

Promoter hypermethylation and reduced expression of *RASSF1A* are frequent molecular alterations of endometrial carcinoma

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Alterations in the regulation of the RAS–MAPK pathway are frequent in endometrial carcinoma. *RASSF1A* is a tumor-suppressor gene that can regulate this pathway negatively. *RASSF1A* has been found to be inactivated by promoter methylation in some human tumors. The aim of the study was to assess the immunohistochemical expression of *RASSF1A* in normal endometrium and endometrial carcinoma, and to correlate its expression with *K-RAS* mutations, presence of microsatellite instability, *RASSF1A* promoter methylation, and clinicopathological data. *RASSF1A* immunostaining was evaluated in one tissue microarray constructed from 80 paraffin-embedded samples of normal endometrium, and two tissue microarrays constructed with a total of 157 endometrial carcinomas (one constructed with 95 endometrial carcinomas previously evaluated for *K-RAS* mutations, and microsatellite instability, and another one containing 62 endometrial carcinomas that were also subjected to *RASSF1A* promoter methylation analysis). *RASSF1A* immunostaining was correlated with cell proliferation (Ki67), *Cyclin D1* expression and clinicopathological data. Promoter methylation of *RASSF1A* was assessed by methylation-specific PCR. *RASSF1A* immunostaining was variable during the menstrual cycle in normal endometrium. *RASSF1A* expression was significantly reduced in 48% of endometrial carcinomas, particularly in tumors exhibiting microsatellite instability. *RASSF1A*-promoter methylation was very frequent in endometrial carcinoma (74%), and was frequently associated with reduced expression of *RASSF1A*. *RASSF1A*-promoter hypermethylation was common in advanced-stage endometrial carcinoma. The results suggest that reduced expression of *RASSF1A* may play a role in endometrial carcinogenesis by controlling cell proliferation and apoptosis through the MAPK-signaling pathway.

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Endometrial carcinomas can be classified in two main types;¹ type I are endometrioid adenocarcinomas.² They usually develop in perimenopausal women and are related to estrogen stimulation. Endometrioid adenocarcinoma may show microsatellite instability³ and mutations of the *PTEN*,⁴ *PIK3CA*,⁵ *K-RAS*⁶ and β -catenin genes.^{7,8} Type-II tumors are non-endometrioid carcinomas, which

tend to occur in older women, are unrelated to estrogen stimulation and are frequently associated to p53 mutations and chromosomal instability.⁹

The RAS-signaling pathway plays an important role in tumorigenesis. Mutations in the *RAS* oncogene have been detected in many different types of tumors. The RAS superfamily of small GTP-binding proteins has a fundamental role in cell growth and differentiation, transcriptional regulation and apoptosis. The frequency of *K-RAS* mutations in endometrial carcinoma ranges between 10 to 30%.⁶ In some series, *K-RAS* mutations have been reported to be more frequent in endometrioid adenocarcinoma showing microsatellite instability.⁶ In these tumors, *K-RAS* mutations are typically transitions, which may be preceded by abnormal DNA methylation.

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The fact that in endometrioid adenocarcinoma, microsatellite instability frequently coexists with *K-RAS* methylation-related transitions has led to the suggestion that these two alterations are close related.

The *RASSF1A* gene has been recently identified and mapped to chromosome 3p.21.3. *RASSF1A* is the longest isoform of *RASSF1*, a new group of RAS effectors thought to regulate cell proliferation and apoptosis.¹⁰ Expression of *RASSF1A* has been detected in any normal tissue tested. However, *RASSF1A* transcripts have been found to be lost in several cancer cell lines. Also, re-expression of *RASSF1A* dramatically reduces cell growth of tumor cells, supporting a role for *RASSF1A* as a tumor-suppressor gene. The *RASSF1A* gene is epigenetically inactivated in a large percentage of different human malignancies, including cancers of the lung, breast, kidney, bladder, stomach, ovary and neuroblastoma.¹¹

RASSF1A can either associate with oncogenic/mutant *K-RAS* or with GTP-bound *K-RAS* through heterodimerization with *NORE1*, a RAS-GTP-binding protein.¹² This assumption raises the hypothesis of *K-RAS-BRAF-RASSF1A* genes as partners in the same pathway.

Promoter methylation is a frequent mechanism of gene inactivation of endometrioid adenocarcinoma. It may involve many different genes participating in cell signalling, apoptosis and cell proliferation (*p14*, *PTEN*, *p16*, *TSP-1*, *IGF-2*, estrogen receptor- α , *HIC-1*, *MLH-1*). Inactivation of *hMHL-1* by promoter hypermethylation is present in the majority of endometrioid adenocarcinoma with microsatellite instability, which suggests that abnormal methylation is the first step in the acquisition of the mutator phenotype in endometrial carcinogenesis.¹³ Since altered methylation status is frequent in endometrioid adenocarcinoma with microsatellite instability, and also frequently coexists with *K-RAS* mutations, we may hypothesize that *RASSF1A* promoter hypermethylation could be frequent in endometrial carcinoma, possibly coexisting with any of these phenomena.

