IL-1 β -Induced Increase in Intestinal Epithelial Tight Junction Permeability Is Mediated by MEKK-1 Activation of Canonical NF- κ B Pathway

Rana Al-Sadi, Dongmei Ye, Hamid M. Said, and Thomas Y. Ma

From the Department of Internal Medicine, University of New Mexico School of Medicine and Albuquerque Veterans Affairs Medical Center, Albuquerque, New Mexico; and the Department of Internal Medicine, University of California, Irvine, California

IL-1 β is a proinflammatory cytokine that plays a central role in the inflammatory process of the gut. IL-1 β causes an increase in intestinal epithelial tight junction (TJ) permeability, but the intracellular pathways that mediate intestinal TJ permeability remain unclear. The major aims of this study were to delineate the protein kinases that regulate the IL-1 β modulation of intestinal TJ barrier function and to determine the intracellular mechanisms involved, using filter-grown Caco-2 monolayers as the *in vitro* model system. Our results showed that IL-1 β caused a rapid activation of MEKK-1 and NIK. The knockdown of MEKK-1, but not NIK, inhibited the IL-1 β increase in Caco-2 TJ permeability. IL-1 β caused an activation of both canonical and noncanonical NF- κ B pathways; MEKK-1 regulated the activation of the canonical pathway, while NIK regulated the activation of the noncanonical pathway. Inhibition of MEKK-1 activation of the canonical pathway prevented the IL-1 β increase in TJ permeability. Our data also indicated that inhibitory κ B kinase was the catalytic subunit primarily involved in canonical pathway activation and TJ barrier opening. MEKK-1 also played an essential role in myosin light chain kinase gene activation. In conclusion, our data show for the first time that MEKK-1 plays an integral role in IL-1 β modulation of Caco-2 TJ barrier function by regulating the activation of the canonical NF- κ B pathway and the MLCK gene. (*Am J Pathol* 2010, 177:2310–2322; DOI: 10.2353/ajpath.2010.100371)

Defective intestinal epithelial tight junction (TJ) barrier has been implicated to be an important pathogenic factor

in number of inflammatory conditions of the gut and systemic inflammatory conditions, including Crohn's disease (CD), postinfectious irritable bowel syndrome, nonsteroidal anti-inflammatory drug associated enteritis, ulcerative colitis, heat stroke, alcoholic hepatitis, and various infectious diarrheal syndromes.^{1–5} It has been postulated that the defective intestinal TJ barrier allows paracellular permeation of noxious luminal antigens that propagate and contribute to the inflammatory response.^{5–7} It is well-established that patients with CD have a defective intestinal TJ barrier manifested by an increase in intestinal permeability.^{1,2,5,8} Intestinal permeability studies in healthy first degree relatives of CD patients (an at risk population to develop CD) showed that the healthy relatives also had an abnormal increase in intestinal permeability, prompting the investigators to conclude that the increase in intestinal permeability “is a primary defect that may be an etiological factor in this disease.”^{1,8}

Interleukin-1 β (IL-1 β) is one of the first cytokines to be discovered and has been shown to play a central role in intestinal inflammation in CD.^{9–12} A direct correlation exists between elevated levels of IL-1 β and severity of intestinal inflammation in CD.^{11–14} Patients with CD also have an imbalance between the level of IL-1 β and its naturally occurring antagonist IL-1 receptor antagonist (IL-1ra) such that they have deficiency of anti-inflammatory form of IL-1 and excess production of IL-1 β .^{15,16} In addition, CD patients have increased incidence of IL-1 β gene polymorphism that determines the severity of intestinal inflammation.^{17,18} IL-1 β antagonists have been shown to be effective in the treatment of immune-mediated inflammation in mice and are currently being

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Address reprint requests to Thomas Y. Ma, M.D., Ph.D., Internal Medicine-Gastroenterology, MSC10 5550, University of New Mexico, Albuquerque, NM 87131-0001. E-mail: tma@salud.unm.edu.

developed for clinical usage.^{19,20} Previous studies have shown that IL-1 β causes an increase in intestinal TJ permeability, and it has been postulated that the defect in intestinal TJ barrier contributes to the development of intestinal inflammation.^{21,22}

