Vaccination strategy to target lysyl oxidase-like 4 in dendritic cell based immunotherapy for head and neck cancer

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Abstract. Overexpression of lysyl oxidase (LOX) is associated with the invasive potential of metastatic breast and head and neck cancer (HNC) cells and reduced metastasis-free and overall survival. Recently, we have demonstrated up-regulation of a new member of the LOX family, lysyl oxidase-like 4 (LOXL4), in invasive HNC revealed a significant correlation between LOXL4 expression and local lymph node metastases and higher tumour stages. The objective of this study was to examine whether cellular LOXL4 may provide an effective target for cell-mediated immunotherapy in invasive tumours associated with LOXL4 overexpression. As a feasibility study we expressed LOXL4 mRNA in immature dendritic cells derived from human peripheral blood mononuclear cells (PBMC). LOXL4 protein expression was ascertained using Western blotting and immunocytochemistry with polyclonal rabbit anti-LOXL4 antibody. The successfully transfected immature dendritic cells (DCs) were induced to mature with GM-CSF, IL-4, IL-18, TNF-α, IL-6, and PGE2, and then used to stimulate T cell enriched non-adherent fraction of PBMC. LOXL4 specific T cell stimulation induced cytotoxic T lymphocyte (CTL) response was monitored using IFN-γ secretion from the non-adherent PBMC fraction exposed to mature, LOXL4 transfected DCs acting as the antigen presenting target cells. LOXL4-DC stimulated T cells produced higher IFN-γ secretion compared to unstimulated T cells and T cells stimulated with untransfected DCs, in the presence of the pan-DR-epitope (PADRE). These initial results demonstrated the potential for LOXL4-transfected DCs to serve as efficient tumour vaccine and support their suitability as a vaccination strategy applicable to cancer patients with tumour specific up-regulation of LOXL4.

Introduction

A number of cancer-related antigens have been identified to contain epitopes that can be recognized by specialized immune cells, predominantly by cytotoxic T lymphocytes (CTL). The possibility of tumour-specific therapy has led to continued efforts to develop antigen specific immunotherapeutic strategies for different malignancies.

The essential role of the extracellular matrix (ECM) amine oxidase, lysyl oxidase (LOX), in breast tumour cell invasion was reported from our and other laboratories (1,2). We have also demonstrated that inhibition of LOX expression and catalytic activity reduced the migratory and invasive potential of these cells. Additional evidence indicates that this active LOX induced FAK/Scr/paxillin mediated mechanism was critical in vivo role for LOX in invasive breast tumours was further supported by data in a mouse model for breast cancer (4) and our studies of breast cancer patients (5). LOX is a copper dependent amine oxidase that catalyzes the oxidation of lysine to semialdehyde, the precursor to the covalent cross linkages that stabilizes elastin and collagen fibers in the ECM (6,7). Micro- and tissue array and a prospective head and neck cancer studies have shown the expression of LOX to be elevated in hypoxic HNC (8). Additionally, LOX expression was found associated with lower distant metastasis-free survival and reduced overall survival in HNC patients (4). Furthermore, LOX regulation by hypoxia-inducible factor and association with hypoxia was reported in human breast and HNC (4,9). Inhibition of LOX activity eliminated metastasis in mice with breast tumours indicating that LOX is a candidate therapeutic target for preventing and treating metastases.

Multiple studies reported tumour-associated expression for other members of the LOX family, including LOXL (10),
LOXL1, LOXL3 and LOXL4 (2). We have found elevated LOXL4 in head and neck squamous cell carcinoma (HNSCC) compared to normal squamous epithelium (11). Results of our recent study demonstrated an overexpression of the LOXL4 transcript in 74% of invasive HNSCC and 90% of both primary and metastatic HNSCC cell lines tested, and a significant correlation between LOXL4 expression and local lymph node metastases versus primary tumour types and higher tumour stages (12). Immunohistochemistry demonstrated cellular overexpression of the LOXL4 protein that correlated with the increased mRNA transcription in HNSCC cells. These results strongly support an important role for LOXL4 in the molecular pathogenesis of invasive head and neck carcinomas and demonstrate that both LOX and LOXL4 are potential immunotherapeutic targets.

