Surface Expression of Minichromosome Maintenance Proteins Provides a Novel Method for Detecting Patients at Risk for Developing Adenocarcinoma in Barrett’s Esophagus

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ABSTRACT

Purpose: The purpose is to determine whether a novel cell cycle marker, minichromosome maintenance protein 2 (Mcm2), predicted esophageal adenocarcinoma (AC) risk in Barrett’s esophagus (BE) and whether this could be used in combination with a surface sampling method.

Experimental Design: Archival specimens [30 normal squamous esophagus (NE), 20 gastric antrum (GA), 13 duodenum (D2), 62 BE ± dysplasia, and 16 (AC)] were stained for Mcm2. Sequential biopsies from nine patients who developed AC during surveillance were compared with 18 matched controls who did not progress. Prospective endoscopic cytological brushings (61 NE, 90 BE ± dysplasia, and 11 AC) were scored as Mcm2 positive or negative.

Results: Mcm2 was not expressed on the luminal surface of NE, GA, and D2. In BE, the percentage of surface cells expressing Mcm2 correlated highly with the degree of dysplasia ($P < 0.0001$). In patients who developed AC, biopsies before dysplasia had higher Mcm2 expression than the matched control patients (mean, 28.4 and 3.4% positive cells, respectively, $P < 0.0001$). In the prospective cohort, the histopathological diagnosis of dysplasia or AC and the Mcm2-positive brushings were concordant in 91% of the patients ($P < 0.0005$) and the results correlated with the frequency of cases with surface expression of Mcm2 in the retrospective study ($P < 0.0001$).

Conclusions: Surface expression of Mcm2 can be used to detect dysplasia and AC, as well as patients with BE at risk for subsequent development of dysplasia and AC. A brushing technique combined with Mcm2 staining has the potential to be exploited in surveillance and screening protocols.

INTRODUCTION

The incidence of esophageal AC$^2$ has increased by 350% in the Western world over the last 30 years (1). Most patients present with dysphagia, and despite advances in multimodal treatment, the 5-year survival rates of symptomatic esophageal carcinomas remain <10% (2, 3). Because the major determinant of outcome is the stage of the cancer at presentation (4), early detection of esophageal cancer is essential to significantly improve survival rates. The identification of a multistage process of cancer development, akin to that described in colon carcinogenesis, provides the rationale for endoscopic surveillance in patients with diagnosed BE (5–8). Several studies have shown a significant improvement in the 5-year survival of patients with surveillance-detected AC (9, 10).

The clinical effectiveness of surveillance for patients with BE is currently hampered by the inadequate surveillance methods used. These are prone to sampling bias and the highly subjective histopathological diagnosis of dysplasia, with high inter- and intraobserver variability (11). Furthermore, we know that only a minority of patients with BE are diagnosed (12). As a result, there has been a suggestion that screening endoscopy should be performed in people with heartburn who are most at risk for developing BE and AC (13). However, the current dependence on endoscopic techniques means that screening is unlikely to be cost effective or feasible at the current time.

In an attempt to improve current surveillance methods, investigators have tried to identify biomarkers to stratify BE patients according to their cancer risk (7, 14). No single oncogene or tumor suppressor gene has been shown to be sensitive enough. Therefore, an alternative approach would be to use markers for the final common pathway after genetic instability, which is the loss of proliferative control. Until recently, assessment of cell proliferation on fixed tissue has relied on markers with disadvantages such as Ki67 and the proliferating cell nuclear antigen (15, 16). Now with an increased understanding of the proteins controlling the cell cycle, alternative markers are becoming available (17). The MCM protein complex binds to the origins of replication to ensure a single round of DNA replication. MCMs are expressed in all cycling cells throughout the cell cycle and are degraded when mammalian cells exit the...
cell cycle into quiescent, differentiated, and senescent states (18, 19). As a result of the sensitivity of antibodies to MCMs as markers of cell cycle state and because of the aberrant entry into the cell cycle in dysplasia and malignancy, MCM proteins have been proposed as candidate markers for cancer screening, surveillance, and as prognostic markers (20–24). Williams et al. (23) demonstrated that Mcm2 and Mcm5 surface positivity correlated with the severity of dysplasia. These observations have led to the rationale for using surface sampling techniques in combination with MCM staining to detect dysplasia and malignancy (24). This approach has yielded promising results for cervical and colon cancer screening (22, 23).

