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Author(s)	Betsuyaku, Tomoko; Tanino, Mishie; Nagai, Katsura; Nasuhara, Yasuyuki; Nishimura, Masaharu; Senior, Robert M.
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Extracellular Matrix Metalloproteinase Inducer Is Increased in Smokers' Bronchoalveolar Lavage Fluid

Tomoko Betsuyaku, Mishie Tanino, Katsura Nagai, Yasuyuki Nasuhara, Masaharu Nishimura, and Robert M. Senior

First Department of Medicine, Hokkaido University School of Medicine, Sapporo, Japan, and the Departments of Medicine and Cell Biology and Physiology, Washington University School of Medicine, St. Louis, Missouri

Extracellular matrix metalloproteinase inducer (EMMPRIN), also called basigin, is present in the lung during development, but its expression in normal adult lung is minimal. Increases of EMMPRIN have been found in various forms of experimental lung injury. To determine whether EMMPRIN might be involved in alveolar injury/repair associated with smoking, we developed an ELISA for EMMPRIN and applied it to bronchoalveolar lavage fluids from never-smokers ($n = 7$), former smokers ($n = 16$), and current smokers ($n = 58$). The smoker groups included subjects with emphysema, as determined by high-resolution chest computed tomography. EMMPRIN levels were significantly elevated in current and former smokers (315 ± 20 and 175 ± 15 pg/ml SEM, respectively, compared with 31 ± 7 pg/ml in never-smokers), but the EMMPRIN levels of smokers with emphysema were not different from smokers without emphysema. Immunohistochemistry of smokers' lung tissue showed EMMPRIN in bronchiolar epithelium and alveolar macrophages, but EMMPRIN mRNA in alveolar macrophages was not different between current and never-smokers. Matrix metalloproteinase-1 was also detectable in the bronchoalveolar lavage fluid from some smokers but not in never-smokers. These findings indicate that smoking is associated with increased intrapulmonary EMMPRIN. Whether EMMPRIN is involved in smoking-induced lung pathology remains to be determined.

Keywords: smoking; emphysema; matrix metalloproteinases; alveolar macrophages

Extracellular matrix metalloproteinase (MMP) inducer (EMMPRIN) is a member of the IgG superfamily (1). EMMPRIN is known by other names: CD147 and M6 in humans; basigin and gp42 in mouse; OX-47 and CE9 in rat; and HT7, neurothelin, and 5A11 in chicken. EMMPRIN is strongly expressed in many tissues during development, including lung (2), but only at low levels in normal adult lung and other tissues (3, 4), suggesting a physiologic role in tissue differentiation and development. EMMPRIN was originally identified on the surface of tumor cells and was termed tumor cell-derived collagenase stimulatory factor (5) because it induced collagenase in adjacent stromal cells (6, 7). Its possible role in tumor cell invasion and metastasis is supported by studies showing increased dissemination of tumor cells with high levels of basigin expression (8). Its upregulation in tissues

has been also identified in various nonmalignant conditions, including rheumatoid arthritis (9, 10), chronic liver disease (11), heart failure (12, 13), ischemic myocardial injury (14), atherosclerosis (15), and experimental ventilator-induced lung injury (16).

MMPs are a family of neutral proteinases that degrade extracellular matrix and modify other substrates (17). MMPs have been implicated in various injury and/or repair processes in the lungs (18, 19) and other organs. MMPs are regulated at both pretranscriptional and post-transcriptional levels, and their activities can also be regulated by endogenous inhibitors, including the tissue inhibitors of MMPs, α_2 -macroglobulin, and reversion-inducing cysteine-rich protein with Kazal motifs (RECK) (20). Because several MMPs are increased in various types of human alveolar pathology (21, 22) and in experimental bleomycin-induced lung injury (23–25), we recently investigated the expression of basigin (murine EMMPRIN) after bleomycin. We found basigin in fibroinflammatory areas, increased basigin in bronchoalveolar lavage (BAL) fluids, and basigin mRNA upregulated in bronchiolar epithelial cells (26).

