Differences in activity and phosphorylation of MAPK enzymes in esophageal squamous cells of GERD patients with and without Barrett’s esophagus

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GERD patients with and without Barrett’s esophagus.

Summary

We hypothesized that, in esophageal squamous epithelial cells, there are differences among individuals in the signal transduction pathways activated by acid reflux that might underlie the development of Barrett’s esophagus. To explore that hypothesis, we immortalized nonneoplastic, esophageal squamous cells from patients with gastroesophageal reflux disease (GERD) with (NES-B3T) and without (NES-G2T) Barrett’s esophagus and used those cells to study acid effects on MAPK proteins. During endoscopy in patients with GERD with and without Barrett’s esophagus, we took biopsy specimens from the distal squa

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We have hypothesized that, in esophageal squamous epithelial cells, there are differences among individuals in signal transduction pathways activated by acid reflux that might underlie the development of Barrett’s esophagus. In an earlier report, we described the effects of acid perfusion on the ERK1/2 pathways in the esophageal squamous epithelium of patients with GERD with and without Barrett’s esophagus (25). We found significant differences between those patient groups in baseline and acid-induced levels of ERK1/2 in esophageal squamous epithelium (25). The present study was designed to explore the mechanisms underlying those differences. To do so, we have immortalized nonneoplastic esophageal squamous epithelial cells from patients with GERD through the stable incorporation of human telomerase reverse transcriptase (hTERT). Using those cell lines and biopsies of esophageal squamous epithelium taken from patients with GERD with and without Barrett’s esophagus during endoscopic examinations, we investigated the effects of acid exposure on the activity and phosphorylation of MEK1/2 and the expression levels of MKP-1 both in vitro and in vivo.

MATERIALS AND METHODS

Isolation and culture of esophageal squamous cells. This study was approved by the institutional review board on human studies at the Dallas VA Medical Center. During endoscopic examinations, samples of the squamous epithelium of the distal esophagus were taken from patients with GERD with and without long-segment Barrett’s esophagus (defined as ≥3 cm of specialized intestinal metaplasia in the distal esophagus) with the use of jumbo biopsy forceps (Olympus FB-50K-1). Biopsy tissues were placed in cold Hank’s salt solution containing 1% antibiotic-antimycotic (both from Invitrogen, Carlsbad, CA) and processed within 2 h of collection. Six biopsies were placed into a conical tube containing trypsin (Invitrogen) at 37°C for 30 min. Following trypsinization, biopsies were dispersed into a single-cell suspension and cocultivated with mouse embryonic Albino Swiss 3T3 [American Type Culture Center (ATCC), CCL-92] fibroblast cells (1–3 × 10⁶ cells/100-mm plate) pretreated with mitomycin C (10 µg/ml for 2 h), in a supplemented 3:1 mixture of DMEM/Ham’s F12 medium (Invitrogen) as previously described (13, 18). For subculturing, fibroblast feeder cells were removed with EDTA washes (0.02% in 1× PBS, pH 8.0; Sigma, St. Louis, MO) before trypsinization of the epithelial cells. Squamous cells were resuspended in fresh medium containing 5% trypsin inhibitor (Sigma), reseeded at a 1:8 split ratio, and grown at 37°C in 5% CO₂ with fresh media changes three times weekly. The number of population doublings (PD) per passage was calculated as log [(number of cells harvested/number of cells plated)/log2].

Retroviral vectors and transductions. Esophageal squamous cells were retrovirally infected with hTERT using the Cre-lox recombination system as previously described by our laboratory (8). Cells at ~30–50% confluence were infected twice with the retroviral vector containing hTERT flanked by lox-P sites (hTERTloxP) in the presence of 4 µg/ml of Polybrene (Sigma) at PDs 10–16. Esophageal squamous epithelial cell lines derived from patients who had GERD with Barrett’s esophagus (NES-B3T) and without Barrett’s esophagus (NES-G2T) containing the hTERTloxP vector were selected with 500 ng/ml puromycin for 7–9 days (Fig. 1).

Telomerase activity. Telomerase activity was measured in cultured cells before and after the introduction of hTERT with the TRAP-eze Telomerase Detection kit (Intergen, Burlington, MA) as previously described (7, 8). Telomerase-expressing cervical cancer-derived HeLa cells were used as positive controls. Lysis buffer was used as a negative control.