If cell cycle markers could be combined with a simple surface sampling technique, then this may provide a significant advance for identifying individuals at risk for the development of esophageal AC. This study had three main aims. Firstly, to determine the magnitude and pattern of expression of Mcm2 (24), as a prototype MCM protein, in histological samples from patients undergoing upper gastrointestinal endoscopy. Secondly, to determine whether aberrant surface expression of Mcm2 is predictive of cancer development using a retrospective case-control study. Thirdly, to determine whether detection of Mcm2 could be combined with a brushing technique and liquid-based cytology as a novel strategy for predicting those BE patients at risk for the development of AC.

PATIENTS AND METHODS

Patient and Tissue Collection. Approval was obtained from the Local Research Ethics Committee for the retrospective and prospective parts of this study.

Retrospective Cross-Sectional Study. Archival blocks were obtained from patients who had attended Addenbrooke’s Hospital (Cambridge, United Kingdom). We studied 37 BE negative for dysplasia, 28 for LGD, 12 for HGD, and 17 with AC. Sections of normal tissue from non-BE and BE patients’ NE (n = 30), the second part of the D2 (n = 13), and GA (n = 20) were used as control.

Retrospective Case-Control Study. Serial biopsies from nine patients whose AC or HGD was detected through a surveillance program (followed up for a median of 6 years, range from 3 to 13 years) were compared with those from 18 controls, matched for age and length of follow-up (with a median of 5 years, range from 3 to 7 years) who did not progress. The cancer patients had at least two endoscopies before the development of HGD, and they all progressed through the no dysplasia-LGD-HGD sequence. The control cases had no more than a diagnosis of focal LGD in one of their surveillance endoscopies. Hence, control patients with multifocal LGD at any time point of their surveillance program were excluded from the study.

Prospective Study. A total of 162 patients attending Addenbrooke’s Hospital for an endoscopy for either heartburn symptoms, routine BE surveillance, or for assessment of dysplasia/AC as part of a tertiary referral service was recruited for esophageal brushings before biopsy (61 NE, 90 BE ± dysplasia, and 11 AC).

All patients with BE, in either the retrospective or prospective parts of the study, had an endoscopically visible columnar-lined segment and a histopathological diagnosis of specialized intestinal metaplasia. Esophagitis, occurring either in isolation or in conjunction with BE, was graded according to the Los Angeles classification with histopathological confirmation according to the established criteria (25). All patients had diagnostic biopsies taken 2 cm above the GEJ and in addition patients with BE had surveillance biopsies taken (each quadrant every 2 cm; Ref. 26).

Cytology Brushings. For all patients recruited prospectively, brush cytology was performed 2 cm above the GEJ using a disposable brush (Boston Scientific, Watertown, MA) under direct endoscopic vision. The brush was then put into PreservCyt solution (Cytyc Corporation, Boxborough MA), and from this, a monolayer of cells was obtained using a ThinPrep 2000 (Cytyc Corporation). One slide was stained conventionally with a Papanicolaou stain and analyzed for cytological abnormalities by an expert cytopathologist (M. O.). The second slide was stained for Mcm2. Only a subset of the samples (61 NE, 21 BE, 37 LGDs, 3 HGD, and 7 AC) was analyzed for routine cytology as it quickly became apparent that the results obtained showed variable correlation with the histopathological diagnosis (see “Results”).