Bone marrow-derived plasmacytoid and myeloid dendritic cells (DCs) are the most potent antigen-presenting cells capable of activating native T cells (13-15). Loading DCs ex vivo with tumour antigens in the form of mRNA can stimulate potent anti-tumour immunity in tumour-bearing mice (16,17). Suggestive evidence of clinically related responses was first seen in phase I clinical trials in patients with prostate cancer immunized with prostate-specific antigen mRNA transfected DCs and in patients with renal cancer immunized with autologous tumour RNA transfected DCs (18-20). Since then, many patients have been treated with antigen-loaded DCs in different tumour types with varying protocols for the generation of DCs and application of the vaccine. While these therapies did not yet fully materialize their potential, all DC vaccines proved to be safe. In this feasibility study, we transfected cloned LOXL4 mRNA into immature dendritic cells and examined whether cellular expression of LOXL4 may provide an effective target for cell-mediated immunotherapy.

Materials and methods

Human subjects. Peripheral blood mononuclear cells (PBMC) were harvested from healthy volunteers using peripheral blood leukapheresis. All cells used in these experiments were obtained from human subjects, following informed consent through protocols approved by the ethics committee at our institution (AZ: A128/01; Kiel, Germany).

DC generation from peripheral blood. For DC culture, we adopted techniques described by Romani et al (21) and Morse et al (22) and implemented some modifications, particularly in the use of serum-free media and supplements. Briefly, peripheral blood mononuclear cells (PBMC) were incubated at 2x10^8 cells/30 ml in serum-free AIM-V medium (Life Technologies, Grand Diagnostics, St. Louis, MO, USA), and cells were resuspended over polysucrose/sodium diatrizoate (Histopaque; Sigma was further separated by density-gradient centrifugation.

Concentrated leukocyte fraction was generated through a 2-h restricted peripheral blood leukapheresis processing 6-8 l. Immature and mature DCs were characterized by flow cytometry. Briefly, 3x10^5 cells in 100 μl PBS were incubated with 5 μl each of monoclonal antibodies or isotype controls (Caltag, Hamburg, Germany) for 20 min in the dark at room temperature. Cells were washed three times in PBS and analysed by flow cytometry (FACSCalibur, Becton Dickinson, Heidelberg, Germany) and with the CellQuest/WinMDI software. DC preparations fulfilling certain phenotypic criteria as outlined in the results section were used for subsequent experiments.

Cloning strategy and sequence analysis. A purified double-stranded full length LOXL4 cDNA fragment was ligated into the pCR2.1 Topo cloning vector and amplified in E. coli competent cells during overnight incubation. Plasmid DNA was extracted and prepared for cycle-sequencing reaction by PCR using Dye Terminator (Applied Biosystems, Weiterstadt, Germany). Sequencing was performed using the M13 primer binding sites of the pCR2.1 Topo vector. The reaction mixture contained 1 μg plasmid-DNA, 10 pmol M13 upstream and downstream-primer mix, 5% (v/v) dimethylsulfoxide, and 4 μl premix (Applied Biosystems) in a total volume of 20 μl. PCR was performed for 25 cycles at 55°C for 15 sec, and 60°C for 4 min (two-step-anneling). After amplification the PCR product was extracted using absolute ethanol, redissolved in TSR-buffer (Applied Biosystems) and subjected to sequencing (ABI PRISM-310, Applied Biosystems). The sequence obtained was compared with sequences deposited in the Human Gene Bank Database, NCIB (accession no. AY036093).

Generation of LOXL4/pSp73/A64/NotI and in vitro transcription of LOXL4 mRNA. To generate in vitro transcribed LOXL4 mRNA, the LOXL4 cDNA was subcloned into the vector pSp73/A64/NotI, kindly provided by David Boczkowski (23). A 2277-bp fragment containing the entire coding sequence for LOXL4 was isolated from pCR2.1 Topo vector. LOXL4 cDNA was excised and inserted into pSp73/A64/NotI with HindIII (A'AGCT_T) and XhoI (C'TCGA_G). Subcloning generated the plasmid LOXL4/pSp73/A64/NotI, which contains the Amp®r, a T7 promoter and the PolyA-sequence. After linearization with SpeI, mRNA was generated by in vitro transcription with T7 RNA polymerase (Ambion mMessage mMachine kit, Austin, TX, USA). As a control DNA encoding jelly fish green fluorescent protein (eGFP) was used in some experiments.