The main aim of the present study was to assess the frequency of reduced expression of *RASSF1A* and *RASSF1A*-promoter hypermethylation in endometrial carcinoma in correlation with the presence of microsatellite instability, *K-RAS* mutations and clinicopathological data. Also, the purpose of this study was to assess immunohistochemical expression of *RASSF1A* in normal endometrium, in different phases of the menstrual cycle, using microarray technology.

Materials and methods

Tissue Microarrays

Three tissue microarrays were constructed. The first tissue microarray was constructed from 80

paraffin-embedded samples of normal endometrium in different phases of the menstrual cycle, obtained from the surgical pathology files from Hospital Universitari Arnau de Vilanova de Lleida. The second tissue microarray was composed of 95 endometrial carcinomas, previously evaluated for *K-RAS* mutations and microsatellite instability.^{3,6} The third tissue microarray was constructed from 62 endometrial carcinomas that were also subjected to *RASSF1A*-promoter methylation (see below). Overall, the two last tissue microarrays comprised 157 endometrial carcinomas, obtained from the surgical pathology files of Hospital Universitari Arnau de Vilanova de Lleida and Hospital de Sant Pau, Barcelona, Spain. They included 47 grade-I endometrioid adenocarcinomas, 58 grade-II endometrioid adenocarcinomas, 26 grade III endometrioid adenocarcinomas, 15 serous carcinomas, four clear-cell carcinomas and seven mixed Müllerian malignant tumors. One hundred and eight tumors were stage-I, 15 stage-II carcinoma, 22 stage-III and one was a stage-IV carcinoma. Staging information was incomplete for 11 cases. The study was approved by the local ethical committee. Specific informed consent was obtained from all subjects.

A Tissue Arrayer device (Beecher Instrument, MD, USA) was used to construct tissue microarrays. Briefly, all the samples were histologically reviewed and representative areas were marked in the corresponding paraffin blocks. Two selected cylinders (0.6 mm in largest diameter) from two different areas were included in each case. Control normal tissues from the same endometrial carcinoma specimens were also included.

Immunohistochemical Study

Tissue-microarray blocks were sectioned at a thickness of 3 μ m, dried for 16 h at 56° before being de-waxed in xylene and rehydrated through a series of graded ethanol concentrations, followed by wash with phosphate-buffered saline. Antigen retrieval was achieved by heat treatment in a pressure cooker for 2 min in EDTA (pH 8.9). Before staining sections, endogenous peroxidase activity was blocked. The antibodies used were as follows: anti-*RASSF1A* (1:500 dilution; eB114-20H1; eBioscience), Ki67 (1:100 dilution; MIB-1; Dako) and cyclin D1 (1:25 dilution; DCS6; Dako). After incubation, the reaction was visualized with the EnVision Detection kit (Dako), using diaminobenzidine chromogen as a substrate. Sections were counterstained with hematoxylin. Appropriate positive and negative controls were also tested.

Immunohistochemical results were evaluated by two pathologists following uniform pre-established criteria. *RASSF1A* immunoexpression was graded semi-quantitatively by considering the percentage and intensity of the staining. A histological score

was obtained from each sample, which ranged from 0 (no immunoreaction) to 300 (maximum immunoreactivity). The score was obtained by applying the following formula: Histoscore = $1 \times (\% \text{ light staining}) + 2 \times (\% \text{ moderate staining}) + 3 \times (\% \text{ strong staining})$. The reliability of such score for interpretation of immunohistochemical staining in endometrial carcinoma tissue microarrays has been proven previously.^{14–17} Since each tissue microarray included two different tumor cylinders from each case, immunohistochemical evaluation was conducted after examining both samples. Finally, the percentage of positive nuclei in each case was used to assess the cellular proliferation (Ki67) and the cyclin D1 expression.

The reproducibility of tissue-microarray immunostaining was confirmed by comparing results from tissue microarray with those obtained in sections from the corresponding paraffin blocks of 37 randomly selected cases. The overall concordance was 89.2%. The Kappa index of agreement between the two methods ranged from 0.68 to 0.83.