Immunostaining. Tissue sections were deparaffinized in xylene and rehydrated through alcohol solutions, water, and finally with TBS-Tween. Washing using TBS-Tween was performed between each step. An antigen-retrieval step was performed by pressure cooking samples for 3 min in 0.01 M Tris-sodium citrate buffer at pH 6.0. Staining was performed using the Dako autostainer (DakoCytomation Ltd., Ely, United Kingdom) and the staining kit Dako ChemMate for increased reproducibility. Blocking of nonspecific binding was performed using 10% normal goat serum (v:v) and 10% BSA (w:v) for 30 min at room temperature. The samples were incubated with monoclonal Mcm2 antibody (21, 22) at 1/10 or MIB-1 (anti-Ki67 antibody; DakoCytomation Ltd.) at 1/100 dilution in antibody diluent (DakoCytomation Ltd.) for 1 h at room temperature. Endogenous peroxidase activity was blocked with peroxidase-blocking solution (DakoCytomation Ltd.) for 5 min. The secondary and the peroxidase-linked antibodies were each incubated for 30 min at room temperature followed by 3,3’-diaminobenzidine substrate (DakoCytomation Ltd.) for 10 min. Sections were counterstained with hematoxylin. A negative control was performed by omission of the primary antibody.

For the cytological samples, the fixative was removed by a wash in 100% methanol for 5 min, and the cells were hydrated in TBS for 5 min and then washed in TBS-Tween. The cells were permeabilized in 4 mM sodium deoxycholate in TBS. The staining procedure was the same as for paraffin sections except that the primary antibody was applied for 2 h. A negative control was done by omission of the primary antibody using a monolayer slide made from a BE AC cell line SEG-1 (gift from Dr. David Beer, University of Michigan Medical School).

Scoring Immunostaining. The epithelial surface was defined as the superficial four layers of cells in the squamous esophagus and the most superficial layer of columnar cells for the glandular tissues. Biopsies without surface epithelium were excluded from the analysis. All of the surface cells per biopsy were counted up to a maximum of 600 to determine
the frequency of Mcm2 expression. Only cells with diffuse nuclear staining were considered as positive. The immunopositive epithelial cells were expressed as a percentage of the total number of epithelial cells counted. The brushings were scored as either positive or negative for Mcm2 staining. Any positive cell was considered as abnormal (see "Results," Figs. 1 and 2), therefore, the brushing was positive if only one positive cell was present.

**Statistical Analysis.** The frequency of Mcm2 staining are expressed as a mean ± SE. The Jonckheere-Terpstra test (27) and the \( \chi^2 \) test for trend were used to assess the correlation between Mcm2 expression and the degree of dysplasia. The Jonckheere-Terpstra test is a nonparametric statistic test, testing for a shift in ordered distributions (Mcm2 surface expression) when stratified by ordered categories (NE, BE, LGD, HGD, and AC). The Mann-Whitney test was used to identify specific differences between groups. In all cases, \( P < 0.05 \) was required for significance (28, 29).

**RESULTS**

**Mcm2 Surface Staining Correlates with the Grade of Dysplasia.** In normal control tissues (NE, GA, and D2), Mcm2 was not expressed at the luminal surface (Figs. 1, A–C, and 2). In NE, expression was confined to the parabasal and suprabasal compartments of the epithelium (Fig. 1A). In GA and D2, expression was restricted to the glands and lower crypts (Fig. 1, B–G; \( \times 100 \)), and the top panel demonstrates a high magnification view of the epithelial surface from the corresponding section (\( \times 400 \)).

For comparison, in BE the mean surface expression of Mcm2 and Ki67 was respectively 6.8 and 4.9% of epithelial cells in nondysplastic samples, 32.9 and 20.3% in LGD, 49.7 and 34.2% in HGD, and 66.5 and 59.9% in the AC. Hence, as expected, the Mcm2 antibody was more sensitive than Ki67, although this was difference was not statistically significant.
Interestingly, Mcm2 staining in patients with nondysplastic BE was dichotomous (unlike for Ki67, data not shown). Forty-four percent of BE samples were positive for Mcm2 surface expression with a mean of 15.5% positive cells/biopsy and 56% of patients were negative for surface Mcm2 expression (Fig. 2).