Recent studies from several laboratories have shown that proinflammatory cytokines (including IL-1 β , TNF- α , and IFN- γ) cause an increase in intestinal epithelial TJ permeability.^{22–27} The cytokine-induced increase in intestinal TJ permeability has been postulated to be an important factor in the development of intestinal inflammation.^{2,5,7,28} The role of cytokine-induced alteration in intestinal permeability as a pathogenic factor of intestinal inflammation has been supported by animal studies showing that the preservation of intestinal TJ barrier function prevents the development of intestinal inflammation.^{7,28–31} In IL-10-deficient mice (IL-10^{–/–}), a commonly used murine model of inflammatory bowel disease, the development of intestinal inflammation was preceded by an initial increase in intestinal permeability,³² suggesting a possible cause-and-effect relationship. The inhibition of intestinal TJ barrier defect by oral administration of TJ barrier enhancing agent AT-1001 (a zonulin peptide inhibitor) prevented the development of enterocolitis in IL-10^{–/–} mice, leading the authors to conclude that the “abnormal small intestinal permeability not only precedes the development of colitis but is etiologically important.”³¹ Similarly, other investigators have shown that the maintenance of intestinal TJ barrier function in various murine models of intestinal inflammation also prevents the development of intestinal inflammation and its clinical sequelae.^{29,30,33} Consistent with the above animal studies, clinical studies have also revealed that the therapeutic re-tightening of intestinal TJ barrier is associated with more rapid improvement and resolution of active CD and prolonged clinical remission.^{34–36} Conversely, persistent increase in intestinal permeability after medical therapy was predictive of poor clinical outcome and early exacerbation of the disease.^{35,36} Together, these studies suggested that the therapeutic preservation or re-tightening of intestinal TJ barrier has important potential clinical implications.

Myosin light chain kinase (MLCK) gene and protein play a central role in IL-1 β modulation of intestinal epithelial TJ barrier.^{2,5,7,28} Previous studies have shown that the IL-1 β -induced increase in intestinal epithelial TJ permeability was mediated by an increase in MLCK mRNA transcription and protein synthesis.²² The inhibition of MLCK transcription or protein synthesis completely inhibited the IL-1 β -induced increase in Caco-2 TJ permeability.²² The intracellular processes or protein kinase pathways that mediate the IL-1 β alteration in MLCK gene activity or increase in intestinal epithelial TJ permeability are unknown. Mitogen activated protein kinase kinases (MAP3 kinase) are recruited by IL-1 β receptor complex and play a crucial role in the regulation of variety of biological activities in intestinal epithelial cells.^{37–39} However, their involvement in TJ barrier regulation remains unknown. IL-1 β has been shown to activate MAP3 kinases including MEKK-1 and NIK in different cell types.^{40–43} The major aim of this study was to examine the

regulatory role of MAP3 kinases MEKK-1 and NIK in IL-1 β -induced increase in intestinal epithelial TJ permeability, using filter-grown Caco-2 intestinal epithelial monolayers as an *in vitro* model of intestinal epithelium. Our data show for the first time that the IL-1 β -induced increase in intestinal epithelial TJ permeability is mediated by MEKK-1 activation. Our data also suggest that the IL-1 β effect on Caco-2 TJ barrier is due to MEKK-1 regulation of canonical NF- κ B pathways and MLCK gene activity.

Materials and Methods

Chemicals

Cell culture media (DMEM), trypsin, FBS, glutamine, penicillin, streptomycin, and PBS were purchased from GIBCO-BRL (Grand Island, NY). Anti-MEKK-1, NIK, IKK- α , IKK- β , I κ B- α , MLCK, and anti- β -actin antibodies were obtained from Sigma (St. Louis, MO). Anti-phospho-MEKK1, phospho-NIK, phospho-IKK α/β antibodies were obtained from Santa Cruz Biotechnology (Santa Cruz, CA). Anti-NF- κ B p65 and p100/p52 antibodies were purchased from Abcam (Cambridge, MA). Horseradish peroxidase-conjugated secondary antibodies for Western blot analysis were purchased from Invitrogen (San Francisco, CA). siRNA of MEKK-1, NIK, IKK α , IKK β , p65, and p100 and transfection reagents were obtained from Dharmacon (Lafayette, CO). All other chemicals were purchased from Sigma, VWR (West Chester, PA) or Fisher Scientific (Pittsburgh, PA).

Cell Cultures

Caco-2 cells (passage 20) were purchased from the American Type Culture Collection (Rockville, MD) and maintained at 37°C in a culture medium composed of DMEM with 4.5 mg/ml glucose, 50 U/ml penicillin, 50 U/ml streptomycin, 4 mmol/L glutamine, 25 mmol/L HEPES, and 10% FBS. The cells were kept at 37°C in a 5% CO₂ environment. Culture medium was changed every 2 days. Caco-2 cells were subcultured after partial digestion with 0.25% trypsin and 0.9 mmol/L EDTA in Ca²⁺- and Mg²⁺-free PBS.^{21,22}

Determination of Epithelial Monolayer Resistance and Paracellular Permeability

An epithelial voltometer (World Precision Instruments, Sarasota, FL) was used for measurements of the trans-epithelial electrical resistance (TER) of the filter-grown Caco-2 intestinal monolayers as previously reported. The effect of IL-1 β on Caco-2 paracellular permeability was determined using an established paracellular marker inulin (m.w. = 5000 g/mol).² For determination of mucosal-to-serosal flux rates of inulin, Caco-2-plated filters having epithelial resistance of 400–500 $\Omega \cdot \text{cm}^2$ were used. Known concentrations of inulin (2 $\mu\text{mol/L}$) and its radioactive tracer were added to the apical solution.