Contact inhibition and anchorage-dependent cell growth assay. 50,000 telomerase-immortalized NES-B3T and 120,000 telomerase-immortalized NES-G2T cells were plated in six-well plates and harvested at multiple time points ranging from 1 to 16 days. Cells were harvested with 0.05% trypsin and counted by Coulter counter.

Fig. 1. Telomerase-immortalized esophageal squamous epithelial cells in culture. Low (4×) and high (10×) power magnifications of esophageal squamous epithelial cells (white arrows) grown with ATCC Albino Swiss 3T3 fibroblast feeder cells (black arrows). A: NES-G2T squamous cells from a patient with gastroesophageal reflux disease (GERD) without Barrett’s esophagus. B: NES-B3T squamous cells from a patient with GERD with Barrett’s esophagus.
An in vitro marker for neoplastic cells is the ability to grow in soft agar. A 12-well soft agar plate was created with two layers: the bottom layer had 0.5% (wt/vol) of Noble agar (Sigma, A5431) in DMEM/F12 supplemented with 20% FBS (Atlanta Biologicals, Norcross, GA). The top layer consisted of cells with 0.3% agar in DMEM/F12 and 20% FBS. 5,000 telomerase-immortalized NES-B3T and NES-G2T cells were plated in triplicate. For positive controls, 1,000 SEG-1 cancer cells were also plated in triplicate as previously described (8). Cells were fed weekly with 200 µl of media containing 10% serum. Plates were examined daily for 3 wk. Cells were imaged using a Bio-Rad Molecular Imager (Bio-Rad, Hercules, CA). All experiments were performed in duplicate.

Western blotting and MEK activity assays. Cells and tissues were lysed in 1× cell lysis buffer (Cell Signaling Technology, Beverly, MA). Equal amounts of protein were separated by SDS-polyacrylamide gel electrophoresis; protein concentrations were determined using the BCA-200 Protein Assay kit (Pierce, Rockford, IL). After separation and transfer to nitrocellulose membranes, the membranes were incubated with primary antibodies to p53 and p21 (Oncogene, San Diego, CA), cytokeratins (CK) 13 and 4 (Novocastra, Newcastle upon Tyne, UK), phospho-MEK1/2 (serines 217/221), phospho-ERK1/2, or MKP-1 (Cell Signaling Technology). Horseradish peroxidase secondary antibodies were used, and chemiluminescence was determined using the Super Signal West Dura detection system (Pierce). Membranes were stripped using Restore Stripping Buffer (Pierce) and were reprobed with total ERK1/2 or total MEK1/2 (Cell Signaling Technology). Tubulin (Sigma) was used to confirm equal loading. Doxorubicin-treated MCF-7 cells were used as controls for p53 and p21. Neonatal human foreskin keratinocytes were used as a control for cytokeratins 4 and 13. MEK1/2 activity was determined using a commercially available immunoblot assay per the manufacturer’s instructions (Sigma). In brief, equal amounts of protein lysate were immunoprecipitated with 2 µl of anti-MEK antibody and EZView Red Protein A affinity gel beads for 4 h at 4°C. The supernatant was then removed and the pellet washed three times with 1× wash buffer. Assay buffer and 1 µg of ERK2 substrate were added and incubated with the pellet for 30 min at 30°C. The reaction was terminated with 12 µl of 4× SDS sample buffer followed by Western blotting for phosphorylated ERK1/2 β-tubulin (Sigma) was used to confirm that equal amounts of protein lysates were added to the reaction. All Western blots were performed in duplicate.

UV irradiation. Telomerase-immortalized NES-B3T and NES-G2T squamous epithelial cells were cultured overnight in 100-mm plates as described above. Cells were rinsed several times with 1× PBS and exposed to 200 J/m² UV-B irradiation as previously described (8). Protein was harvested at 6, 24, and 48 h after irradiation. Nonirradiated cells served as controls.

Cytogenetic analysis. Cytogenetic preparations were made following conventional procedures. Briefly, metaphase NES-B3T (PD 53.9) and NES-G2T (PD 64) cells were obtained by Colcemid arrest followed by hypotonic treatment with prewarmed 0.075 M KCl. They were then fixed and washed in freshly made modified Carnoy’s fixative (3:1 absolute methanol:glacial acetic acid), spread onto pre-cleaned, wet microscope slides, and air dried. Trypsin G-banding was performed following a modified methodology (21).

Cell lines and acid exposure. NES-B3T and NES-G2T cell lines were cultured in DMEM/F12 with supplements (growth media). Cells were cultured either in neutral full growth media (pH 7.2) or in acidic full growth media (brought to a pH of 5.0 with 1 M HCl) for 3, 5, or 10 min, after which cells were collected for analyses.