Methylation-Specific PCR

Genomic DNA was isolated with the DNAeasy tissue kit (Qiagen GmbH, Hilden, Germany), from normal and tumor tissue from 57 patients with endometrial carcinoma. Samples were frozen after surgery and corresponded to 57 of the 62 endometrial carcinoma samples of the third tissue microarray. A specific informed consent was used and the study was approved by the local ethical committee. DNA was subjected to bisulfite treatment. Briefly, 1 μg of DNA was denatured by sodium hydroxide and modified by sodium bisulfite. DNA samples were then purified and treated again with sodium hydroxide, precipitated with ethanol and resuspended in water (EZ DNA Methylation kit; ZYMO RESEARCH, USA). *RASSF1A*-promoter methylation status was determined following protocols from previous studies.¹⁸ DNA was amplified with primers MF (GTGTTAACGCGTTGCGTATC) and MR (AACCCCGCAACTGAAAAACGA) for methylated DNA and UF (TTTGTTGGAGTGTGTTAATGTG) and UR (CAAACCCACAAAATAAAACAA) for unmethylated DNA. Each PCR reaction contained $1 \times$ buffer, 1.5 mM MgCl_2 and 1 U/ μl TaqGold (Applied Biosystems Inc., Santa Clara, CA, USA); 0.2 mM dNTPs (Biotools B&M Labs, SA, Madrid, Spain) and 0.2 μM of each primer. PCR conditions were as follows: an initial 10-min denaturation at 96°C followed by two cycles of 30 s at 95°C, 30 s at 67°C, 30 s at 72°C; an additional two cycles of 30 s at 95°C, 30 s at 65°C, 30 s at 72°C, a further two cycles of 30 s at 95°C, 30 s at 63°C, 30 s at 72°C cycles and a final extension step of 10 min at 72°C. Each PCR product was directly loaded on 3% agarose gel, stained with ethidium bromide and visualized under UV illumination.

Statistical Analysis

Statistical analysis was performed with a database using SPSS for Windows (version 11.5; SPSS Inc., Chicago, IL, USA). Immunohistochemical results were compared by Mann–Whitney *U* and Student's *t*-test when applicable. Associations between qualitative variables were assessed by χ^2 - or Fisher's exact tests. Correlations between quantitative variables were established through Pearson and Spearman ρ tests. Statistical significance was set at $P \leq 0.05$.

Results

Immunohistochemistry

In normal endometrium, *RASSF1A* was evaluated in all the 80 cases that were included in the first tissue microarray. *RASSF1A* immunostaining varied according to the different phases of the menstrual cycle. *RASSF1A* immunoreexpression was significantly higher in the proliferative phase (mean Histoscore, 45.96) than in the secretory endometrium (mean Histoscore, 31.58) ($P = 0.001$) (Figure 1). Staining was cytoplasmic. Stromal cells were positive in both, the proliferative and the secretory phases.

In endometrial carcinoma, *RASSF1A* was evaluated in 148 out of the 157 cases that were included in the second and third tissue microarrays. Six cases were excluded as there was no representative tumor tissue in the tissue microarray sections, and three cases were missed in the construction of tissue microarray. Overall, *RASSF1A* immunoreexpression was significantly reduced in endometrial carcinoma as compared with normal endometrium. Mean *RASSF1A* Histoscore was 38.28 in endometrial carcinoma and 70.00 in normal endometrium ($P = 0.007$). We observed negative staining for *RASSF1A* in 71 cases of endometrial carcinoma (48%), whereas remaining 77 cases were positive (52%). Staining was cytoplasmic and heterogeneous, and Histoscore ranged from 10 to 290 (Figures 2 and 3). Nuclear immunostaining was observed in 11 cases.

A slightly reduced *RASSF1A* immunostaining was observed in non-endometrioid endometrial carcinomas (mean Histoscore, 25.25) as compared with endometrioid endometrial carcinoma (mean Histoscore, 30.18). However, these differences were not statistically significant ($P = 0.457$). The correlation between *RASSF1A* immunoreexpression and histological grade or pathological stage did not get any significant difference ($P = 0.771$ and $P = 0.303$, respectively). There was no significant correlation between cell proliferation (Ki67) and *RASSF1A* immunoreexpression ($r = 0.088$, $P = 0.446$). Cyclin D1 nuclear expression was observed in 64.79% cases of endometrial carcinoma, and no correlation was found with *RASSF1A* protein expression ($P = 0.635$).