Cigarette smoking is a major risk factor for the development of emphysema. Although the molecular mechanisms linking smoking to the pathogenesis of emphysema have not been fully clarified, MMPs have been implicated in its development (27–30). Therefore, we hypothesized that EMMPRIN may be upregulated in smokers' lungs and may be involved in the intrapulmonary expression of MMPs associated with smoking. To examine this possibility, we have developed an ELISA for EMMPRIN and have applied it to BAL fluids from never-smokers and smokers. We have observed that EMMPRIN levels are significantly increased in smokers' BAL fluid compared with never-smokers.

METHODS

Subjects

We recruited 81 volunteers (aged 32–80 years), including 58 current smokers, 16 former smokers who had not smoked for at least 1 year, and 7 individuals who had never smoked. The subjects had no respiratory symptoms and denied acute respiratory infections within the preceding 2 months. The serum α -1 antitrypsin concentration was more than 160 mg/dl in all subjects. Measurements of VC, FEV₁, and diffusing capacity were performed on all subjects (CHESTAC-55V; Chest Co., Tokyo, Japan). High-resolution chest computed tomography scans were performed on all current and former smokers to screen for emphysematous changes, as previously described (31). Three pulmonary physicians who were blinded to any information about the subjects independently evaluated the computed tomography scans for the presence of emphysema. Emphysema was considered present only if all three physicians agreed, regardless of severity. Informed consent was obtained from each subject, and the Ethics Committee of Hokkaido University School of Medicine approved the study.

Sequential BAL and Processing of BAL Fluids

Current smokers were asked to refrain from smoking for at least 12 hours before the BAL procedure to reduce acute effects of smoking. Four separate 50-ml aliquots of sterile 0.9% saline were instilled into a segment of the right middle lobe through a wedged flexible fiberoptic

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Correspondence and requests for reprints should be addressed to Tomoko Betsuyaku, M.D., Ph.D., First Department of Medicine, Hokkaido University School of Medicine, N-15, W-7, Kita-ku, Sapporo 060-8638, Japan. E-mail: bytomoko@med.hokudai.ac.jp

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bronchoscope (Olympus BF-B3R, Tokyo, Japan) and then gently aspirated. The fluid returned from the first 50-ml aliquot was discarded to reduce the contribution from large airways (32). The fluid from the other three aliquots was combined and filtered through several layers of gauze to remove excess mucus and debris, and a sample was taken for cell count and differential. The fluid was then centrifuged at 1,500 rpm for 5 minutes at 4°C. The cell-free supernatant was aliquoted and stored at -70°C. The albumin concentration was measured as previously described (31).