Patients and acid exposure. Patients scheduled for elective endoscopy at the Dallas VA Medical Center who had GERD with and without long-segment Barrett’s esophagus were invited to participate in the study. During endoscopy, three samples of the squamous mucosa were taken using a jumbo biopsy forceps (Olympus FB-50K-1) before and after perfusion of the distal esophagus with 20 ml of 0.1 N HCl for 3 min using the technique previously described by our laboratory (25). None of the patients with GERD had erosive esophagitis at the time of enrollment in the study.

Statistical analyses. Quantitative data are expressed as the mean ± SE. Statistical analyses were performed using the paired Student’s t-test with the Prism statistical software package (GraphPad Software, San Diego, CA). P values ≤0.05 were considered significant for all analyses.

RESULTS

Establishment of immortalized NES-B3T and NES-G2T esophageal squamous cell lines. The log phase doubling times of hTERT-immortalized NES-B3T and NES-G2T esophageal squamous cells were 24–26 h and 36–40 h per PD, respectively. To date, NES-B3T cells have been cultured beyond 100 PDs, and NES-G2T cells have been cultured to 80 PDs, both without evidence of senescence. The TRAP-eze detection kit demonstrated substantial telomerase activity after the introduction of hTERT in both cell lines (Fig. 2A).

NES-B3T and NES-G2T cells express squamous cell cytokeratins and display minimal chromosomal abnormalities. NES-B3T and NES-G2T cell lines showed immunoblot evidence for expression of both CK 4 and CK 13, which are markers of esophageal squamous cell differentiation (3, 5, 20) (Fig. 2B). To identify structural and numerical chromosomal abnormalities, we used conventional G-banding analysis. Our analysis of 20 metaphase cells (~PD 64) for NES-G2T revealed a 47,XY,+t(8)q(10) in 19 cells with one cell containing a 47,XY,+5 karyotype. Our analysis of NES-B3T (~PD 54) revealed a 47,XX,+20 karyotype as the sole abnormality. No other consistent numerical or structural changes were identified.

NES-B3T and NES-G2T cells are nontumorigenic. Immortalization can be the first step in in vitro transformation (14). Therefore, we determined the degree of in vitro tumorigenicity for our NES-B3T and NES-G2T cells. Unlike transformed cells, our cell lines demonstrate cell-contact inhibition (Fig. 3A). Transformed cells also exhibit anchorage independent growth, as evidenced by their ability to form colonies in soft agar. As shown in Fig. 3B, the SEG-1 cancer cell line developed multiple colonies in soft agar, whereas NES-B3T and NES-G2T cells showed no growth after 3 wk. These findings suggest that our long-term cultures of NES-B3T and NES-G2T cells are nontumorigenic although the ultimate evidence that cells are not transformed is the demonstration of their inability to form tumors after injection into athymic nude mice.

NES-B3T and NES-G2T cells express p53 and p21 after UV irradiation. Viral oncoproteins used to immortalize human cells commonly disrupt the p53 cell cycle checkpoint (6). In contrast, telomerase-immortalized epidermal keratinocytes, mammary epithelial cells, esophageal squamous, and Barrett’s epithelial cells maintain appropriate p53 cell cycle checkpoint responses (8, 13, 18). We determined whether our NES-B3T and NES-G2T cells could mount an appropriate p53 response to UV-induced DNA damage (Fig. 4). NES-B3T and NES-G2T cells were harvested at various time points after irradiation with 200 J/m² of UVB. This single dose of UV exposure increased p53 protein expression levels within 6 h, and levels remained elevated at 48 h (Fig. 4). In both cell lines, expression of p21, a downstream effector of p53, increased at 24 h after

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UV exposure and remained elevated at 48 h after exposure (Fig. 4).

Baseline levels of phosphorylated ERK1/2 are lower in NES-G2T than in NES-B3T cells, and acid exposure increases ERK1/2 phosphorylation only in NES-G2T cells. In an earlier study, we showed that the esophageal squamous epithelium of patients with GERD without Barrett’s esophagus had lower baseline levels of phosphorylated ERK1/2 than that of patients with GERD with Barrett’s esophagus (25). In addition, we found that esophageal acid perfusion activated ERK1/2 in the squamous epithelium of patients with GERD without Barrett’s esophagus, but not in those with Barrett’s esophagus (25). In the present study, we explored whether our NES-B3T and NES-G2T cells recapitulated those differences in vitro. In accordance with our in vivo data, we found that NES-G2T cells exhibited lower baseline levels of phosphorylated ERK1/2 than