Assessment of Protein Expression by Western Blot Analysis

Caco-2 monolayers were treated with IL-1 β (10 ng/ml) for varying time periods. At the end of the experimental period, Caco-2 monolayers were immediately rinsed with ice-cold PBS, and cells were lysed with lysis buffer (50 mmol/L Tris-HCl, pH 7.5, 150 mmol/L NaCl, 500 μ mol/L NaF, 2 mmol/L EDTA, 100 μ mol/L vanadate, 100 μ mol/L PMSF, 1 μ g/ml leupeptin, 1 μ g/ml pepstatin A, 40 mmol/L paranitrophenyl phosphate, 1 μ g/ml aprotinin, and 1% Triton X-100) and scraped, and the cell lysates were placed in Microfuge tubes. Cell lysates were centrifuged to yield a clear lysate. Supernatant was collected, and protein measurement was performed using Bio-Rad Protein Assay kit (Bio-Rad Laboratories). Laemmli gel loading buffer was added to the lysate containing 10–20 μ g of protein and boiled for 7 minutes, after which proteins were separated on SDS-PAGE gel. Proteins from the gel were transferred to the membrane (*Trans-Blot* Transfer Medium, Nitrocellulose Membrane; Bio-Rad Laboratories) overnight. The membrane was incubated for 2 hours in blocking solution (5% dry milk in TBS-Tween 20 buffer). The membrane was incubated with appropriate primary antibodies in blocking solution. After being washed in TBS-1% Tween buffer, the membrane was incubated in appropriate secondary antibodies and developed using the Santa Cruz Western Blotting Luminol Reagents (Santa Cruz Biotechnology, Santa Cruz, CA) on the Kodak BioMax MS film (Fisher Scientific, Pittsburgh, PA).

siRNA of MEKK-1, NIK, IKK- α , IKK- β , p65, and p100/p52

Targeted siRNAs were obtained from Dharmacon, Inc. (Chicago, IL). Caco-2 monolayers were transiently transfected using DharmaFect transfection reagent (Lafayette, Co).²² Briefly, 5×10^5 cells per filter were seeded into a 12-well transwell plate and grown to confluency. Caco-2 monolayers were then washed with PBS twice and 1.0 ml Opti-MEM medium was added to the apical compartment of each filter and 1.5 ml were added to the basolateral compartment of each filter. siRNA of interest (5 nmol/L) and 2 μ l of DharmaFect reagent were preincubated in Opti-MEM. After 5 minutes of incubation, two solutions were mixed and incubated for another 20 minutes, and the mixture was added to the apical compartment of each filter. The IL-1 β experiments were carried out 96 hours after transfection. The efficiency of silencing was confirmed by Western blot analysis.

Nuclear Extracts and ELISA for Transcription Factor Activation

Filter-grown Caco-2 monolayers were treated with IL-1 β (10 ng/ml) for 30 minutes. Caco-2 monolayers were washed with ice-cold PBS, scraped, collected, and cen-

trifuged at 14,000 rpm for 30 seconds. The cell pellets were resuspended in 200 μ l of buffer A (in millimoles: 10 HEPES-KOH, 1.5 MgCl₂, 10 KCl, 0.5 DTT, and 0.2 PMSF [pH 7.9]), and incubated on ice for 15 minutes. After centrifugation at 14,000 rpm for 30 s, pelleted nuclei were resuspended in 30 μ l of buffer C (in millimoles: 20 HEPES-KOH [25% glycerol], 420 NaCl, 1.5 MgCl₂, 0.2 EDTA, 0.5 DTT, and 0.2 PMSF [pH 7.9]). After incubation on ice for 20 minutes, the lysates were centrifuged at 14,000 rpm for 20 minutes. Protein concentrations were determined using the Bradford method. The NF- κ B p65 and p52 DNA-binding assay was performed using *Trans-AM* ELISA-based kits from Active Motif according to the manufacturer's protocol. In brief, the binding reactions contained 1 pM biotinylated probe (Integrated DNA Technologies) and 5 μ g of nuclear extract in complete binding buffer with a total volume of 50 μ l. After 30 minutes of incubation, the solution was transferred to an individual well on 96-well plate and incubated for 1 hour. Appropriate antibody (2 μ g/ml) was added to the well to bind the target protein in nuclear extract. After incubation for 1 hour, the antibody was removed, and 100 μ l of horseradish peroxidase-conjugated secondary antibody was added to the well and incubated for 1 hour. Subsequently, 100 μ l of developing solution was added for 2–10 minutes, and 100 μ l of stop solution were added. The absorbance at 450 nm was determined using the SpectraMax 190 (Molecular Devices).