Quantification of EMMPRIN by ELISA

An enzyme immunoassay was developed for the quantification of EMMPRIN in BAL fluid. The wells of microtiter plates (Immulon #2; Dynatech Laboratories, Inc., Chantilly, VA) were coated overnight by incubation at room temperature with 1 µg/ml of goat polyclonal antibody raised against recombinant extracellular domain of human EMMPRIN (R&D Systems, Abingdon, UK). After three washes with wash buffer (0.05% Tween 20 in phosphate-buffered saline [PBS], pH 7.4), the plate was blocked by PBS containing 1% bovine serum albumin (BSA), 5% sucrose, and 0.05% NaN₃ and was incubated at room temperature for a minimum of 1 hour. For the generation of a standard curve, recombinant human EMMPRIN (R&D Systems) was diluted with 0.1% BSA and 0.05% Tween 20 in TBS (pH 7.3) in polypropylene tubes to final concentrations of 1,000, 500, 250, 125, 62.5, 31.3, 15.6, and 7.8 pg/ml. The EMMPRIN used to establish the standard curve showed a single band at approximately 40 kD by Western blotting with the primary antibody, using an enhanced chemiluminescence Western blotting analysis system (Amersham, Aylesbury, UK) as previously described (26) (Figure 1). The plate was then washed three times with wash buffer, and 100 µL of unconcentrated BAL fluid or standard were added per well, followed by incubation for 2 hours at room temperature. After three washes with wash buffer, the plate was incubated with 100 µL of biotinylated antiextracellular domain of human EMMPRIN antibody (R&D Systems) at 400 ng/ml for 2 hours at room temperature. Then the plate was washed three times with wash buffer and 100 µL of streptavidin horseradish peroxidase (R&D Systems), diluted according to the manufacturer's direction, was added to each well. After 20 minutes, the plate was washed three times and incubated with 100 µL of a 1:1 mixture of H₂O₂ and tetramethylbenzidine (R&D Systems) in the dark, and the reaction was stopped with 1 M of sulfuric acid within 30 minutes. The absorbance values were determined at 450 nm using a microplate reader (Wallac 1,420, Turku, Finland). A linear response was obtained with EMMPRIN standards from 7.8 to 1,000 pg/ml (Figure 2A). The ELISA had an interbatch coefficient of variation of 8% (n = 9) and an intrabatch coefficient of variation of 14% (n = 10), thus providing precise and sensitive detection of EMMPRIN. All assays were done in duplicate, and the mean values were used. All BAL fluid samples were on the standard curve when measured unconcentrated.

To test the reliability and accuracy of the EMMPRIN ELISA, selected BAL fluids were spiked with 125, 250, 500, and 1,000 pg of recombinant EMMPRIN and then assayed for EMMPRIN content. As shown in Figure 2B, assays of spiked BAL fluid consistently reflect the sum of the EMMPRIN in the starting sample plus the EMMPRIN spike.

Immunohistochemistry

Lung tissue was obtained from nine male smokers at the time of lobectomy for removal of primary lung cancer. These included subjects with and without airflow obstruction by pulmonary function testing. The tissue specimens were fixed in 10% phosphate-buffered formalin and embedded in paraffin. Five-micrometer sections were deparaffinized in xylene and rehydrated with graded alcohols. Antigenic activity was retrieved by incubation in 10 mM of citrate buffer (pH 6.0) in a microwave for 5 minutes. After washing in PBS, the slides were processed for immunohistochemistry using a catalyzed signal amplification (CSA) kit (DAKO Japan, Kyoto, Japan) according to the manufacturer's protocol. The primary antibody was anti-EMMPRIN monoclonal antibody (CHEMICON, Temecula, CA) diluted at 1:500 with PBS.

Quantitative Reverse Transcription-Polymerase Chain Reaction for EMMPRIN

Alveolar macrophages (AMs) recovered from BAL fluids were adjusted to 1 × 10⁶ per ml and plated on poly-L-lysine-coated plastic wells

Figure 1. Western blot of recombinant EMMPRIN. The recombinant human EMMPRIN (40 ng) used to establish the standard curve for the EMMPRIN ELISA showed a single band at approximately 40 kDa detected by the goat polyclonal antibody raised against recombinant extracellular domain of human EMMPRIN.

(Corning, NY) for 1 hour at 37°C in RPMI-1640 medium (GIBCO, Grand Island, NY). After two washes in PBS, the adherent cells were used for purification of total RNA with guanidium thiocyanate using an Isogen RNA extraction kit (Nippon Gene, Tokyo, Japan). Reverse transcription was performed in the presence of Maloney leukemia virus reverse transcriptase (Epicentre Technologies, Madison, WI), and

Figure 2. Standard curve of the ELISA for EMMPRIN. (A) A linear response was obtained with concentrations of recombinant EMMPRIN from 7.8 to 1,000 pg/ml ($r = 0.99$). The concentration of EMMPRIN in BAL fluid was calculated from the standard curve run in parallel with the fluid. (B) One hundred microliters of BAL fluid were spiked with 125, 250, 500, and 1,000 pg of exogenous recombinant EMMPRIN, and the samples were assayed for EMMPRIN (closed circles) together with 100 µL of 0.1% BSA, 0.05% Tween 20 in tris-buffered saline (pH 7.3) to which 125, 250, 500, and 1,000 pg of exogenous recombinant EMMPRIN was added (open circles). The two sets of samples yielded results that consistently differed by the content of EMMPRIN in the starting BAL fluid. OD = optical density.

