Molecular Backgrounds of ERAP1 Downregulation in Cervical Carcinoma

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1. Introduction

Cervical carcinoma is the third most common cause of cancer-related death among women worldwide. It is caused by infection of the uterine cervix epithelium by oncogenic types of the human papillomavirus (HPV), followed by viral persistence and progressive malignant transformation, leading to a spectrum of premalignant lesions (cervical intraepithelial neoplasia; CIN) and, ultimately, cervical carcinoma [1]. The antigen processing machinery (APM) and human leukocyte antigen (HLA) class I-mediated peptide presentation are important determinants of the processing and presentation of HPV-derived peptides and are therefore significant factors in recognition and subsequent lysis of cervical carcinoma cells by cytotoxic T lymphocytes [2].

Downregulation of various APM components and of HLA class I has been shown to be associated with detrimental survival among cervical carcinoma patients [3]; in particular, downregulation of ERAP1 (endoplasmic reticulum aminopeptidase associated with antigen presentation 1) has been demonstrated to be an independent predictor of overall and disease-free survival [3]. ERAP1 is responsible for length-specific N-terminal trimming of peptides (derived from intracellular proteins) before presentation by HLA class I molecules [4, 5]. It is therefore hypothesized to be an important determinant of the repertoire of presented peptides, as has been shown in various ERAP1 downregulated mouse models [5–7]. Moreover, genetic variation in the ERAPI gene is associated with both increased cervical carcinoma risk and decreased survival among patients [8–10]. These data suggest that ERAP1 is an important factor in tumor immunogenicity and cervical carcinogenesis [11]. In addition, downregulation of ERAP1 protein expression (occurring in approximately 15% of cases) is associated with worse clinical outcome in cervical carcinoma patients [3].

Although several studies have shown that various viral proteins, including the HPV E7 oncoprotein, can interfere with APM-related cellular processes (physically or at the transcriptional level) [12], the mechanisms leading to ERAP1 downregulation in cervical carcinoma remain largely
unknown. Recent data suggest that a single nucleotide polymorphism (SNP) in the gene is associated with downregulation of the corresponding protein [9]; however, the existence of a direct causal relation between this SNP and downregulation remains to be established.

To address the issue of ERAP1 downregulation in cervical carcinoma, we have examined ERAP1 mRNA expression in tumor cells from a panel of cervical carcinoma lesions with known ERAP1 downregulation at the protein level. Subsequently, we performed an evaluation of possible molecular mechanisms for inactivation of the ERAP1 gene in flow sorted tumor cells from this series of clinical specimens.

2. Materials and Methods

2.1. Tumor Specimens. From 109 patients with cervical carcinoma who underwent radical hysterectomy with bilateral pelvic lymphadenectomy (by the same surgical team) between 1985 and 1999, formalin-fixed, paraffin-embedded tissue blocks were retrieved from the archives of the Department of Pathology, Leiden University Medical Center, Netherlands. All patients were inhabitants of Netherlands and had not received preoperative radiotherapy or chemotherapy. Mean age was 48.5 years, the youngest patient being 24 years and the oldest 87 years at the time of surgery. The use of clinical material was approved by the institutional review board according to the guidelines of the Dutch Federation of Medical Research Associations; specific patient consent for this study was waived as patients had given general consent for use of operative specimens for research purposes at the time of surgery.

2.2. Tumor Dissociation, Staining, Flow Sorting, and DNA Extraction. Formalin-fixed, paraffin-embedded cervical carcinoma tissue blocks (identified by previous immunohistochemical staining of the tissue microarray) were trimmed if necessary to remove normal epithelium and CIN. The remaining tumor tissue was dissociated, stained for keratin, vimentin, and DNA as previously described [13]. The samples were analysed using a FACS Calibur (BD Biosciences, San Jose, CA). The vimentin-negative keratin-positive (V−K+) fraction, which represents the epithelial cell (tumor cell) subpopulation, and vimentin-positive keratin-negative (V+K−) fraction, consisting of normal (stromal) cells, were flow sorted using a FACS Vantage (BD Biosciences, San Jose, CA). As previously described, this method yields a high range of purity of the different sorted factions [13]. Sorted cells were resuspended in isolation buffer (0.3 mg/mL proteinase K in 10 mM Tris-HCl (pH 8.3), 1 mM EDTA, and 0.5% Tween 20) at a concentration of 1000 cells/µL. After overnight incubation at 56°C and 5 min inactivation at 100°C, DNA samples were stored at −20°C until further analysis [14].

2.3. DNA Mutation Analysis. DNA amplification and sequencing were performed on flow sorted tumor and normal fractions as previously described with CAACCTCGACGGCTTC-CCCGG (nucleotide position 54 → 73 of GenBank accession number NM_016442) and GGCCTAGCTCCC-CCAGGACC (nucleotide position 276 → 257) as forward and reverse primers, respectively [15]. All sequences were visually analyzed with Mutation Surveyor DNA variant analysis software (Version 2.6i; Softgenetics, State College, PA).