RNA transfection and maturation of human DC. Transfection of immature DCs with RNA was performed by electroporation (Gene Pulser II, Bio-Rad Laboratories, Munich, Germany). In brief, DCs were washed twice in PBS, counted, and centrifuged at 300 x g for 10 min. Subsequently, DCs were resuspended at a concentration of 1x10^6 cells/ml in Opti-MEM medium (Invitrogen, Life Technologies, Karlsruhe, Germany). RNA (10 μg) (eGFP mRNA or LOXL4 mRNA) and 2x10^6 immature DCs were mixed in Opti-MEM for a 0.4 cm fusion chamber (Bio-Rad Lab.). Electroporation was performed...
under various conditions. The best electroporation results (high transfection rate, viability of cells ≥80% within the first 2 h) were obtained using the following settings: 0.6 kV/cm, 260 mV, 0% modulation, 40 Hz, one square-wave pulse of 16 ms. Subsequently, cells were allowed to rest for 5 min before transfer into fresh culture medium for a 2-h reconstitution phase in a humidified incubator at 37°C/5% CO2. Viability of cells was assessed by trypan blue dye exclusion in a hematocytometer before and 2, 24 and 48 h after electroporation.

Maturation of the transfected immature DCs was performed by culturing for 1 day in serum-free medium supplemented with GM-CSF, IL-4, IL-1β, TNF-α (10 ng/ml; CellConcepts), IL-6 (1000 U/ml; CellConcepts), and PGE2 (1 μg/ml; Sigma-Aldrich, Deisenhofen, Germany).

Western blot analysis and immunocytochemistry. For Western blot analysis, DCs were homogenised and cell lysates were subjected to SDS-PAGE. After electrophoresis the proteins were transferred to positively charged nitrocellulose membranes (Millipore, Eschborn, Germany).

For immunocytochemistry, DCs were fixed with ice-cold methanol for 10 min and blocked with PBS (pH 7.3) containing 1% (w/v) BSA (Sigma) for 1 h at room temperature. A primary affinity purified polyclonal rabbit anti-LOXL4 antibody from our own production (12) was added to the cell preparations for 1 h at room temperature, followed by incubation with a biotin-conjugated swine anti-rabbit IgG secondary antibody (Dako, Hamburg, Germany) at room temperature for 1 h. The optimum concentration of the primary antibody (1:400) was determined using serial dilutions of highly purified synthetic peptide (Eurogentec, Seraing, Belgium). When diluted 1:400, the antibody detected LOXL4 at a minimal concentration of 0.1 μg/ml in Western blotting, and stained a single band in the Western blot analysis of total protein extracts from the carcinoma cells. Negative controls were obtained by omission of the primary antibody and by incubating the primary antibody with specific blocking peptide in 5-10-fold molar excess before staining.

Generation of human LOXL4 antigen specific T cells and INF-γ secretion assay. LOXL4 specific T cells were generated using a protocol adopted from Heiser et al (18); in brief, the T cell-enriched non-adherent fraction of PBMC obtained after the DC plastic adherence step was used. Non-adherent PBMC were cultured in serum-free medium supplemented with recombinant human IL-2 (20 U/ml; CellConcepts) and recombinant human IL-7 (10 ng/ml; CellConcepts). Cells were stimulated weekly at least four times with autologous LOXL4-mRNA-transfected DCs in absence or presence of the pan-DR-epitope (PADRE, Pharmexa, Denmark) in two separate experiments.

Target cells (LOXL4-mRNA-transfected DC, unstimulated T cells or untransfected DC in absence or presence of PADRE) were incubated with CTL at E:T ratio of 100:1, 30:1 and 10:1 in 96-well U-bottom plates for 12 h at 37°C in 5% CO2. Subsequently, 150 μl of supernatant was harvested and used in cytokine secretion assay and analysed by FACS array according to manufacturer's protocol (CBA FLEX system, BD Biosciences, San Jose, USA).

Results

In vitro transcription of the LOXL4 mRNA. To clarify that the LOXL4 sequence derived from the LOXL4/pCR2.1 clone was accurate, plasmid DNA was sequenced. The sequencing data revealed the full coding and additional 440 bp non-coding sequences at both ends of the LOXL4 mRNA. Both end sequences exhibited identity to human LOXL4 cDNA (Accession no. AY036093). In order to produce LOXL4 mRNA, the cloning vector LOXL4/pSP73-Sph/A64/NotI was linearized with SpeI and in vitro transcription was performed using T7 RNA polymerase. The resulting material was subjected to gel electrophoresis that revealed two bands of expected size for the plasmid and the LOXL4 mRNA.

RNA transfection of DCs. Transfection of mRNA into DCs was performed using electroporation. In order to establish the efficiency of electroporation, mRNA encoding eGFP was used. Successful transfection resulted in translation of the mRNA by the DCs resulting in fluorescent eGFP expression, that was detected using flow cytometry (Fig. 1). Optimized electroporation using reproducible settings resulted in a transfection rate of >80%.