Immunostaining of NF- κ B p65 and p52 Proteins

Cellular localization of NF- κ B p65 and p52 was assessed by immunofluorescent antibody labeling.²³ At the end of the experimental period, filter-grown Caco-2 monolayers were washed twice in cold PBS and were fixed with 2% paraformaldehyde for 20 minutes. After being permeabilized with 0.1% Triton X-100 in PBS at room temperature for 20 minutes, Caco-2 monolayers were then incubated in blocking solution composed of bovine serum albumin and normal donkey serum in PBS for 1 hour. Cells were then labeled with primary antibodies in blocking solution overnight at 4°C. After being washed with PBS, the cells were incubated in FITC and Cy-3-conjugated secondary antibodies for 1 hour at room temperature. ProLong Gold antifade reagent (Invitrogen, CA) was used to mount the filters onto the coverslips. Immunolocalizations of NF- κ B p65 was visualized using a Confocal fluorescence microscope (LSM 510, University of New Mexico Imaging center) equipped with a Hamamatsu digital camera (Hamamatsu Photonics, Hamamatsu, Japan). Images were processed with LSM software (Zeiss, Germany).

RNA Isolation and Reverse Transcription

Caco-2 cells (5×10^5 per filter) were seeded into six-well transwell permeable inserts and grown to confluency. Filter-grown Caco-2 cells were then treated with appropriate experimental reagents for desired time periods.

At the end of the experimental period, cells were washed twice with ice-cold PBS. Total RNA was isolated using Qiagen RNeasy Kit (Qiagen, ML) according to the manufacturer's protocol. Total RNA concentration was determined by absorbance at 260/280 nm using SpectraMax 190 (Molecular Devices). The reverse transcription (RT) was carried out using the GeneAmp Gold RNA PCR core kit (Applied Biosystems, Foster City, CA). Two micrograms of total RNA from each sample were reverse transcribed into cDNA in a 40- μ l reaction containing 1 $\times$ RT-PCR buffer, 2.5 mmol/L MgCl₂, 250 μ mol/L of each dNTP, 20 U RNase inhibitor, 10 mmol/L DTT, 1.25 μ mol/L random hexamer, and 30 U multiscribe RT. The RT reactions were performed in a thermocycler (MyCycler, Bio-Rad, Hercules, CA) at 25°C for 10 minutes, 42°C for 30 minutes, and 95°C for 5 minutes.

Quantification of Gene Expression Using Real-Time PCR

The real-time PCRs were carried out using ABI prism 7900 sequence detection system and Taqman universal PCR master mix kit (Applied Biosystems, Branchburg, NJ) as previously described.⁴⁴ Each real-time PCR reaction contained 10 μ l RT reaction mix, 25 μ l 2 $\times$ Taqman universal PCR master mix, 0.2 μ mol/L probe, and 0.6 μ mol/L primers. Primer and probe design for the real-time PCR was made with Primer Express version 2 from Applied Biosystems. [The primers used in this study are as follows: MLCK specific primer pairs consisted of 5'-AGGAAGGCAGCATGAGGTTT-3' (forward), 5'-GCTTTCAGCAGGCAGAGGTAA-3' (reverse); probe specific for MLCK consisted of FAM 5'-TGAAGATGCTGGCTCC-3' TAMRA; the internal control GAPDH-specific primer pairs consisted of 5' CCACCCATGGCAAATTCC-3' (forward), 5'-TGGGATTTCATTGATGACCAG-3' (reverse); probe specific for GAPDH consisted of JOE 5'-TGGCACCGTCAAGGCTGAGAACG-3' TAMRA]. All runs were performed according to the default PCR protocol (50°C for 2 minutes, 95°C for 10 minutes, 40 cycles of 95°C for 15 s, and 60°C for 1 minute). For each sample, real-time PCR reactions were performed in triplicate, and the average threshold cycle was calculated. A standard curve was generated to convert the threshold cycle to copy numbers. Expression of MLCK mRNA was normalized with GAPDH mRNA expression. The average copy number of MLCK mRNA expression in control samples was set to 1.0. The relative expression of MLCK mRNA in treated samples was determined as a fold increase compared with control samples.

Transfection of MLCK DNA and Measurement of Promoter Activity

The MLCK promoter region was cloned using Genome-Walker system (Clontech, CA). A 2091-bp DNA fragment (–2109 to –18) was amplified by PCR.⁴⁴ The amplification condition was 1 cycle at 94°C for 2 minutes, followed