Fig. 2. A: telomerase activity in parent and human telomerase reverse transcriptase (hTERT)-infected esophageal squamous cell lines. HeLa cells served as a positive (+) control; lysis buffer served as a negative (−) control. TRAP assay shows a characteristic, six-base-pair ladder in cells with telomerase activity, which include the positive control and the NES-B3 and NES-G2 cells infected with hTERT. B: both the NES-G2T and NES-B3T cell lines express the squamous cytokeratin (CK4 and CK13); β-tubulin served as a loading control. Skin derived keratinocytes (Ker) served as a positive control.

Fig. 3. Telomerase-immortalized esophageal squamous cells exhibit contact inhibition and do not grow in soft agar. A: NES-G2T and NES-B3T cells demonstrate early log phase growth, but, with increasing confluence, the cells demonstrate a growth plateau. B: NES-G2T and NES-B3T cells show no colonies after 3 wk in soft agar in contrast to SEG-1 cancer cells, which exhibit numerous colonies.
At 3 and 5 min of acid exposure, ERK1/2 phosphorylation increased in the NES-G2T cells; phospho-ERK1/2 levels were slightly decreased following 10 min of acid exposure (Fig. 5). In contrast, acid exposure decreased levels of phospho-ERK1/2 in NES-B3T cells in a duration-dependent manner (Fig. 5). Thus our esophageal cell lines appear to be good in vitro models that recapitulate the differences in baseline and acid-induced ERK1/2 phosphorylation that we had observed in our earlier study of primary biopsy tissues.

Acid exposure increases MEK1/2 phosphorylation at activating sites (ser217/221) in both NES-G2T and NES-B3T cells. We next used our cells to investigate the mechanisms underlying the disparate effects of acid on ERK1/2 phosphorylation. We found no apparent differences in expression of MKP-1 at baseline or after acid exposure between NES-B3T and NES-G2T cells (data not shown). Thus differences in MKP-1 do not appear to underlie the differences in acid-induced phospho-ERK1/2 phosphorylation that we had observed in our earlier study of primary biopsy tissues.

Acid exposure increases MEK1/2 phosphorylation at activating sites (serines 217/221) in both NES-G2T and NES-B3T cells. We next sought to determine the effects of acid on MEK1/2 phosphorylation sites (serines 217/221) that activate the kinase activity, using total MEK1/2 and tubulin as controls. Compared with nonacid exposed controls, we found a marked increase in phospho-MEK1/2 (ser 217/221) in both NES-G2T and NES-B3T cells at 3 and 5 min of acid exposure; there was no change in phospho-MEK1/2 expression levels by 10 min of acid exposure in either cell line (Fig. 5). These data suggest that failure of acid to cause MEK1/2 phosphorylation at activating sites is not the mechanism underlying the lack of acid-induced phosphorylation of ERK1/2 in NES-B3T cells.

NES-B3T cells have higher levels of MEK1 phosphorylation at an inhibitory site (threonine 286) than NES-G2T cells. MEK1 and MEK2 are highly homologous proteins, but MEK1 has threonine located at position 286, whereas MEK2 has valine at that position. MEK1 phosphorylation at threonine 286 inhibits the kinase activity of the enzyme (19, 23). Compared with NES-G2T cells, NES-B3T cells exhibited higher levels of MEK1 phosphorylation at the threonine 286 inhibitory site at baseline and at all time points after acid exposure (Fig. 6A). Acid exposure had no apparent effect on the levels of this phosphorylated form of MEK1 in either cell line (Fig. 6A). These data suggest, but do not prove, that it is the high basal levels of phospho-MEK1 threonine 286 that prevent ERK1/2 phosphorylation following acid exposure in NES-B3T cells.

Acid exposure increases MEK1/2 activity in NES-G2T cells but not in NES-B3T cells. To confirm that the acid-induced increase in ERK phosphorylation at 3 and 5 min that we observed in our NES-G2T cells is due to an acid-induced increase in MEK1/2 kinase activity, we immunoprecipitated the MEK1/2 kinases before and after acid exposure and performed an in vitro kinase assay using exogenous ERK2 as the substrate. As expected, acid increased MEK1/2 activity in NES-G2T cells but not in NES-B3T cells (Fig. 6B).
Acid exposure increases MEK1/2 phosphorylation at activating sites (serines 217/221) in squamous mucosa from patients with GERD with and without Barrett’s esophagus. Having found that acid exposure increases MEK1/2 phosphorylation at activating sites (serines 217/221) in both the NES-G2T and NES-B3T cell lines, we sought to confirm that these same effects occur in vivo by obtaining endoscopic biopsy specimens of esophageal squamous epithelium before and after acid perfusion in seven patients with GERD with Barrett’s esophagus and five patients with GERD without Barrett’s esophagus.

