Aberrant expression of minichromosome maintenance proteins 2 and 5, and Ki-67 in dysplastic squamous oesophageal epithelium and Barrett’s mucosa

J J Going, W N Keith, L Neilson, K Stoever, R C Stuart, G H Williams

Background: Minichromosome maintenance (Mcm) proteins are essential for eukaryotic DNA replication, and their expression implies potential for cell proliferation. Expression is dysregulated in dysplastic states but data for oesophageal squamous mucosa and Barrett’s mucosa have not been published.

Aim: To test the hypothesis that Mcm proteins are downregulated together with the proliferation marker Ki-67 in differentiating epithelial compartments of non-dysplastic squamous and Barrett’s epithelium, and that this process does not occur in dysplastic mucosae.

Methods and cases: Forty-five patients with Barrett’s oesophagus included 20 with glandular dysplasia (10 low grade, eight high grade, two both, and four with invasive adenocarcinoma). Twenty-five other patients included 12 with oesophageal squamous dysplasia (three low grade, six high grade, three both, and four with invasive squamous carcinoma). Formalin fixed paraffin-embedded tissue sections from biopsy series and resections were immunostained using antibodies to Mcm2, Mcm5, and Ki-67. Percentage of nuclei positive for Mcm2, Mcm5, and Ki-67 was estimated and scored from 0 to 6 as: 0, none +; 1, <10%+; 2, 10–30%+; 3, 30–70%+; 4, 70–90%+; 5, >90%+; 6, all+. Four separate epithelial strata were scored: in squamous epithelium the basal layer and thirds to the surface, in Barrett’s mucosa the luminal surface, upper and lower crypt, and deep glands.

Results: In non-dysplastic squamous epithelium and Barrett’s mucosa, high level expression of Mcm2, Mcm5, and Ki-67 proteins was largely confined to the proliferative compartments and downregulated in differentiated compartments. Expression persisted up to the mucosal surface in dysplastic squamous epithelium and dysplastic Barrett’s mucosa.

Conclusions: Persistent expression of Mcm2 and Mcm5, and Ki-67 proteins in luminal compartments of dysplastic oesophageal squamous epithelium and dysplastic Barrett’s mucosa may be diagnostic markers and imply disruption of cell cycle control and differentiation in these dysplastic epithelia.

Lethality of symptomatic oesophageal cancer motivates screening4–6 for earlier disease, treatable by surgical resection or mucosal ablation by laser,7 argon beam,8 or photodynamic therapy.9 Population screening for squamous oesophageal carcinoma is only practised where the incidence is high (for example, Japan and China). Dysplasia and carcinoma surveillance by endoscopy and biopsy of Barrett’s patients in Western populations is also undertaken, but with uncertain benefit.10 It is a substantial commitment, rigorous definition of Barrett’s dysplasia is difficult, and inter- and intra-pathologist agreement is imperfect.11–13 Even four quadrant sampling with jumbo biopsy forceps does not guarantee that all significant dysplasia will be detected,14 and the natural history of oesophageal glandular dysplasia is obscure.

Abnormal proliferation and differentiation typify epithelial dysplasia. Normal oesophageal squamous epithelial cells divide slowly in the basal layer, proliferate suprabasally, and mature towards the luminal surface.15 In Barrett’s mucosa, despite its partially intestinal phenotype, proliferation and differentiation patterns resemble gastric mucosa, with maximal proliferation in a crypt zone beneath the mucosal surface16 and differentiation into deep glands and characteristic cell populations on the mucosal surface (in normal small intestine, stem cells in the crypts of Lieberkühn feed a proliferative compartment from which differentiating enterocytes and goblet cells migrate to the villi17 while Paneth cells migrate basally).

Proliferation and differentiation compartments break down in dysplastic epithelia. “Dysplastic” cells adjacent to an invasive carcinoma probably represent the neoplastic clone from which the carcinoma emerged. Dysplasia alone implies an increased cancer risk, and motivates eradication or increased intensity of surveillance. Difficulty in reliably recognising and grading dysplasia is therefore therapeutically relevant, and improved methods for doing so are desirable.