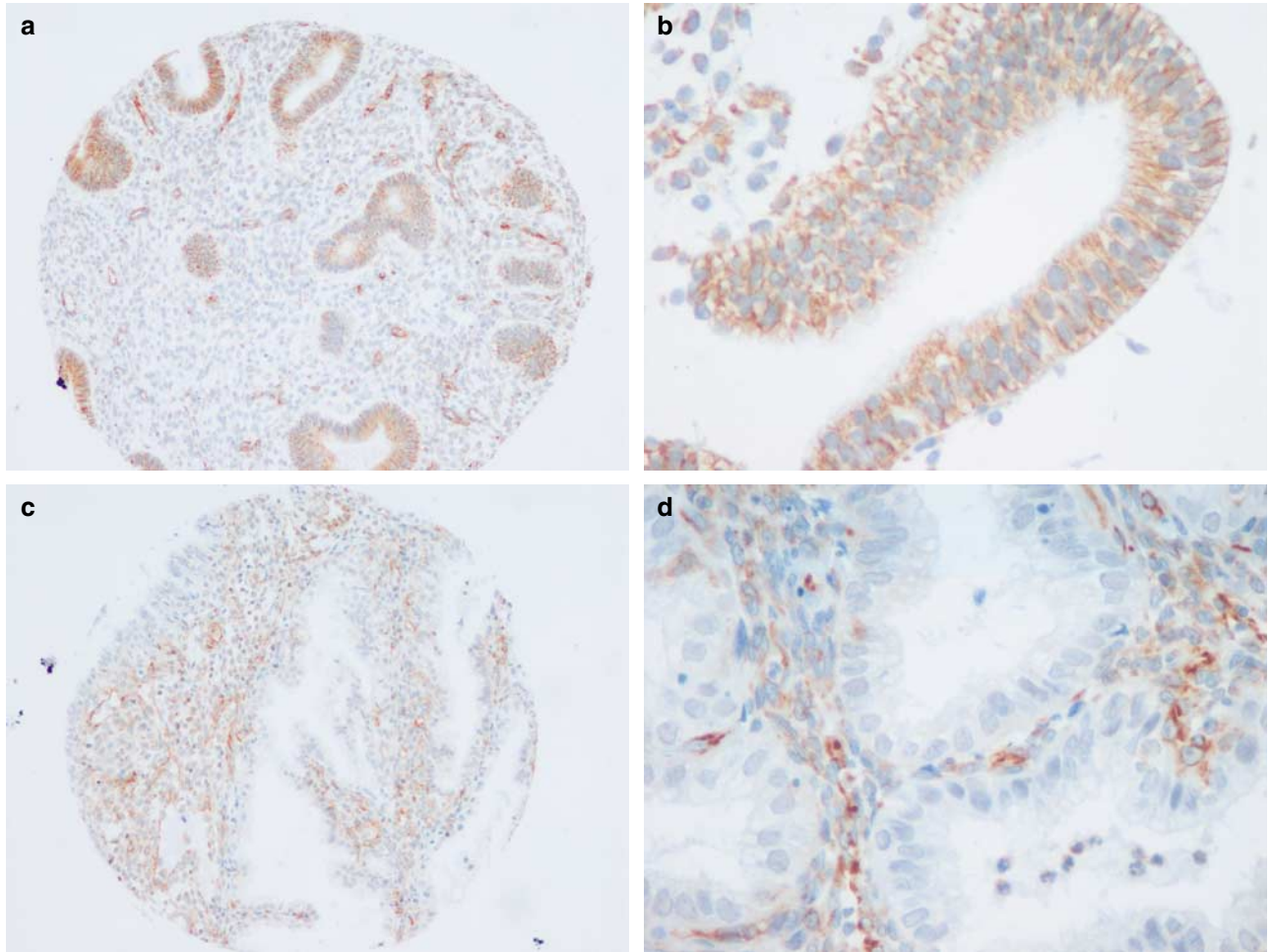


Figure 1 RASSF1A immunostaining was higher in the proliferative phase (a, b) than in the secretory phase (c, d).

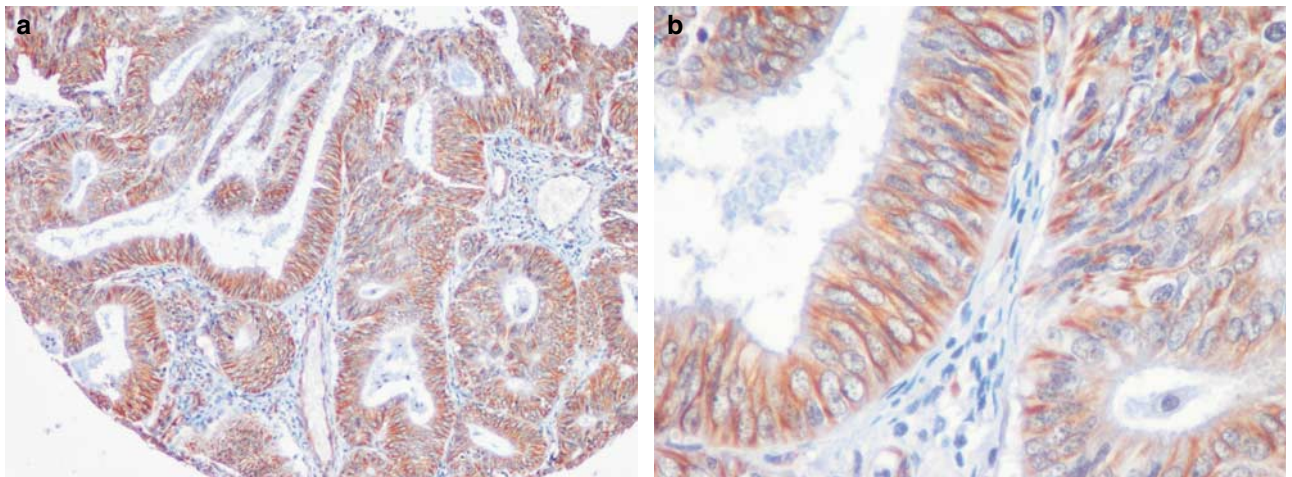


Figure 2 RASSF1A immunoeexpression in an endometrial carcinoma without *RASSF1A*-promoter hypermethylation. Notice the strong cytoplasmic staining (a, b).