In agreement with our in vitro data, we found that acid significantly increased phospho-MEK1/2 levels at serines 217/221 in the esophageal squamous epithelium of patients with GERD both with \( [0.41 \pm 0.08 \text{ to } 0.63 \pm 0.28 \text{ RIUs, } P = 0.04] \) and without \( [0.52 \pm 0.10 \text{ to } 0.87 \pm 0.42 \text{ RIUs, } P = 0.03] \) Barrett’s esophagus (Fig. 7). Using the same patient samples, we determined MEK1/2 activity by performing Western blot for the active, diphosphorylated form of ERK1/2. ERK1/2 phosphorylation data from some of these patients were included in our previous publication (25). In agreement with our previous findings, we found that acid significantly increased phosphorylated ERK1/2 expression in the esophageal squamous mucosa from patients who had GERD without Barrett’s esophagus \( [0.59 \pm 0.09 \text{ to } 0.85 \pm 0.05, P = 0.03] \) but not in those with Barrett’s esophagus \( [0.76 \pm 0.07 \text{ to } 0.85 \pm 0.08, P = 0.43] \) (Fig. 8).

Phosphorylated levels of MEK1 at threonine 286 are found in the esophageal squamous epithelium of patients with GERD with Barrett’s esophagus but not of those without Barrett’s esophagus. Our in vitro data demonstrated that levels of phospho-MEK1 at the inhibitory site (thr 286) were not altered by acid exposure. We detected this inhibitory form of phospho-MEK1 in esophageal squamous epithelial biopsies from eight patients who had GERD with Barrett’s esophagus but not in eight patients who had GERD without Barrett’s esophagus (Fig. 9).

**DISCUSSION**

This study adds support to our hypothesis that, in esophageal squamous epithelial cells, there are differences among individuals in signal transduction pathways activated by acid reflux that might underlie the development of Barrett’s esophagus. To explore that hypothesis, we established stable, telomerase-immortalized, but not transformed, esophageal squamous epithelial cell lines from patients with GERD with and without Barrett’s esophagus. We have shown that our cells maintain morphological and immunohistochemical characteristics of esophageal squamous epithelium, including the expression of squamous epithelial cell markers like cytokeratins 4 and 13 (3, 5, 20). Unlike transformed cells, our cells maintain growth inhibition with cell-to-cell contact, anchorage-dependent growth, and an intact p53 cell cycle checkpoint. Thus our cell lines maintain a number of normal features desirable for an in vitro model of nonneoplastic esophageal squamous epithelial cells with which to study early events in the pathogenesis of metaplasia.

We first set out to determine whether our squamous cell lines could recapitulate our in vivo finding from an earlier study that acid causes activation of ERK1/2 in the esophageal squamous epithelium of patients who have GERD without Barrett’s esophagus but not in patients with Barrett’s esophagus. As in that study, we found that baseline levels of the active, diphosphorylated form of ERK1/2 were lower in squamous cells from a patient who had GERD without Barrett’s esophagus (NES-G2T cells) than in those from a patient with Barrett’s esophagus (NES-B3T cells) and that acid exposure increased phospho-ERK1/2 levels in NES-G2T cells but not in NES-B3T cells.
This in vitro recapitulation of in vivo findings supports the use of our telomerase-immortalized cells as reasonable models for investigating mechanisms underlying differences in acid-induced ERK1/2 phosphorylation in patients with and without Barrett’s esophagus.

Even if acid exposure causes ERK1/2 phosphorylation in esophageal squamous cells, ERK1/2 could be rapidly dephosphorylated, and thus inactivated, if the cells have high levels of MKPs such as MKP-1 (24). There is a strong, inverse correlation between levels of MKP-1 and phospho-ERK1/2, and MKP-1 expression can be induced by a variety of cell stresses (10, 11, 27). We found that acid exposure caused no apparent change in MKP-1 expression in either NES-B3T or NES-G2T cells, suggesting that differences in MKP-1 expression are not responsible for the differences that we observed in acid-induced ERK1/2 phosphorylation. Nevertheless, our findings do not exclude the possibility that other types of MKPs (e.g., serine/threonine phosphatases such as PP2A, protein Tyr phosphatases such as PTP-SL) may affect acid-stimulated ERK1/2 phosphorylation levels in esophageal squamous cells. Further investigations are needed in this area (1, 17, 27).