Using the optimized settings, LOXL4 mRNA was successfully transfected into immature DCs. LOXL4 transfected DCs showed the same viability as GFP transfected DCs. Expression of the LOXL4 protein in DCs cells was analyzed by two different methods. Western blot analysis of cell extracts revealed expression of the correct size LOXL4 in DCs (Fig. 2A, lane 2). LOXL4 expression and cellular localization was further confirmed by immunocytochemical (ICC) staining using polyclonal rabbit anti-LOXL4 in transfected DCs (Fig. 1B). The majority of cells displayed a positive cytoplasmatic ICC staining, and no staining of non-transfected DCs. Non-transfected DCs on Western blotting and ICC (Fig. 2A) using an unrelated control antibody and secondary antibody alone were negative for LOXL4 staining (data not presented).
Phenotyping of DCs. Immature DCs were generated from PBMC derived from healthy volunteers. By flow cytometry, DCs were analyzed for the presence of antigens known to be characteristic for monocyte-derived DCs (23). Immature monocyte-derived DCs were positive for HLA-DR, CD1a, CD80, CD86 and negative for CD14 and CD83. After electroporation and maturation using TNF-α and LPS induction, the DCs displayed enhanced expression of HLA-DR, CD1a, CD80, CD86, and also stained positive for CD40 and CD83 which has been specified to be characteristic profile (14) for mature DCs (Fig. 2C displays data for CD86, CD83 and HLA-DR).

Antigen-presenting capacity of LOXL4-RNA-transfected DCs. To evaluate the antigen-presenting capacity of LOXL4-RNA-transfected DCs, non-adherent PBMC obtained from healthy volunteers were stimulated with autologous LOXL4-RNA-transfected DCs. Subsequently, INF-γ secretion, a marker for activated antigen specific T cells, into the supernatant was determined. Results from these experiments demonstrate that LOXL4-specific T cells have been successfully induced (Fig. 3). Stimulation of non-adherent PBMC with LOXL4-RNA-transfected DCs resulted in a 26% increased INF-γ secretion after stimulation with LOXL4-positive target cells compared to untransfected DCs at E:T ratio of 100:1. The presence of PADRE further enhanced T helper cell activation increasing INF-γ secretion by 116% compared to LOXL4-negative target cells. Stimulation with LOXL4-RNA-transfected DCs at E:T ratio of 30:1 and 10:1 and stimulation with untransfected DCs failed to stimulate PBMC to secrete INF-γ. In the presence of PADRE, unspecific increase of INF-γ secretion was observed in these controls.
Expression in the normal mucosa. As the human lysyl oxidase—
identified in head and neck cancer are self-proteins with
antitumour immunity. However, most tumour antigens
targets for specific cytotoxic T lymphocytes mediating
tumour immunity by priming T cells to tumour antigens.
It is well accepted that DCs are capable of inducing anti-
tumour immunity by priming T cells to tumour antigens. Many tumour antigens have been identified as potential
targets for specific cytotoxic T lymphocytes mediating
antitumour immunity. However, most tumour antigens
identified in head and neck cancer are self-proteins with
expression in the normal mucosa. As the human lysyl oxidase-
like protein 4 was not detected in healthy oral epithelia, and
up-regulation of LOXL4 expression in invasive HNC was
associated with significant correlation to higher tumour
stages and metastasizes, provide substantial evidence that
targeting LOXL4 is a promising approach of DC based
vaccination therapy (12).
We successfully transfected LOXL4 mRNA into dendritic
cells derived from PBMC, ascertained LOXL4 protein
expression in these cells and demonstrated that the LOXL4
expressing DCs stimulate the T cell enriched non-adherent
fraction of PBMC. Using INF-γ secretion assay for monitoring
LOXL4-specific CTL induction our results support the use of
LOXL4 mRNA-transfected DCs as a promising vaccination
tool for head and neck cancer therapy.

**RNA transfection of DCs.** Transfer of exogenous genetic
material into foreign cells can be accomplished actively by
physical or chemical methods or passively by diffusion. The
RNA transfection method provides the insertion of genetic
information into the cytoplasm for direct translation of genes
into proteins without affecting the genome. Therefore this
technology, while efficient, does not carry the high risk of
gene therapy. The advantages of using mRNA loaded DCs are
3-fold: first, unlike synthetic peptides encoding HLA
restricted epitopes, mRNA spanning the entire coding sequence
of a protein, LOXL4 in our case, is likely to encode multiple
epitopes corresponding to many different HLA-restricted
alleles; second, as previously shown, mRNA transfected DCs
can function as specific targets in CTL assays and provide a
general platform to monitor CTL responses without the need
to culture and label tumour cells as targets (14,17,18,20); third,
RNA pulsed DC vaccines can be used to stimulate CTL from
virtually all cancer patients regardless of their HLA genotype.