Because normal epithelial tissues such as GA and D2 do not express surface Mcm2 and because Mcm2 surface expression correlates with dysplasia, we hypothesized that the population of nondysplastic BE patients with Mcm2 surface-positive staining might be more likely to progress down the dysplastic pathway than those who have no surface expression. This hypothesis formed the basis of the case-control study.

**Mcm2 Surface Expression Allows Detection of Patients at Risk for Progression to AC.** For each of the nine cases of AC, Mcm2 surface expression in biopsies before the diagnosis of dysplasia was higher than for the matched controls (Fig. 3). The mean expression of Mcm2 was 28.4% of total surface epithelial cells for BE biopsies in the group destined to develop AC and 3.4% for the control group (Fig. 3). The level of Mcm2 expression was maintained throughout the length of follow-up for the control group, but the expression levels in the AC group increased with time as the degree of dysplasia increased (Fig. 3). The correlation between the surface expression of Mcm2 and the degree of mucosal inflammation (data not shown, $P = 1$), and the concordance between histopathology and cytology for dysplasia was 48%, in keeping with other studies (30), compared with 91% for Mcm2 staining.

**DISCUSSION**

In this study, we have demonstrated that aberrant surface expression of Mcm2 increases as patients progress along the metaplasia-dysplasia-carcinoma sequence of esophageal AC. The case-control study suggests that patients with surface expression of Mcm2 in metaplastic BE may be more at risk for malignant progression. Because Mcm2 expression occurs on the surface of potentially premalignant tissues, but not on the surface of normal gastrointestinal tissues, a surface-sampling brush methodology may be applicable for detection of dysplasia and cases of BE destined to progress to AC. We have demonstrated comparable frequency of Mcm2 expression in endoscopic brushings versus multiple biopsies.

A recent publication demonstrated that Mcm2 and Mcm5 expression was increased in dysplastic BE (31). Interestingly, the pattern of Mcm2 and Mcm5 staining in histopathological sections of BE resembled that of gastric type mucosa with maximal expression in the crypts. However, it is not clear whether the presence of intestinal metaplasia was used in the study as part of the diagnostic criteria for BE. Here, we have restricted the definition of BE to the intestinal subtype containing goblet cells, which is also the type that confers the highest
risk for malignancy (32), and the expression pattern of Mcm2 in BE resembles more that of D2 with a maximum expression in the lower crypts.

Cytology alone is likely to be a poor screening or surveillance test for BE and associated dysplasia, as shown by our own results (Table 1), and as demonstrated in previous studies (33). The lack of cytological detection of intestinal metaplasia (28%) may be partly because of the variable distribution of goblet cells in BE. The assessment of LGD was difficult in some cases because of marked reactive changes attributable to inflammation. The air drying and spreading artifacts were partially overcome by using liquid-based cytology as opposed to conventional-based cytological techniques. Cytology was highly sensitive for the detection of HGD/AC, with a failure of detection only occurring because of inadequate material for assessment (2 of 7 AC cases missed).

However, analysis of Mcm2 expression in conjunction with cytology increases the diagnostic yield significantly. Given that the positive Mcm2 staining is localized to the nucleus, it is easy to distinguish true positive columnar cells from background staining in the liquid-based cytology monolayer (Fig. 5). The shape, size, and low background seen in squamous cells makes them easy to distinguish from columnar cells (Fig. 5). In our prospective study, any cells obtained from the brushings with nuclear staining were counted as positive. This makes scoring positive and negative brushings very simple and hence appealing for clinical practice. Furthermore, the exfoliative cytology technique used in this study enables a greater surface area of esophageal mucosa to be sampled compared with a conventional biopsy series. In the future, with the advent of newer cytological sampling devices, it should be possible to sample an even greater surface area of mucosa and to apply these techniques nonendoscopically (34). This has the potential to change radically the cost-effectiveness of esophageal screening and surveillance programs.