by 43 cycles at 94°C for 1 minute, 50°C for 1 minute, and 72°C for 2 minutes and 1 cycle at 72°C for 5 minutes. The resultant PCR product was digested with HindIII and KpnI and inserted into pGL3-basic luciferase reporter vector (Promega). The sequence was confirmed by DNA services at the University of New Mexico. MLCK promoter was transiently transfected into Caco-2 cells using transfection reagent lipofectamine 2000 (Life Technologies). Renilla luciferase vector (pRL-TK, Promega) was cotransfected with each plasmid construct as an internal control. Cells (5 $\times$ 10⁵ per filter) were seeded into a six-well transwell plate and grown to confluency. Caco-2 monolayers were then washed with PBS twice and 1.0 ml Opti-MEM medium was added to the apical compartment of each filter and 1.5 ml were added to the basolateral compartment of each filter. One microgram of each plasmid construct and 0.25 μ g pRL-TK or 2 μ l lipofectamine 2000 was preincubated in 250 μ l Opti-MEM, respectively. After 5 minutes of incubation, two solutions were mixed and incubated for another 20 minutes, and the mixture was added to the apical compartment of each filter. After incubation for 3 hours at 37°C, 500 μ l DMEM containing 10% FBS were added to both sides of the filter to reach a 2.5% final concentration of FBS. Subsequently, media were replaced with normal Caco-2 growth media 16 hours after transfection. Specific experiments were carried out 48 hours after transfection. At the completion of specific experimental treatments, Caco-2 cells were washed twice with 1 ml ice-cold PBS, followed by the addition of 400 μ l 1 $\times$ passive lysis buffer, incubated at room temperature for 15 minutes, scraped and transferred into an Eppendorf tube, and centrifuged for 15 seconds at 13,000 rpm in a microcentrifuge. Luciferase activity was determined using the dual luciferase assay kit (Promega). Twenty microliters of the supernatant were used for each assay. Luciferase values were determined by Lumat LB 9507 (EG&G Berthold, Oak Ridge, TN). The value of reporter luciferase activities were then divided by that of renilla luciferase activities to normalize for differences in transfection efficiencies. The average activity value of the

Figure 1. Time course effect of IL-1 β on Caco-2 MEKK-1 and NIK activation. **A:** Time course effect of IL-1 β (10 ng/ml) on Caco-2 MEKK-1 phosphorylation (total MEKK-1 was used for equal protein loading). **B:** Time course effect of IL-1 β on NIK phosphorylation (total NIK was used for equal protein loading). IL-1 β caused a time-dependent increase in Caco-2 MEKK-1 and NIK activation.

Figure 2. Effect of siRNA-induced MEKK-1 and NIK knockdown of IL-1 β -induced increase in Caco-2 TJ permeability. **A:** MEKK-1 siRNA transfection resulted in a near complete depletion in MEKK-1 protein expression as determined by Western blot analysis. **B:** MEKK-1 siRNA transfection prevented the IL-1 β -induced drop in Caco-2 TER (means $\pm$ SE, $n = 4$). * $P < 0.01$ versus control; ** $P < 0.01$ versus IL-1 β treatment. **C:** MEKK-1 siRNA transfection prevented the IL-1 β -induced increase in inulin flux (means $\pm$ SE, $n = 4$). * $P < 0.001$ versus control; ** $P < 0.001$ versus IL-1 β treatment. **D:** NIK siRNA transfection resulted in a near complete depletion in NIK protein expression. **E:** NIK siRNA transfection did not prevent the IL-1 β -induced drop in Caco-2 TER (means $\pm$ SE, $n = 4$). * $P < 0.01$ versus control. **F:** NIK siRNA transfection did not prevent the IL-1 β -induced increase in inulin flux (means $\pm$ SE, $n = 4$). * $P < 0.001$ versus control.

control samples was set to 1.0. The luciferase activity of MLCK promoter in treated samples was determined relative to the control samples.

Statistical Analysis

Results are expressed as means $\pm$ SE. Statistical significance of differences between mean values was as-

sessed with Student's *t*-tests for unpaired data and analysis of variance analysis whenever was required. All reported significance levels represent two-tailed *P* values. A *P* value of <0.05 was used to indicate statistical significance. Each experiment was performed in triplicate or quadruplicates ($n = 3$ or 4) and all experiments were repeated at least three times to ensure reproducibility.

Figure 3. Effect of IL-1 β (10 ng/ml) on Caco-2 NF- κ B pathways (p65 and p52) activation. **A:** IL-1 β caused the degradation of I κ B- α expression (30-minute experimental period) as assessed by Western blot analysis. **B:** ELISA-based DNA binding assay of NF- κ B p65. IL-1 β treatment caused a significant increase in Caco-2-NF- κ B p65 binding to the DNA probe. * $P < 0.001$ versus control. **C:** IL-1 β caused activation of p100 and generation of p52 (30 minutes experimental period) as assessed by Western blot analysis. **D:** ELISA-based DNA binding assay of NF- κ B p52. IL-1 β treatment caused a significant increase in Caco-2-NF- κ B p52 binding to the DNA probe. * $P < 0.001$ versus control.