**LOXL4 as a target tumour antigen and its expression in DCs.**
We have previously observed significant correlation between
LOXL4 expression and regional lymph node metastases versus primary tumour types (p<0.01) and higher tumour
stages (p<0.01). As up-regulated LOXL4 is characteristic
for head and neck tumour cells, while its expression is
undetected or very low in normal squamous epithelia and
similarly low in other normal tissue types (1,11,12), LOXL4
is an excellent candidate tumour antigen for immune therapy
for head and neck cancer and potentially for other tumour
types over-expressing LOXL4 (1,2,5). The target LOXL4
cDNA, was originally generated and cloned into the pCR2.1
Topo vector (8) that we have modified for our expression
needs. We have developed an efficient electroporation protocol
using eGFP mRNA monitored by flow cytometry. Our results
revealed that the electroporation-based mRNA delivery system
was highly efficient and displayed low cytotoxicity against
DCs under optimal electrical settings. These observations were
consistent with other reports showing that the electroporation-
based mRNA delivery system is superior to the commonly
used techniques of lipid based transfection or passive pulsing
by providing better mRNA transfection efficiency (24-28).

**Discussion**

Following the optimized protocol, we have generated LOXL4
electroporated DCs and confirmed the specific expression of
the LOXL4 protein in these cells using Western blot analysis
and immunocytochemistry. Results from these analyses
revealed LOXL4 protein synthesis in DCs 24 h and 48 h after
cell transfection (Fig. 2).

**Maturation of DCs in vivo.** The cellular basis of the vaccine, immature DCs were generated and cultured from PBMC. The activation of DCs is a crucial step in the initiation of adaptive
immunity because it links peripheral events initiated by the
encounter with pathogens to the activation and expansion of
antigen-specific T lymphocytes during their typical migration
from the skin to lymphoid tissues. It is well recognized that a
wide variety of microbial products and endogenous signals
can trigger DC activation (29,30). Most researchers believe
that activated DCs applied in vaccines need to be matured
*in vitro* before application to enrich the cell's migration
capacity and to avoid the tolerance inducing potential of
immature DCs to self-antigens (18,31-33). In our study,
matured DCs were generated and cultured from PBMC, ascertained LOXL4 protein
expression in these cells and demonstrated that the LOXL4
expressing DCs stimulate the T cell enriched non-adherent
fraction of PBMC. Using INF-γ secretion assay for monitoring
LOXL4-specific CTL induction our results support the use of
LOXL4 mRNA-transfected DCs as a promising vaccination
tool for head and neck cancer therapy.

**Function in vitro assessment of the vaccine.** RNA-transfected
DCs expressing different target antigens have been shown to
successfully trigger antigen-specific T cell responses *in vitro*.
The induction of a strong immune response by antigen specific
cytotoxic T lymphocytes, the effector cells for tumour cell
killing, is the primary goal for any vaccination strategy (29,34).
In our study, LOXL4-stimulated T cells showed increased
secretion of INF-γ when targeting LOXL4-positive T cells.
The *in vitro* stimulation of T cells with LOXL4-transfected
DCs and the resulting induction of LOXL4-specific CTL
provided evidence for the potential of these DCs to serve as
tumour vaccine. The background signal on LOXL4-negative
target cells might be due to the fact that the cells we used
have been obtained directly from stimulated cell culture and
were still secreting some INF-γ. As known from preclinical
and clinical trials, PADRE induces important co-stimulatory
signals that activate helper T cells (35). When combined with
LOXL4-stimulation, a PADRE-induced helper T cell
response significantly enhanced the LOXL4-specific T cell
response referring to LOXL4 specific CTL induction. This
activating effect was not observed at E:T ratio lower than
100:1.

In conclusion, we have demonstrated the successful
generation of a DC-based vaccine targeting the novel antigen,
LOXL4. Our results provide evidence for the *in vitro* induction
of LOXL4-specific CTL as a functional proof towards the
development of LOXL4-based anticancer vaccine. This
vaccine may serve as part of therapeutic strategies applicable
for cancer patients with tumour cell specific up-regulation of
the LOXL4 gene, particularly those with advanced stages of HNSCC.

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