Finally, RASSF1A immunoeexpression was correlated with microsatellite instability and *K-RAS* mutations in 67 cases of endometrial carcinoma

from the second tissue microarray. No significant correlation was found when we considered total absence of RASSF1A expression (Histscore 0) as

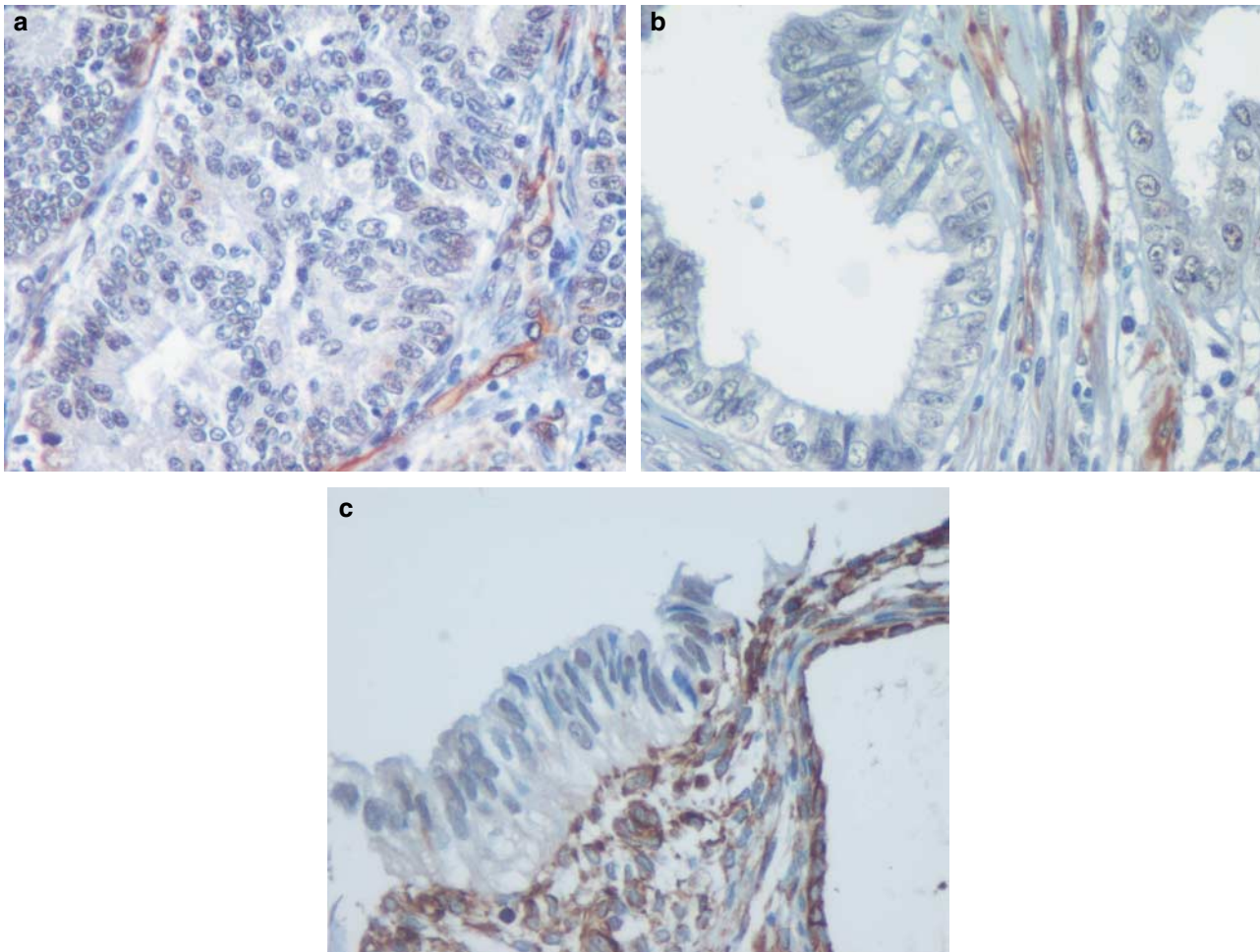


Figure 3 RASSF1A immunostaining in an endometrial carcinoma with *RASSF1A*-promoter hypermethylation. Notice that the cytoplasmic staining (a, b) is lower than in the case illustrated in Figure 2a and b. The corresponding normal tissue, exhibited positive staining (c).

negative ($P=0.510$ and $P=0.294$). However, association between RASSF1A immunostaining and presence of microsatellite instability turned out to be significant when we considered as negative those endometrial carcinoma with a Histocore less than 50. In other words, 18 of the 20 endometrial carcinomas with microsatellite instability exhibited Histocores less than 50 ($P=0.042$) (Table 1). Moreover, seven from the 10 cases with *K-RAS* mutations were negative for RASSF1A immunorexpression (Histocore less than 50), but differences were not statistically significant ($P=0.540$) (Table 1).