In eukaryotic cells, initiation of DNA synthesis at specific sites (origin firing) is tightly restricted to permit duplication of the genome once only per cell cycle.18 Initiation factors including the origin recognition complex, Cdc6,19 and minichromosome maintenance (Mcm) proteins which assemble during G1 into pre-replicative complexes (pre-RCs) at replication origins to establish competence for DNA replication in S phase. Activated Cdc7/Dbf4 kinase and S phase promoting cyclin dependent kinases trigger unwinding of replication origins and establish bidirectional replication forks, and disassembly of pre-RCs during replication prevents reinitiation of DNA replication within a single cycle.20

When mammalian cells exit the cell cycle into quiescent, differentiated, and senescent states, the Cdc6 and Mcm components of the pre-RCs are downregulated,21,22 and dysregulated expression of these proteins is characteristic of both dysplastic cervical squamous epithelium23 and urothelium.24 It seemed appropriate therefore to evaluate their potential as markers of dysplasia and dysregulated cell cycle control in normal oesophageal squamous epithelium, Barrett’s mucosa.

Abbreviations: Mcm proteins, minichromosome maintenance proteins; pre-RCs, pre-replicative complexes; mAb, monoclonal antibody.
without dysplasia, and in their dysplastic and neoplastic counterparts.

MATERIALS AND METHODS

Production of antibodies

Anti-Mcm2 monoclonal antibody (mAb) (mouse IgG1 isotype) was raised against a fragment of human Mcm2 (amino acids 725–888; BM28, Transduction Laboratories, Lexington, Kentucky, USA). The specificity of the anti-Mcm2 mAb was established by immunoblot, immunofluorescence, and immunoprecipitation assays. Rabbit polyclonal antibodies were raised against a fragment of human Mcm5 (amino acids 372–590) and purified by affinity chromatography over a column prepared by linking the immunogen to Sulfolink Gel (Pierce, Rockford, Illinois, USA). Specificity of anti-Mcm2 mAb and anti-Mcm5 polyclonal antibody was established by immunoblot, immunofluorescence, and immunoprecipitation assays. The anti-Ki-67 mAb MIB1 was supplied by Dako (Ely, Cambridge, UK).

Immunohistochemistry

Preliminary testing of six different antibodies at different dilutions and antigen retrieval schedules including enzymatic digestion and microwave heating in citrate and EDTA buffer was performed on formalin fixed paraffin embedded tissue sections. Good results were obtained with antibodies against Mcm proteins 2 and 5 using EDTA buffer and microwave heating antigen retrieval. Both Mcm antibodies were used diluted 1:4000 and conventional three stage streptavidin/biotin and peroxidase with diaminobenzidine/H2O2 detection. Antibody MIB1 was applied diluted 1:100 following microwave antigen retrieval and detected as for Mcm2/Mcm5.

Cases

Seventy patients were studied. Twenty five patients without Barrett’s oesophagus included 13 with no squamous dysplasia, three with low grade squamous dysplasia, six with high grade squamous dysplasia, and three with both. Four patients in this group also had invasive squamous carcinoma. Forty five patients with Barrett’s oesophagus included 25 without dysplasia, 10 with low grade glandular dysplasia, eight with high grade dysplasia, and two with both. Four patients in this group also had invasive adenocarcinoma.

Patients were from cohorts undergoing diagnostic endoscopy for upper gastrointestinal symptoms, enrolled in a yearly surveillance of Barrett’s oesophagus, or having surgical resection of oesophageal carcinoma. Dysplastic changes were...
assessed on haematoxylin and eosin stained sections of paraffin embedded endoscopic biopsies and tissue blocks from resection specimens which were chosen from pathology department archives to represent a range of morphologies from normal (non-dysplastic) oesophageal squamous epithelium through low and high grade squamous dysplasia to invasive squamous carcinoma. Similarly, examples of Barrett’s mucosa without dysplasia, low grade and high grade dysplasia in Barrett’s mucosa, and invasive Barrett’s adenocarcinoma were selected for study, and 4 \mu m sections were immuno-stained as described.

Scoring immunocytochemistry
A semiquantitative scoring scheme was designed to describe the immunostaining observed. Cell nuclei were positive or negative for Mcm2, Mcm5, or Ki-67. All scoring was done by one specialist upper gastrointestinal pathologist (JJG). Within each separate mucosal compartment, the estimated percentage of positive cells was allocated to scoring bands as follows: 0, none+; 1, <10%+; 2, 10–30%+; 3, 30–70%+; 4, 70–90%+; 5, >90%+; and 6, all+. Four compartments were recognised in oesophageal squamous epithelium: the most basal single layer of cells, and the thickness of the epithelium above that divided into parabasal, middle, and luminal thirds. In Barrett’s mucosa, four strata again were defined: the surface epithelium between “crypts”, the underlying “crypts” or “pits” divided into upper and lower halves, and the deepest layer, a differentiated glandular zone. These compartments correspond to those defined by Lauwers and colleagues\textsuperscript{18} in their study of cell proliferation in Barrett’s mucosa.