2.4. RNA Isolation and Quantitative Real-Time PCR (QRT-PCR). RNA was isolated from macrodissected formalin-fixed, paraffin-embedded material and cDNA synthesis, qPCR, and analysis were performed as described previously [16]. Expression of ERAP1-specific mRNA was determined by QRT-PCR in an iCycler iQ (Bio-Rad Laboratories Inc., Hercules, CA) using CATCGGTTGGATGGATAAGAA (nucleotide position 3101 → 3121 of GenBank accession number NM_016442) and CATCCGTGGTCGTCAGCTTT (nucleotide position 3193 → 3175) as forward and reverse primers, respectively. Besides ERAP1, expression of the housekeeping gene beta-actin (ACTB) was also measured with primer sequences as described previously [17]. Expression of these genes was measured using the Sybr Green method (Eurogentec, Seraing, Belgium) according to the manufacturer’s instructions. For each measured gene, the required calibration curve was determined using a serial dilution of pooled cDNA from established cervical cancer cell lines (SiHa, Caski, C33, and Hela). Each measurement was performed in duplicate.

For the analysis, ERAP1 mRNA expression levels were calculated relative to the ACTB housekeeping gene for both ERAP1 downregulated cases and control specimens. Mean ERAP1 mRNA expression levels of both groups were compared using the independent samples t-test.

2.5. Microsatellite and Allelic Status Analysis. To investigate loss or retention of heterozygosity (LOH and ROH, resp.) of the ERAP1 gene locus on chromosome 5q15, in DNA isolated from flow sorted samples of the 16 ERAP1 downregulated cases, 4 microsatellite markers (D5S433, D5S1463, D5S107, and D5S2094) spanning the genomic region from 5q21 to 5q14 were amplified using 6-carboxy-2′,4′,4′,5′,5′,7′-hexachlorofluorescein- (HEX-) labeled primer pairs (Genome Database, http://www.gdb.org/). As no suitable microsatellite marker could be identified in the ERAP1 gene itself, 5 previously described single nucleotide polymorphisms (SNPs) (codons 56, 127, 276, 528, and 730 of the gene itself, 5 previously described single nucleotide polymorphisms (SNPs) (codons 56, 127, 276, 528, and 730 of the ERAP1 gene) were investigated using TaqMan SNP Genotyping Assays (Applied Biosystems, Inc., Foster City, CA) as described previously [8]. PCR, electrophoresis, and analysis were performed as described previously [18].

Since flow sorting yields highly pure stromal and tumor cell fractions, the threshold for complete LOH in analysis of the microsatellite markers was set high [19]. Comparing normal and tumor DNA, a reduction of at least 80% of one allele was defined as complete LOH, corresponding to an allelic imbalance factor (AIF) of ≥5. Retention of heterozygosity was defined as less than 30% reduction of one allele (AIF ≤ 1.4). The range between 1.5 and 5 was defined as allelic imbalance (AI). Haplotype loss was defined as loss of at least three markers.
3. Results

3.1. ERAP1 Protein and mRNA Expression. Results of immunohistochemical staining of the tissue microarray using an anti-ERAP1 antibody have been described previously [3]. In short, 16 cases (15%) showed partial loss of ERAP1 expression while normal expression was observed in the remaining cases. Moreover, ERAP1 downregulation was shown to be an independent prognostic parameter for shorter overall and disease-free survival ($P = 0.037$ and $P = 0.049$, resp.).

As it was unclear whether downregulation of ERAP1 was influenced by factors at a pre- or posttranscriptional level, mRNA was isolated from 14 cases with ERAP1 downregulation and from 22 randomly selected cases with normal ERAP1 expression levels. Distribution of the ERAP1 mRNA expression levels in both groups is demonstrated in Figure 1. A 21% downregulation of ERAP1 mRNA expression was found in the group with ERAP1 protein downregulation as compared to the group with normal protein expression ($P = 0.001$). Even when the cases with the lowest mRNA expression levels were excluded from analysis, the observed difference was still significant ($P = 0.003$).

To investigate the underlying mechanisms of ERAP1 protein and mRNA downregulation, the role of various molecular factors was tested.

3.2. Genotype Analysis. The results of analysis of various ERAP1 SNPs in relation to protein expression have been described previously [9]. Recently, a mutation in codon 349 (nucleotide position 1045; c.1045A>G) of the ERAP1 gene was reported in melanoma cell lines [20]; to investigate association of this mutation with protein expression, mutation analysis on DNA from pure tumor cell fractions (obtained by flow sorting) of the 16 ERAP1 downregulated cases was performed. The ERAP1-349 mutation was not found in any of the specimens with ERAP1 protein downregulation (data not shown).