RASSF1A-Promoter Hypermethylation

RASSF1A-promoter hypermethylation was detected in 42 of the 57 (74%) endometrial carcinoma samples that were tested (Figure 4). In some cases, the degree of promoter methylation varied in different areas of the tumor (Figure 4, case 2, T1 and T2). Promoter methylation was not detected in normal tissue samples from the same patients with

Table 1 RASSF1A immunorexpression according to microsatellite instability and *K-RAS* mutational status

	<i>Microsatellite instability</i>		<i>K-RAS mutations</i>	
	<i>Positive</i>	<i>Negative</i>	<i>Positive</i>	<i>Negative</i>
RASSF1A positive	2	16	3	15
RASSF1A negative (Hscore < 50)	18	31	7	42
Total	20	47	10	57

endometrial carcinoma. Interestingly, the frequency of *RASSF1A*-promoter methylation increased in correlation with pathological stage (Table 2). *RASSF1A*-promoter methylation was observed in 67% of stage-IA or stage-IB endometrial carcinoma cases (18/26) as compared with 80% of stage-IC endometrial carcinomas (10/14) and 100% of stage-IIIa endometrial carcinomas (6/6). The vast majority of *RASSF1A* unmethylated tumors were stage-IA endometrial carcinomas (54.5%) (6/11), and only two

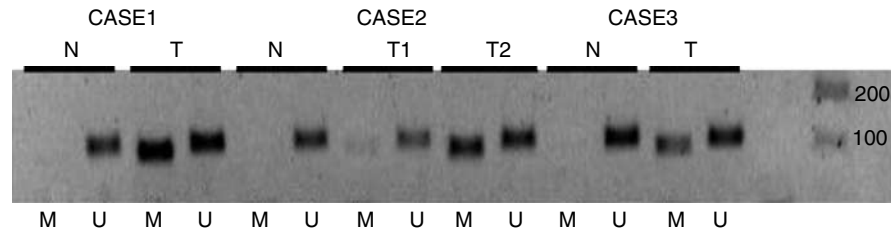


Figure 4 MS-PCR for the *RASSF1A* gene in endometrial carcinoma and normal tissue. U, PCR product with primers specific for unmethylated DNA; M, PCR product with primers specific for methylated DNA. *RASSF1A* is methylated in the three cases of endometrial carcinoma. Notice that the degree of promoter hypermethylation differed between two different areas of one tumor (T1 and T2) of case 2.

Table 2 *RASSF1A* methylation status according to pathological stage

Stage	<i>RASSF1A</i> methylated	<i>RASSF1A</i> unmethylated	Total
IA	5	6	11
IB	13	3	16
IC	10	4	14
IIA	2	0	2
IIB	6	2	8
IIIA	6	0	6

of the 16 stage-II or stage-III endometrial carcinomas were unmethylated (12%). *RASSF1A*-promoter hypermethylation was more frequent among endometrioid carcinomas (75%) (37 of 49 cases) as compared with non-endometrioid carcinomas (62.5%) (5 of 8 cases), but these differences were not statistically significant ($P=0.422$). *RASSF1A* was also analyzed in correlation with histological grade. Grade-I and grade-II endometrial carcinomas were more frequently methylated (71 and 86%, respectively) than grade-III tumors (57%). However, the differences were not significant ($P=0.152$).

Correlation between *RASSF1A* Immunostaining and *RASSF1A*-Promoter Hypermethylation

Finally, we assessed the correlation between *RASSF1A* immunoexpression and the methylation status of the promoter region of the gene in 53 of the 57 initial cases studied for *RASSF1A*-promoter hypermethylation. *RASSF1A* was methylated in 26 of 27 cases negative for *RASSF1A* immunostaining (Table 3). As expected, *RASSF1A* protein expression was dramatically reduced in cases with *RASSF1A*-promoter methylation (mean HistoScore, 30.38) as compared with unmethylated cases (mean: 101.92), and differences were statistically significant ($P=0.000$). However, 14 cases exhibited *RASSF1A* immunostaining in the presence of *RASSF1A*-promoter methylation. Interestingly, *RASSF1A* staining in these cases was heterogeneous and weaker. A comparison between cases with positive or negative *RASSF1A*-promoter hypermethylation among endometrial carcinoma with