Results

Role of MAP3 Kinases MEKK-1 and NIK in IL-1 β -Induced Increase in Caco-2 TJ Permeability

In the following studies, the effect of IL-1 β on activation of MAP3 kinases MEKK-1 and NIK was determined in Caco-2 cells. The time-course effect of IL-1 β (10 ng/ml) on MEKK-1 activation in filter-grown Caco-2 monolayers was assessed by MEKK-1 phosphorylation. IL-1 β produced a time-dependent increase in MEKK-1 phosphorylation in Caco-2 cells, starting at about 10 minutes and continuing up to 60 minutes as determined by phospho-MEKK-1 immunoblotting (Figure 1A). IL-1 β (10 ng/ml) also caused a rapid increase in NIK phosphorylation (peaking at 5 minutes) as assessed by phospho-NIK immunoblotting (Figure 1B). IL-1 β did not affect the total MEKK-1 or NIK level (Figure 1). These results indicated that IL-1 β induces a rapid phosphorylation of both MEKK-1 and NIK in Caco-2 cells.

Next, to assess the involvement of MEKK-1 and/or NIK in IL-1 β -induced increase in Caco-2 TJ permeability,

MEKK-1 and NIK expression was selectively knocked-down via siRNA transfection of filter-grown Caco-2 cells. The MEKK-1 siRNA transfection resulted in a near-complete depletion of MEKK-1 expression in Caco-2 cells (Figure 2A); the siRNA induced knockdown of MEKK-1 inhibited the IL-1 β -induced drop in Caco-2 TER and increase in mucosal-to-serosal flux of paracellular marker inulin (Figure 2, B and C). In contrast, the siRNA-induced knockdown of NIK (Figure 2D) did not affect the IL-1 β -induced drop in Caco-2 TER (Figure 2E) or the increase in *trans*-epithelial flux of inulin (Figure 2F). These data suggested that MEKK-1 but not NIK was required for the IL-1 β -induced increase in Caco-2 TJ permeability.

MAP3 Kinase Regulation of Canonical and Noncanonical NF- κ B Pathways

To examine the downstream processes involved in MAP3 kinase modulation of Caco-2 TJ permeability, the regulatory role of MEKK1 and NIK on NF- κ B pathway activation was investigated. MAP3 kinases have been shown to

Figure 4. Effect of siRNA-induced MEKK-1 knockdown of IL-1 β -activation of NF- κ B p65 and p52. **A:** MEKK-1 siRNA transfection prevented the IL-1 β -induced degradation of I κ B- α as assessed by Western blot analysis. **B:** MEKK-1 silencing inhibited the IL-1 β -induced binding of p65 to its binding site on the DNA probe as measured by DNA ELISA-binding assay. * P < 0.001 versus control; ** P < 0.001 versus IL-1 β treatment. **C:** MEKK-1 siRNA transfection did not prevent the IL-1 β generation of p52. **D:** MEKK-1 silencing did not inhibit the IL-1 β -induced binding of p52 to its binding site on DNA probe. * P < 0.001 versus control. **E:** Effect of MEKK-1 siRNA transfection on IL-1 β -induced nuclear translocation of p65 and p52 as viewed by confocal microscopy. Magnification, $\times 40$.

regulate the activation of both canonical and noncanonical NF- κ B pathways.⁴⁵ The activation of canonical pathway involves the degradation of inhibitory κ B ($I\kappa$ B) proteins, which results in the activation and nuclear translocation of NF- κ B heterodimer p65/p50.^{45,46} The activation of noncanonical pathway results in the phosphorylation and processing of p100 into p52 subunit and activation and nuclear translocation of p52/Rel B dimer.⁴⁵ The activation of canonical and noncanonical pathways is distinguished by the differential activation of dimers p65/p50 and p52/Rel B, respectively.^{45,46} In the following studies, IL-1 β effect on canonical and/or noncanonical pathway activation was examined in filter-grown Caco-2 cells. The IL-1 β effect on canonical pathway was determined by degradation of $I\kappa$ B- α and activation of NF- κ B p65 subunit, while the effect on noncanonical pathway was determined by generation and activation of p52 subunit. IL-1 β caused a rapid degradation of $I\kappa$ B- α and a cytoplasmic-to-nuclear translocation of p65 subunit in Caco-2 cells (Figures 3A and 4E). The IL-1 β -induced nuclear translocation of p65 was accompanied by an

increase in binding of activated p65 to the κ B binding site on the oligonucleotide probe as determined by ELISA binding assay (3B). IL-1 β also caused an increase in p52 generation (Figure 3C), nuclear translocation of p52 (Figure 4E), and an increase in binding of the activated p52 subunit to the κ B binding site on the oligonucleotide probe (Figure 3D). These results suggested that IL-1 β causes activation of both canonical and noncanonical NF- κ B pathways in Caco-2 cells.