5' exonuclease-based fluorogenic polymerase chain reaction was performed using an ABI PRISM 7,700 Sequence Detector (PE Applied Biosystems, Foster City, CA) as described (33). Oligonucleotide primers for EMMPRIN were derived from exon 5 of the human EMMPRIN gene sequence (GenBank accession number: L20471) (34). The primers and probe for this quantification system designed to meet specific criteria by using Primer Express software (PE Applied Biosystems) were as follows: forward, 5'-ACCTGGCCGCCCTCTG -3'; reverse, 5'-TCGTAAGTAGAAGATGATGGTGACC-3'; and probe, 5'-FAM-CCTTCCATGGCATCGTGGCTGAGGT-TAMRA-3'.

The predicted size of the amplified EMMPRIN DNA product was 83 bp, and specific amplification was confirmed by ethidium bromide staining of the polymerase chain reaction product on an agarose gel. Primers and a labeled probe for glyceraldehyde-3-phosphatase-dehydrogenase (GAPDH) cDNA as an endogenous control were purchased from PE Applied Biosystems (TaqMan human GAPDH control reagents). The relative amount of EMMPRIN mRNA in the samples was normalized by GAPDH mRNA.

Interstitial Collagenase (MMP-1)

MMP-1 levels in BAL fluids were measured by a solid-phase enzyme immunoassay (MMP-1 Immunoassay Kit; CHEMICON) that recognizes the zymogen and active forms of MMP-1 as well as forms complexed with tissue inhibitor of metalloproteinase-1. Importantly, the assay does not recognize MMP-8, the collagenase expressed by neutrophils. The assay's range of detection was 0.11–3.6 ng/ml. Because MMP-1 was not detected in any unconcentrated BAL fluid, the fluids were concentrated 15-fold using Centricon filters having a molecular weight cutoff of 10,000 (Amicon Inc., Beverly, MA). Each fluid was measured in duplicate, and the results were averaged.

Statistical Analysis

Results are expressed as mean $\pm$ SEM unless otherwise stated. Statistical analyses were performed on the data using one-factor analysis of variance among four groups and Student's unpaired *t* test between two groups when appropriate. The Spearman's rank correlation was calculated to assess the correlation between the data. In all tests, *p* values of less than 0.05 were considered statistically significant.

RESULTS

Characteristics of the Subjects

The subjects were classified into five groups according to smoking history and presence of emphysema on computed tomography scans (Table 1). There were no significant differences in age among the groups. Emphysema, as detected on computed tomography scans, was less than 25% of the total lung area in most subjects categorized in the emphysema groups. The FEV₁/FVC was significantly lower in the current smokers with emphysema than in never-smokers and current smokers without emphysema. Table 2 summarizes findings in the BAL fluids. The concentrations of albumin, cell numbers, and differentials in BAL fluids were not statistically different among the five groups, but the percentage recovery of instilled fluid was significantly lower in current and former smokers with emphysema than in never-smokers (*p* < 0.05).

TABLE 1. CHARACTERISTICS OF SUBJECTS

	Never-smokers	Former Smokers		Current Smokers	
		Emphysema (-)	Emphysema (+)	Emphysema (-)	Emphysema (+)
Number of subjects	7	9	7	26	32
Age, yr	64 $\pm$ 3	66 $\pm$ 3	55 $\pm$ 4	53 $\pm$ 2	56 $\pm$ 2
Pack-year	0	34 $\pm$ 8	50 $\pm$ 10	48 $\pm$ 4	46 $\pm$ 4
FEV ₁ /FVC, %	84 $\pm$ 2	79 $\pm$ 2	71 $\pm$ 5	79 $\pm$ 1	71 $\pm$ 2 [†]

[†] *p* < 0.05 vs. never-smokers.