Table 3 Correlation between methylation and *RASSF1A* immunoexpression

	<i>RASSF1A</i> methylated	<i>RASSF1A</i> unmethylated	Total
<i>RASSF1A</i> -positive immunostaining	14	12	26
<i>RASSF1A</i> -negative immunostaining	26	1	27

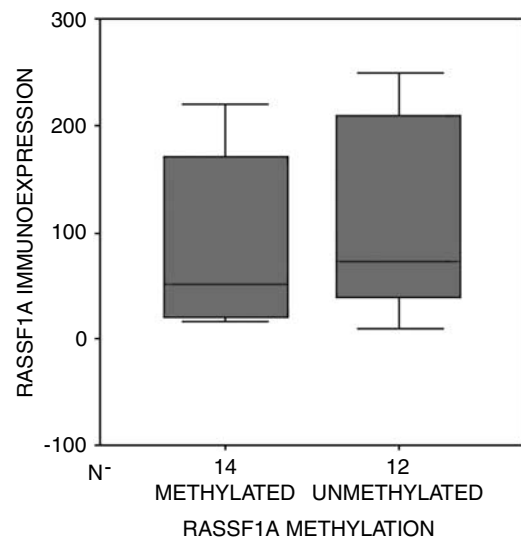


Figure 5 Box-plot comparison of *RASSF1A* immunoexpression and methylation status.

positive *RASSF1A* staining was performed. Interestingly, endometrial carcinoma cases with *RASSF1A*-promoter methylation showed reduced *RASSF1A* immunoexpression (mean HistoScore, 86.09) as compared with unmethylated cases (mean HistoScore, 110.42) (Figure 5), but these differences were not significant ($P=0.274$).

Discussion

Epigenetic inactivation is defined as a change imposed onto the functionality of a gene that does not involve an alteration of its coding sequence. Transcriptional silencing by hypermethylation of

CpG islands in the promoter regions of tumor-suppressor genes is becoming a common phenomenon in carcinogenesis. Although this phenomenon is seen in a wide spectrum of tumors, it is a frequent mechanism of tumor-suppressor gene inactivation in endometrial carcinoma. *De novo* methylation of the *RASSF1A* promoter is one of the most frequent inactivation events detected in human cancer and leads to silencing of *RASSF1A* expression.¹¹ The C-terminus of RASSF1A is homologous to the mammalian RAS-effector protein NORE1 and encodes a RAS-association domain. Three major splice variants, RASSF1A, RASSF1B and RASSF1C, are transcribed from two different CpG islands, separated approximately by 3.5 kb.¹⁹

In the present study, we have assessed the immunohistochemical expression and the presence of promoter hypermethylation of the *RASSF1A* gene in a series of endometrial carcinomas, and detected reduced expression of RASSF1A in 48% of the cases and promoter hypermethylation in 73%. Moreover, there was a statistically significant association between both the phenomena. Although *RASSF1A*-promoter hypermethylation has previously been evaluated in endometrial carcinoma, this is the first study in the literature to demonstrate that it is associated with reduced RASSF1A protein expression, in a large series of endometrial carcinoma and a tissue-microarray approach. In other studies, the frequency of *RASSF1A*-promoter hypermethylation in endometrial carcinoma has been reported to be between 30 and 85%, as found by methylation-specific PCR (MS-PCR).^{20,21} Moreover, in a recent study Kang *et al* found promoter hypermethylation in two of four endometrial carcinoma cell lines tested for promoter hypermethylation (*AN3-CA*, *HEC-1-B*, *KLE*, *RL95-2*). They also demonstrated that endometrial carcinoma cell lines with promoter hypermethylation, treated with 5-aza-2-deoxycytidine, a drug that inhibits DNA methylation, restored *RASSF1A* expression.

RASSF1A protein expression was heterogeneous in our endometrial carcinoma samples. Some cases were RASSF1A positive in many areas and negative in some foci, and also some RASSF1A-negative samples contained a small proportion of cells showing strong staining. In our opinion, the heterogeneous distribution of *RASSF1A*-promoter hypermethylation is the reason that explains why we detected *RASSF1A*-promoter hypermethylation in tumors with positive immunostaining for RASSF1A. This phenomenon can also be found in other types of tumors and with other genes that are silenced by promoter methylation. For example, it has recently been seen that a heterogeneous protein expression of some genes, including *RASSF1A*, is seen in about 70% of malignant melanoma cases, due to intra-tumoral distribution of promoter hypermethylation.²²

During tumorigenesis, activated *RAS* is usually associated with enhanced proliferation, transformation

and cell survival. *RAS* effectors, like RASSF1A, are supposed to have an inhibitory growth signal, which needs to be inactivated during tumorigenesis. Contradictory results between RASSF1A inactivation and *K-RAS* mutation have been obtained using different types of tumors. They were mutually exclusive events in colorectal and pancreatic cancer,^{23,24} but the correlation was not significant in lung cancer.²⁵ Finally, in thyroid cancer, *RASSF1A* methylation is mutually exclusive with mutations in *BRAF*.²⁶ Since *BRAF* mutations are very infrequent in endometrial carcinoma, we focused our interest in the correlation between *RASSF1A* inactivation and *K-RAS* mutations and microsatellite instability.