None of the biopsies from normal squamous esophagus had positive Mcm2 expression on the surface. In contrast, brushings from 13% of NE patients were Mcm2 positive. Half of these samples were positive squamous cells, and there was no correlation between the degree of inflammation and these results ($P = 1$). It is possible that the positive squamous cells originate from the basal layers and were brushed away from the epithelium as a result of an excessively harsh sampling technique. The other half of the positive brushings in NE patients represented positive glandular cells (two patients had diagnosed gastritis, one had intestinal metaplasia, and the last had a normal endos-
Table 1  A. routine cytology and Mcm2-stained brushings in the diagnosis of BE and associated dysplasia; and B. Mcm2 surface expression in the samples from the retrospective study

<table>
<thead>
<tr>
<th>Histopathological diagnosis in biopsy samples</th>
<th>Cytological diagnosis of dysplasia</th>
<th>Mcm2-positive brushings</th>
<th>Mcm2 surface expression</th>
</tr>
</thead>
<tbody>
<tr>
<td>NE</td>
<td>15% (9/61)</td>
<td>13% (8/61)</td>
<td>0% (0/30)</td>
</tr>
<tr>
<td>BE</td>
<td>28% (6/21)</td>
<td>42% (10/45)</td>
<td>44% (15/34)</td>
</tr>
<tr>
<td>LGD</td>
<td>43% (16/37)</td>
<td>88% (37/42)</td>
<td>85% (17/20)</td>
</tr>
<tr>
<td>HGD</td>
<td>100% (3/3)</td>
<td>100% (3/3)</td>
<td>88% (7/8)</td>
</tr>
<tr>
<td>AC</td>
<td>71% (5/7)</td>
<td>100% (11/11)</td>
<td>100% (16/16)</td>
</tr>
</tbody>
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Presumably, the positive glandular cells had originated from either the gastric cardia or from endoscopically undiagnosed ultrashort BE as 15% of biopsies from the GEJ of asymptomatic patients have been shown to have small foci of intestinal metaplasia (35).

It is also of note that 44% of nondysplastic BE patients had some degree of surface Mcm2 expression present in both the biopsy and the brushing samples. It is generally believed that there is a 0.5–1% risk/year of BE patients progressing to AC (36), and therefore, only a minority of the 44% of patients with Mcm2 surface expression are likely to progress. However, in the case-control study, we demonstrated that the patients who did progress to AC had significantly increased surface expression of Mcm2 at their initial endoscopy, before the development of dysplasia (mean, 28.4% positive cells versus 3.4% for controls, P < 0.0001; Fig. 3). It is interesting to note that of the 44% of nondysplastic BE patients with surface Mcm2 expression, the mean number of positive cells was 15.5%, and only three patients had >25% positive cells. Hence, whether a quantitative assessment of surface Mcm2 expression and the change in the level of Mcm2 expression over time is predictive of cancer development may be worthy of additional investigation. In the future, it is also possible that a combination of Mcm2 with other biomarkers may enable patients to be stratified additionally for risk of progression, although the histopathological diagnosis will remain the gold standard for dysplasia.

A large proportion of patients with BE in our prospective study had LGD. This reflects the tertiary referral of patients with dysplasia in BE in our practice. In the future, there needs to be a prospective study of Mcm2 staining applied to a nonselected surveillance cohort with appropriate follow-up. In addition, with the advent of nonendoscopic cytological sampling devices, it would be valuable to apply our immunocytochemical technique to a cohort of heartburn patients who are subsequently endoscoped. For example, in parts of mainland China, nonendoscopic cytological sampling of the esophagus (balloon cytology) has been used for decades as a screening test for squamous esophageal cancer (37, 38). During the current study, five patients with squamous esophageal carcinoma attended for an endoscopy and had brushings taken. All five of these patients had positive Mcm2 surface expression, although they were excluded from subsequent analysis in view of the squamous cell histology.

In view of the alarming increase in the frequency of AC of the esophagus, inexpensive and effective tests for detection of patients at risk are badly needed. The detection of MCM proteins in samples from the esophageal surface may offer a diagnostic advance.

ACKNOWLEDGMENTS

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REFERENCES


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