3.3. LOH Analysis. As partial or complete LOH has been shown to be an important factor in various human carcinomas, the role of allelic instability in ERAP1 downregulation was assessed. Of the 16 specimens with ERAP1 downregulation, 14 were successfully analysed for LOH (Figure 2). The four microsatellite markers D5S433, D5S1463, D5S107, and D5S2094 exhibited either LOH or allelic imbalance in 21%–50% of cases, with complete LOH occurring in 14%–29% of cases. Overall, haplotype loss occurred in 21% of cases.

4. Discussion

A key pathway in antitumor immunity is the presentation of tumor derived peptides by HLA class I molecules to cytotoxic T lymphocytes. The formation and intracellular transport of these peptides are performed by the various components of the antigen processing machinery (APM). One of these components, ERAP1, has previously been shown to be an important determinant of the repertoire of HLA class I-presented peptides [4, 6, 7, 21, 22]. Furthermore, genetic variation in the ERAP1 gene has been shown to be significantly associated with risk of developing cervical carcinoma and with prognosis of cervical carcinoma patients [8–10]; specifically, a SNP at the ERAP1-127 locus was shown to be an independent predictor of both cervical carcinoma risk and cervical carcinoma-associated survival, possibly due to its location in the peptidase M1 domain of the ERAP protein [9]. Importantly, we have previously shown that ERAP1 downregulation is significantly associated with decreased survival in cervical carcinoma [3]. However, to date the mechanisms leading to ERAP1 downregulation in this tumor type have not been elucidated. Although a previous study has investigated the underlying mechanisms of ERAP1 downregulation in melanoma cell lines [20], no
investigations in clinical specimens have been performed to date. The present study therefore is the first description of such mechanisms in clinical specimens.

Firstly, we investigated whether the processes leading to protein downregulation occur at a pre- or posttranscriptional level by studying ERAP1 mRNA levels in patients with normal and decreased ERAP1 expression. mRNA downregulation was found to be significantly associated with decreased protein expression, which indicates that the factors influencing ERAP1 expression occur at a pretranscriptional level.

The possible pretranscriptional mechanisms of protein downregulation can be both genetic and epigenetic. As previous reports indicate that DNA methylation status is not a contributing factor in ERAP1 downregulation [20, 23], we did not investigate ERAP1 gene methylation in the present study.

At the genetic level, both single nucleotide polymorphisms (SNPs) and mutations can lead to altered expression. In our previous report, 5 SNPs (ERAP1-56, ERAP1-127, ERAP1-276, ERAP1-528, and ERAP1-730) in the ERAP1 gene were genotyped and investigated for association with ERAP1 protein downregulation; genetic variation at ERAP1-127 was found to be significantly associated with ERAP1 protein expression ($P < 0.001$) [9]. Until recently, no ERAP1 mutations had been studied in human tumors. However, a recent study identified a novel ERAP1 mutation in human melanoma cell lines [20]. This mutation could not be found in any of the ERAP1 cases showing low ERAP1 expression, nor in any patients with normal ERAP1 expression.

In addition to SNPs and mutations, loss of heterozygosity has been shown to be an important genetic factor in human carcinomas; in the present study, complete and partial LOH and haplotype loss were found to occur in a considerable proportion of cases with ERAP1 downregulation (up to 50%).

LOH and haplotype loss are relatively frequently occurring phenomena in various human carcinomas and have been found to be related to prognosis in colon carcinoma [24]. Our findings suggest that these phenomena occur frequently in cervical carcinoma with ERAP1 downregulation.

In conjunction with our previous reports on the association of molecular changes with clinical aspects, the present study is part of an ongoing effort to elucidate the effects of interaction between exogenous and endogenous factors in the etiology, pathogenesis, and prognosis of cervical cancer. This field, recently termed molecular pathological epidemiology (MPE), has made considerable progress in colorectal carcinoma [24–29]; we believe that our study is the start of the first MPE-based investigative analysis in cervical carcinogenesis. In our previous report, 5 SNPs (ERAP1-56, ERAP1-127, ERAP1-276, ERAP1-528, and ERAP1-730) in the ERAP1 gene were genotyped and investigated for association with ERAP1 protein downregulation; genetic variation at ERAP1-127 was found to be significantly associated with ERAP1 protein expression ($P < 0.001$) [9]. Until recently, no ERAP1 mutations had been studied in human tumors. However, a recent study identified a novel ERAP1 mutation in human melanoma cell lines [20]. This mutation could not be found in any of the ERAP1 cases showing low ERAP1 expression, nor in any patients with normal ERAP1 expression.

In conclusion, here we present the results of the first investigation of in vivo mechanisms of ERAP1 downregulation in cervical carcinoma. Taken together, the results of this study indicate that ERAP1 protein downregulation occurs at a pretranscriptional level and is associated with both genetic variation in SNPs and loss of heterozygosity. As ERAP1 downregulation has been shown to be a significant predictor of decreased survival in cervical carcinoma, insight into the possible underlying mechanisms of this downregulation may yield novel targets for the designs of antitumor immunotherapies.

**Conflict of Interests**

The authors declare that there is no conflict of interests.

**References**