Reproducibility of scoring and statistical analysis
Reproducibility of scoring was evaluated by “blind” re-scoring by JJG of all sections stained for Mcm2 after six months. Weighted kappa \( \kappa_w \textsuperscript{19} \) was calculated for duplicate scores from 562 separate cellular populations; \( \kappa_w = 0.65 \) implied acceptable agreement. Univariate statistical significance of differences between immunostaining was tested using the non-parametric Mann-Whitney test for two independent samples, two tailed, with correction for ties. Kappa and Mann-Whitney calculations were performed in Analyse-It for Microsoft Excel (Analyse-It Software, Leeds, UK).

RESULTS
Immunostaining with the Mcm2 antibody yielded predominantly nuclear staining. The Mcm5 antibody stained nuclei but also cell membranes in glandular mucosae and tumours. Qualitatively, nuclear staining was similar with the two antibodies. Ki-67 staining was purely nuclear. In non-dysplastic squamous epithelium and Barrett’s mucosa, strong Mcm2, Mcm5, and Ki-67 staining of most to
all nuclei was present in the expected proliferative transit compartment—that is, the suprabasal compartment of squamous epithelium—and in the lower crypt compartment of Barrett’s mucosa. In differentiated compartments—that is, the surface of squamous epithelium and Barrett’s mucosa—and in the small differentiaed deep glands of Barrett’s mucosa, expression was downregulated. In dysplastic squamous epithelium and dysplastic Barrett’s mucosa there was persistence of Mcm2, Mcm5, and Ki-67 expression towards the surface, and in the same case Mcm2 downregulation occurs in invasive squamous carcinoma cells are Mcm2 positive but viable Mcm2 negative carcinoma cells are present within cell nests with morphological features of differentiation (keratinisation).

Barrett’s mucosa, and disruption of this highly organised spatial arrangement in premalignant dysplasia. These disturbances are relevant to the identification of dysplasia in oesophageal squamous epithelium and Barrett’s mucosa, both of which are problematic in individuals and populations. Squamous oesophageal cancer is a target for screening in Far Eastern populations. Barrett’s oesophagus and Barrett’s cancer are relatively common in the West. Patients with Barrett’s oesophagus may be subjected to relatively frequent endoscopy and biopsy (for example, yearly). A sensitive and specific test for dysplasia might allow Barrett’s patients to be screened for dysplasia and divided into a cohort without dysplasia, at low risk of oesophageal adenocarcinoma, for whom less intensive follow up would be safe, and a higher risk group, with dysplasia, for whom more frequent endoscopic and biopsy surveillance could be appropriate. A sensitive test for early dysplastic changes in endoscopic biopsies would be useful to the pathologist although to date various tests that have been proposed have been disappointing in practice.

Abnormal expression of Mcm5 protein in dyskaryotic cervical smears is associated with cervical intraepithelial neoplasia, and immunocytochemistry for Mcm5 protein facilitates detection of dyskaryotic cells in such smears, which may otherwise be a problem if they are present in small numbers.17 Similarly, biochemical assay of these proteins in urine is a sensitive and specific test for the presence of uterine epithelial neoplasia.17 These examples and the present

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<th>Table 1</th>
<th>p values by Mann-Whitney testing of the differences between groups</th>
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<td>Ki67</td>
<td>Mcm5</td>
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<tr>
<td>Squamous mucosa</td>
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<td>Surface</td>
<td>Normal v LGD</td>
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<td>Parabasal</td>
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<td>Basal</td>
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LGD, low grade dysplasia; HGD, high grade dysplasia.
in invasive carcinomas, molecular events associated with differentiation appear capable of switching off Mcm protein expression. These findings support the concept of mutual antagonism between the cellular circuits controlling differentiation and proliferation. As this would usually be understood to prevent further cell division, such events may have therapeutic relevance.

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REFERENCES


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