appropriate clones released GDNF into the medium after addition of the rapamycin derivative AP 21967 (10 nmol/l, ARIAD, Cambridge, MA) but not in the absence of the inducer. These clones were amplified in the absence of inducer to 10¹⁰ cells.

Extraction and modification of MHC I-presented peptides. MHC I-presented peptides were obtained by immune precipitation of MHC I molecules from the amplified cell lines as described³³ using the HLA-A, -B, and -C specific antibody W6/32 immobilized on sepharose, acid treatment and ultrafiltration. As described previously,¹⁷ peptides from wt and transduced cells were concentrated by lyophilization, dissolved in 500 μ l water, modified by guanidination, and then nicotinylated with ¹H₄ and ²H₄ nicotinoyloxy-succinimide (LGC Promochem, Molsheim, France), respectively.

LC-ESI-MS and LC-ESI-MS/MS analyses. The modified peptide extracts were analyzed as described¹² using a reversed phase nanoLC-2D system (Eksigent, Darmstadt, Germany), coupled to a hybrid quadrupole orthogonal acceleration time-of-flight MS/MS (Q-TOF Ultima; Micromass/Waters, Saint-Quentin en Yvelines, France) equipped with a micro-ESI source. Results for mixed transduced and wt samples were recorded in an LC-ESI-MS experiment without fragmentation. For sequence analysis, results for transduced and wt samples were recorded separately in individual LC-ESI-MS/MS experiments. Fragment spectra were analyzed manually and database searches (Proteomics Department at the Hammersmith Campus of Imperial College London, National Center for Biotechnology Information, Expressed Sequence Tag) involved using the Multiple Alignment System for Protein Sequences Based on Three-way Dynamic Programming (MASCOT, <http://www.matrixscience.com>). The NetMHCpan 2.3 Server (<http://www.cbs.dtu.dk/services/NetMHCpan>)¹¹ was used to search for MHC I binding partners of the peptide sequences found.

HLA-class I and class II genotyping. DNA was extracted from HEK293T, HER911, and D407 cells using a salting-out technique. HLA medium resolution typing for HLA-A, -B, -C, DRB1, and DQB1 was performed using PCR-sequence specific oligonucleotide (SSO) Luminex kits (LABType SSO A Locus, LABType SSO B Locus, LABType SSO DRB1, and LABType SSO DQB1; One Lambda, Canoga Park, CA).

Immunogenicity assay. Unfractionated allogeneic peripheral blood mononuclear cells (HLA-A*02:29:02, HLA-B*40:02:44:03, HLA-C*02:02:16:01, HLA-DRB1*07:01:11:01, HLA-DQB1*02:02:03:01, 1 \times 10⁵ cells) purified from a blood sample from a healthy donor were labeled with the carboxyfluorescein succinimidyl ester (CFSE, 10 μ mol/l for 10 minutes at 37 °C) and then incubated with 1 \times 10⁴ target cells (transduced

cells, wt cells, or allogeneic stimulatory peripheral blood mononuclear cells for control). Before incubation with peripheral blood mononuclear cells, target cells were treated with mitomycin C (50 µg/ml) for 30 minutes. After 6 days of coculture, cells were stained with eFluor780-conjugated anti-CD3 (UCHT1; eBiosciences, Paris, France), phycoerythrin-conjugated anti-CD25 and allophycocyanin-conjugated anti-CD4 antibodies, and with 7-aminoactinomycin D to exclude dead cells (BD Biosciences, Le Pont de Claix, France). T cell activation and proliferation was then monitored by flow cytometry using a Canto II flow cytometer (BD Biosciences). Results were analyzed with BD Diva software (BD Biosciences) and are reported as mean values ± SE of at least three independent experiments. One-way analysis of variance followed by the Student-Newman-Keuls test (SigmaStat software; Systat Software, Chicago, IL) were used for statistical analyses and a *P* of <0.05 was considered significant.

Supplementary material

Figure S1. (a) Flow cytometric analysis of the expression of MHC I, MHC II (HLA-DR, HLA-DQ, and HLA-DP), CD80, and CD86 molecules and transduced HER 911 cells. (b) Expression of immune relevant molecules by wt and transduced HEK 293T, HER 911 and D407 cells.

Figure S2. Allogeneic immune responses to wt and transduced cells mediated by MHC II.

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