ERK1/2 is phosphorylated by MEK1/2, whose activity also is regulated by phosphorylation (24). MEK1/2 phosphorylation at serines 217/221 activates, whereas MEK1 phosphorylation at threonine 286 inhibits the catalytic activity of the enzymes. In both NES-G2T and NES-B3T cells, acid increased MEK1/2 phosphorylation at serines 217/221. This suggests that differences in acid-induced ERK1/2 phosphorylation between the cell lines do not result from failure of acid to induce phosphorylation at MEK1/2 activating sites. At the threonine 286 inhibitory site of MEK1, however, NES-B3T cells exhibited higher levels of phosphorylation than NES-G2T cells at baseline, and the levels did not change with acid exposure. This suggests that the lack of acid-induced ERK1/2 phosphorylation in NES-B3T cells is not due to acid-induced increases in MEK1 phosphorylation at threonine 286. We speculate that the high basal levels of phospho-MEK1 threonine 286 in the NES-B3T cells interfere with the efficiency of the MEK1 kinase in phosphorylating and activating ERK1/2 when it is exposed to acid. In support of this contention, MEK1 phosphorylation at threonine 286 in NIH3T3 and HeLa cells has been shown to decrease their levels of phospho-ERK1/2 (19, 23). Nevertheless, our studies do not prove that it is the high levels of threonine 286 phosphorylation that underlie the differences between the cell lines in the degree of phosphorylation of ERK1/2 after acid exposure.

Signals transduced by ERK can result in cellular proliferation, differentiation, or apoptosis depending on the cell type, the duration of the signal, and the degree of MEK activity. In
PC12 cells, for example, transient activation of ERK1/2 signaling by epidermal growth factor causes proliferation, whereas sustained ERK1/2 activation by neural growth factor leads to differentiation (15, 22, 29). Alternatively, if ERK1/2 activation by neural growth factor is inhibited, then apoptosis ensues (22). Other investigators have found that sustained ERK1/2 activation also can induce apoptosis in PC12 cells (31). Moreover, alterations in MEK1 phosphorylation at threonine 286 have been shown to alter ERK1/2 activity and its effects on apoptosis (22, 31).

The biological consequences of our observation that there are differences in acid-induced MEK-ERK activation between esophageal squamous cells from patients with and without Barrett’s esophagus are not clear. However, it is conceivable that those differences underlie the development of Barrett’s metaplasia. Proproliferative signals transduced by ERK1/2 appear to facilitate the healing of ulcerations in the gastrointestinal tract (28). Therefore, a squamous epithelium that responds to acid exposure by activating ERK1/2 might be predisposed to heal acid-peptic injuries through the regeneration of new squamous epithelial cells (a proliferative response). Conversely, a squamous epithelium that does not experience ERK1/2 activation with acid exposure might be unable to heal its acid-peptic injuries through proliferation and instead might be predisposed to heal through the process of metaplasia, in which squamous cells are replaced by columnar ones. Given the key role that this pathway plays in cell survival, further studies to elucidate the consequences of acid-induced differences in its activation are warranted.

To confirm our in vitro findings, we studied the effects of esophageal acid perfusion during endoscopic examination on MEK1/2 phosphorylation levels in the esophageal squamous epithelium of patients with GERD with and without Barrett’s esophagus. As in our cells lines, acid exposure significantly increased MEK1/2 phosphorylation at serines 217/221 in the squamous epithelium of both patient groups, but phospho-MEK1 expression at threonine 286 was detected only in the patients with Barrett’s esophagus. These observations further support the utility of our NES-G2T and NES-B3T cell lines as models for studying GERD effects.

In conclusion, we have shown that acid exposure, both in vitro and in vivo, increases ERK1/2 phosphorylation in esophageal squamous cells from patients with GERD without Barrett’s esophagus but not with Barrett’s esophagus. We have also shown that esophageal squamous cells from patients with GERD with Barrett’s esophagus have higher levels of MEK1 phosphorylation at an inhibitory site and have decreased MEK1/2 activity. We speculate that differences in reflux-induced activation of molecular pathways that regulate cellular proliferation and differentiation in esophageal squamous cells contribute to the pathogenesis of Barrett’s metaplasia.

REFERENCES


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