* *p* < 0.05 vs. current smokers without emphysema.

Mean $\pm$ SE.

EMMPRIN in BAL Fluids

EMMPRIN was detected by ELISA in unconcentrated BAL fluid from all subjects. The EMMPRIN levels were significantly elevated in former and current smokers (175 $\pm$ 15 pg/ml SEM and 315 $\pm$ 20, respectively, compared with 31 $\pm$ 7 pg/ml in life-long never-smokers) (Figure 3). However, the levels were not different between the subjects with and without emphysema among former and current smokers. There was a significant but weak correlation between the EMMPRIN level in BAL fluid and the number of AMs in BAL fluid (*n* = 81, *p* < 0.01, *r* = 0.43) (Figure 4).

Localization of EMMPRIN in Smokers' Lung

By immunohistochemistry, EMMPRIN was found predominantly in AMs, bronchiolar epithelium, and bronchial glands in current smokers' lungs. It was present diffusely in the cytoplasm of AMs and bronchial glands (Figures 5b and 5d) and was concentrated at the apices of bronchiolar epithelial cells (Figure 5c). In contrast, EMMPRIN was weakly seen in AM and was barely detectable in airways in lung tissue from life-long never-smokers (Figure 5a).

EMMPRIN mRNA in AMs

EMMPRIN mRNA relative to GAPDH mRNA was determined in 21 subjects, including 7 never-smokers and 14 current smokers, including 5 with emphysema. EMMPRIN mRNA/GAPDH mRNA in AM in current smokers with emphysema was increased compared with the others, although it did not reach statistical significance because of the small number of subjects tested (Figure 6). There was a positive but weak relationship between the EMMPRIN concentration in BAL fluid and EMMPRIN mRNA levels in AMs (*n* = 21, *p* < 0.05, *r* = 0.43).

MMP-1

MMP-1 was not detectable by immunoassay in concentrated BAL fluids from 7 never-smokers but was measurable in concentrated fluids in seven of 36 current smokers, including three with emphysema (Figure 7).

DISCUSSION

EMMPRIN stimulates the production of MMPs by mesenchymal cells *in vitro*, including MMP-1, MMP-2, MMP-3, MMP-9, and MMP-14 (6–8, 35). It also induces the expression of MMPs by cells *in vivo*. For example, human breast cancer cells transfected with EMMPRIN cDNA injected into mammary fat pads of athymic mice have increased expression of MMP-2 and MMP-9 and are more tumorigenic and invasive than control plasmid-transfected cancer cells (8). MMP-1, and possibly other MMPs, binds to EMMPRIN on tumor cells, an effect that may facilitate MMP activity (36). EMMPRIN has been found to interact with cells via a hemo-

TABLE 2. BRONCHOALVEOLAR LAVAGE FLUIDS

	Never-smokers	Former Smokers		Current Smokers	
		Emphysema (-)	Emphysema (+)	Emphysema (-)	Emphysema (+)
Recovery rate, %	71 $\pm$ 4	52 $\pm$ 5	38 $\pm$ 6 [†]	48 $\pm$ 3 [†]	38 $\pm$ 3 [†]
Albumin, mg/L	56 $\pm$ 10	51 $\pm$ 8	44 $\pm$ 16	36 $\pm$ 4	47 $\pm$ 6
Total cells, $\times 10^{-4}$ /mL	13 $\pm$ 1	9 $\pm$ 2	10 $\pm$ 2	33 $\pm$ 6	30 $\pm$ 6
Macrophages, %	81 $\pm$ 6	88 $\pm$ 2	81 $\pm$ 7	91 $\pm$ 2	91 $\pm$ 2
Neutrophils, %	0 $\pm$ 0	1 $\pm$ 0	3 $\pm$ 2	2 $\pm$ 1	2 $\pm$ 0

[†] *p* < 0.05 vs. never-smokers.