In our series of endometrial carcinoma, seven out of 10 cases with *K-RAS* mutations were negative for RASSF1A immunorexpression. Kang *et al* found *RASSF1A* hypermethylation to be associated very strongly with microsatellite instability, and inversely correlated with *K-RAS/BRAF* mutations in microsatellite instability-negative endometrial carcinomas. Very recently, Pijnenborg *et al* reported *RASSF1A* hypermethylation in 85% of recurrent endometrial carcinoma, without finding any correlation between *RASSF1A*-methylation status and *K-RAS* or *BRAF* mutations. Interestingly, they demonstrated hypermethylation of *RASSF1A* in samples of endometrial hyperplasias (50%), cyclic (21%) and atrophic endometrium (38%). The authors suggest a role of *RASSF1A* methylation as an early event in endometrial carcinogenesis. However, we have not been able to detect *RASSF1A*-promoter hypermethylation in any of the normal tissue samples corresponding to the 57 cases of endometrial carcinoma. Moreover, the staining pattern that we obtained in normal endometrial tissue was dependent on the menstrual cycle phase, which suggests that in normal endometrium, *RASSF1A* is under the regulation of steroid hormone-signaling pathways.

Microsatellite instability occurs in 25–30% of sporadic endometrial carcinoma, and is characteristic of endometrioid adenocarcinoma. The identification of CpG-island methylation in several genes in colonic and gastric carcinomas with microsatellite instability suggested a relationship between altered methylation and the microsatellite-mutator phenotype. In the present study, we analyzed the possible association between *RASSF1A* expression and the microsatellite instability status in our tissue-microarray approach. We observed a statistically significant association between decreased RASSF1A protein expression and microsatellite instability when we considered as negative those tumors exhibiting RASSF1A HistoScore of less than 50. Our results support the data of Kang and co-workers who also demonstrated that *RASSF1A* methylation was strongly associated with microsatellite instability and *hMLH-1* methylation. Similar results were also obtained in colorectal cancer in which a higher frequency of *RASSF1A*-promoter methylation has

been described in microsatellite instability + colorectal cancers as compared with tumors without the microsatellite instability phenotype. The results suggest that the methylation affecting *RASSF1A* promoter may preferentially occur within the context of microsatellite instability + endometrioid adenocarcinoma.

In the present series, we have observed a higher frequency of *RASSF1A*-promoter methylation in correlation with advanced pathological stage and poorly differentiated carcinomas. In agreement with this result, Jo *et al*,²⁷ found that *RASSF1A* methylation was significantly more frequent in endometrial carcinoma in association with advanced stage (III, IV), lymph node involvement and high histological grade. Moreover, they demonstrated higher incidence of recurrences and lower disease-free survival in patients with *RASSF1A* hypermethylation.

Finally, we noted a nuclear immunostaining of *RASSF1A* in some cases of endometrial carcinoma. Biochemical studies have demonstrated that *RASSF1A* is a microtubule-binding protein, which regulates mitotic progression.²⁸ It has also been shown that *RASSF1A* overexpression induces mitotic arrest at the metaphase with aberrant mitotic cells.²⁸ *RASSF1A* regulates mitosis by inhibiting the anaphase-promoting complex through *Cdc20*, and induces G2–M arrest at the prometaphase. Also, *RASSF1A* prevents degradation of cyclin D1 and delays mitotic progression.²⁹ *RASSF1A* is also able to affect other cyclins, such as cyclin D3 or cyclin A, through interaction with transcription factor p120^{E4F}, which induces cell-cycle arrest and provides a mechanistic link with other known tumor-suppressor genes such as *p14^{ARF}*, *Rb* and *p53* that are known to interact with p120^{E4F}.³⁰ To evaluate the possible relationship between *RASSF1A* and the cell-cycle machinery, we checked the relation between *RASSF1A* expression with cyclin D1 expression or proliferation index (Ki67 staining), but found no significant association.

In summary, we report, for the first time, reduced protein expression of *RASSF1A* in endometrial carcinoma using tissue-microarray technology, and reduced *RASSF1A* immunorexpression in cases with promoter hypermethylation. Also, we demonstrate higher frequency of *RASSF1A*-promoter methylation in advanced endometrial carcinoma stage, suggesting that *RASSF1A* could be a useful indicator of tumor aggressiveness in endometrial carcinoma patients.

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Conflict of interest

The authors state no conflict of interest.

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