Next, the regulatory role of MEKK-1 or NIK on IL-1 β induced activation of p65 or p52 subunits was determined by selective silencing of MEKK-1 or NIK. The siRNA induced knockdown of MEKK-1 prevented the IL-1 β -induced degradation of $I\kappa$ B- α and activation of p65 in Caco-2 cells but did not affect the generation or activation of p52 (Figure 4, A–E). In contrast, the silencing of NIK did not affect the activation of p65 but inhibited the generation and activation of p52 subunit (Figure 5, A–E). Together, these data suggested that MEKK-1 was responsible for the activation of canonical pathway and that NIK was involved in the activation of noncanonical path-

Figure 5. Effect of siRNA-induced NIK knockdown on IL-1 β activation of NF- κ B p65 and p52. **A:** NIK siRNA transfection did not prevent the IL-1 β -induced degradation of $I\kappa$ B- α as assessed by Western blot analysis. **B:** NIK silencing did not inhibit the IL-1 β -induced binding of p65 to its binding site on DNA probe as measured by DNA ELISA-binding assay. * $P < 0.001$ versus control. **C:** NIK siRNA transfection prevented the IL-1 β generation of p52. **D:** NIK silencing inhibited the IL-1 β -induced binding of p52 to its binding site on the DNA probe. * $P < 0.001$ versus control; ** $P < 0.001$ versus IL-1 β treatment. **E:** Effect of NIK siRNA transfection on IL-1 β -induced nuclear translocation of p65 and p52 as viewed by confocal microscopy. Magnification, $\times 40$.

way. To determine the role of canonical or noncanonical pathway in IL-1 β -induced increase in Caco-2 TJ permeability, the expression p65 or p100 subunit was selectively knocked-down by siRNA transfection. In the noncanonical pathway, p100 is phosphorylated and processed to generate the p52 subunit. The siRNA induced knockdown of p100 subunit (see Supplemental Figure S1A at <http://ajp.amjpathol.org>) or p52 subunit (data not shown) did not affect the IL-1 β -induced increase in paracellular permeability or drop in TER (see Supplemental Figure S1B at <http://ajp.amjpathol.org>) or increase in inulin flux (see Supplemental Figure S1C at <http://ajp.amjpathol.org>). In contrast, siRNA-induced knockdown of p65 completely inhibited the IL-1 β -induced drop in TER and increase in paracellular permeability, as previously reported by us,²² suggesting that the canonical pathway activation of p65 was required for the IL-1 β increase in Caco-2 TJ permeability.

Role of IKK Catalytic Subunits in IL-1 β -Induced Increase in Caco-2 TJ Permeability

The inhibitory κ B kinases (IKK) are important target of MAP kinases and play a crucial role in NF- κ B pathway activation; however, the regulatory role of IKK in TJ barrier function remains unknown. In the following studies, the involvement of IKK catalytic subunits—IKK- α and IKK- β —in IL-1 β -induced increase in Caco-2 TJ permeability was examined. The IL-1 β effect on IKK- α and IKK- β activation was determined by assessing their phosphorylation by immunoblot analysis. IL-1 β (10 ng/ml) treatment resulted in a rapid phosphorylation of both IKK- α and IKK- β (Figure 6A). The phosphorylation of IKK catalytic subunits reached the peak levels between 5–30 minutes and decreased thereafter to the baseline levels (Figure 6A). The time-course of IKK- α and IKK- β phosphorylation correlated closely with I κ B- α degradation (relative correlation coefficient $r = 0.905$) (Figure 6, B and

C). In the following studies, the role of MEKK-1 and NIK in IKK catalytic subunit activation was examined. The silencing of MEKK-1 inhibited the IL-1 β -induced activation of both IKK- α and IKK- β (Figure 6D). In contrast, NIK silencing did not affect the activation of either IKK- α or IKK- β (Figure 6E), confirming that MEKK-1 but not NIK was involved in the IL-1 β -induced activation of IKK catalytic subunits. To determine the involvement of IKK catalytic subunits in IL-1 β modulation of Caco-2 TJ permeability, the expression of IKK- α or IKK- β was selectively silenced via siRNA transfection. IKK- β siRNA transfection of Caco-2 cells resulted in a near-complete knockdown of IKK- β (Figure 7A). The IKK- β knockdown almost completely inhibited the IL-1 β -induced drop in Caco-2 TER (Figure 7B) and increase in paracellular permeability (Figure 7C), indicating that IKK- β was required for the increase in Caco-2 TJ permeability. The siRNA-induced depletion of IKK- α (Figure 7D) only partially inhibited the IL-1 β induced drop in Caco-2 TER and increase in inulin flux (Figure 7, E and F), suggesting a minor role for IKK- α . Next, the involvement of IKK catalytic subunits in the activation of canonical NF- κ B pathway was determined. The knockdown of IKK- β completely prevented the IL-1 β -induced degradation of I κ B- α (Figure 8A) and activation of p65 subunit (Figure 8B), whereas IKK- α silencing only partially inhibited the IL-1 β -induced I κ B- α degradation (Figure 8C) and p65 activation (Figure 8D). Together with the above data, these results suggested that IKK catalytic subunit regulation of Caco-2 TJ permeability was dependent on canonical pathway activation.