Mean $\pm$ SE.

Figure 3. EMMPRIN concentrations in BAL fluids. EMMPRIN levels were significantly elevated in both former and current smokers (179 ± 17 pg/ml SEM and 319 ± 20 , respectively, compared with 31 ± 7 in never smokers, $p < 0.05$), but the levels were not significantly different between the subjects with and without emphysema in each group (open circles, without emphysema; closed circles, with emphysema).

philic interaction predominantly through the first immunoglobulin domain (37). Also, EMMPRIN colocalizes with $\alpha 3 \beta 1$ and $\alpha 6 \beta 1$ integrins and can be cross-linked, suggesting an interaction with integrins (38).

Although induction of MMPs is clearly an activity of EMMPRIN, EMMPRIN may have other activities. Complete EMMPRIN deficiency in mice, produced by gene targeting, is associated with a high level of early embryonic lethality (39). Deficient mice that survive have poor growth and abnormalities of the reproductive and central nervous systems (40), and most die before 2 months of age with diffuse infiltration of the lungs with lymphocytes and granulocytes (39).

By Western blotting, EMMPRIN is detectable in BAL fluids from normal rats and mice and is increased in BAL fluids from rats with ventilator-induced lung injury (16) and mice with bleomycin-induced lung fibrosis (26). This study is the first to quantify EMMPRIN in BAL fluid and the first report of EMMPRIN in human BAL fluid. The ELISA that we have developed has enabled reproducible measurements of EMMPRIN in unconcentrated BAL fluid from never-smokers. The average concentration in these subjects is approximately 32 pg/ml. The concentration is approximately 10-fold higher among current smokers. Former smokers also have elevated levels, approximately fivefold higher than never-smokers despite not smoking for 1 year or longer before obtaining their BAL fluid, indicating that the mechanisms responsible for increased EMMPRIN associated with smoking persist.

Figure 4. Relationship between AM number in BAL fluid and EMMPRIN concentration. EMMPRIN concentration was weakly correlated to the number of AMs in BAL fluid ($\times 10^{-4}$ per ml) ($n = 81$, $p < 0.01$, $r = 0.43$).

Figure 5. Immunohistochemistry of EMMPRIN in the lungs from never-smokers and smokers. EMMPRIN is weakly present in AM in never-smokers (a) ($\times 200$). In current smokers with emphysema, EMMPRIN is prominent in bronchial glands (b) ($\times 200$), bronchiolar epithelium (c) ($\times 400$), and AMs (d) ($\times 400$). Arrows indicate bronchiolar epithelium and macrophages, respectively.

The protein core of EMMPRIN is approximately 27 kD. Glycosylated forms are 43–66 kD (1), and the capacity to induce MMPs requires glycosylation (5, 37). The glycosylation state of EMMPRIN varies among tissues (40). In experimental lung injury, there are increased high molecular weight forms in addition to an increased quantity of EMMPRIN in the BAL fluid (16, 26). The EMMPRIN in smokers' BAL fluid in this study also showed multiple molecular weights, corresponding to unglycosylated and glycosylated forms, resembling forms seen in experimentally induced lung injury in rodents and, therefore, might be expected to include EMMPRIN with the capacity to induce MMPs.

The molecular basis for upregulation of EMMPRIN expression in various situations, including the association with smoking, is not known. Likewise, the mechanisms by which EMMPRIN induces MMPs have not been fully established. Binding of Sp1 and Sp3 to an Sp1 element in the EMMPRIN promoter appears to be required for regulation of EMMPRIN gene expression in macrophages (41). The levels of Sp1 and Sp3, as well as their activation and binding, are affected by a variety of stimuli so

Figure 6. EMMPRIN mRNA in AM. By real-time polymerase chain reaction, EMMPRIN mRNA in AMs was not different between current and never-smokers (open circles, without emphysema; closed circles, with emphysema).

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