Role of MEKK-1 in the Regulation of MLCK Gene Expression

The above studies suggested that MEKK-1 plays a critical role in IL-1 β modulation of Caco-2 TJ barrier; however, the downstream target of MEKK-1 regulatory action remains unclear. Previous studies from our laboratory

Figure 6. Time-course effect of IL-1 β on Caco-2 IKK catalytic subunit activation. **A:** Time-course effect of IL-1 β (10 ng/ml) on Caco-2 IKK- α and IKK- β phosphorylation. IL-1 β caused a time-dependent increase in Caco-2 IKK- α and IKK- β activation. **B:** Time-course effect of IL-1 β on I κ B- α degradation (β -actin was used for equal protein loading). **C:** Graph of IKK- α /IKK- β activation versus I κ B- α degradation ($r = 0.905$). **D:** MEKK-1 siRNA transfection prevented the IL-1 β -induced phosphorylation of IKK- α and IKK- β as assessed by Western blot analysis. **E:** NIK siRNA transfection did not prevent the IL-1 β -induced phosphorylation of IKK- α and IKK- β .

Figure 7. Effect of siRNA-induced IKK- β or IKK- α knockdown of IL-1 β -induced increase in Caco-2 TJ permeability. **A:** IKK- β siRNA transfection resulted in a near complete depletion in IKK- β protein expression as determined by Western blot analysis. **B:** IKK- β siRNA transfection completely prevented the IL-1 β -induced drop in Caco-2 TER (means $\pm$ SE, $n = 4$). * $P < 0.01$ versus control; ** $P < 0.01$ versus IL-1 β treatment. **C:** IKK- β siRNA transfection prevented the IL-1 β -induced increase in inulin flux (means $\pm$ SE, $n = 4$). * $P < 0.001$ versus control; ** $P < 0.001$ versus IL-1 β treatment. **D:** IKK- α siRNA transfection resulted in a near complete depletion in IKK- α protein expression as determined by Western blot analysis. **E:** IKK- α siRNA transfection partially prevented the IL-1 β -induced drop in Caco-2 TER. * $P < 0.01$ versus control; ** $P < 0.01$ versus IL-1 β treatment. **F:** IKK- α siRNA transfection partially prevented the IL-1 β -induced increase in inulin flux (means $\pm$ SE, $n = 4$). * $P < 0.001$ versus control; ** $P < 0.001$ versus IL-1 β treatment.

indicated that the IL-1 β -induced increase in Caco-2 TJ permeability was mediated by an increase in gene and protein expression of downstream effector protein MLCK.²² In the following studies, we examined the possibility that MEKK-1 signaling cascade was involved in the regulation of MLCK gene activity. The IL-1 β effect on MLCK gene activity was assessed by transfection of Caco-2 cells with plasmid vector encoding the MLCK promoter region and luciferase reporter gene. The IL-1 β

treatment of filter-grown Caco-2 monolayers transfected with the plasmid vector encoding the MLCK promoter region resulted in an increase in MLCK promoter activity (Figure 9A). IL-1 β also caused an increase in MLCK mRNA expression and MLCK protein expression (Figure 9, B and C). The siRNA knockdown of MEKK-1 inhibited the IL-1 β -induced increase in MLCK promoter activity, MLCK mRNA expression, and MLCK protein expression (Figure 9, A–C). The NIK silencing did not affect the

Figure 8. Effect of siRNA IKK subunit knockdown of IL-1 β -induced activation of NF- κ B p65. **A:** IKK- β siRNA transfection completely prevented the IL-1 β -induced degradation of I κ B- α as assessed by Western blot analysis. **B:** IKK- β siRNA transfection inhibited the IL-1 β -induced binding of p65 to its binding site on the DNA probe as measured by DNA ELISA-binding assay. * $P < 0.001$ versus control; ** $P < 0.001$ versus IL-1 β treatment. **C:** IKK- α siRNA transfection partially prevented the IL-1 β -induced degradation of I κ B- α as assessed by Western blot analysis. **D:** IKK- α siRNA transfection partially inhibited the IL-1 β -induced binding of p65 to its binding site on the DNA probe as measured by DNA ELISA-binding assay. * $P < 0.001$ versus control; ** $P < 0.01$ versus IL-1 β treatment.

Figure 9. Effect of siRNA induced knockdown of MEKK-1 on IL-1 β -induced increase in MLCK gene activity and protein expression. **A:** MEKK-1 siRNA transfection resulted in a complete inhibition of IL-1 β -induced increase in MLCK promoter activity. * $P < 0.01$ versus control; ** $P < 0.001$ versus IL-1 β treatment. **B:** siRNA MEKK-1 transfection prevented the IL-1 β -induced increase in MLCK mRNA levels. * $P < 0.01$ versus control; ** $P < 0.01$ versus IL-1 β treatment. **C:** MEKK-1 siRNA transfection prevented the IL-1 β -induced increase in MLCK protein expression. **D:** NIK siRNA transfection did not prevent the IL-1 β -induced increase in MLCK promoter activity. * $P < 0.01$ versus control. **E:** siRNA NIK transfection did not prevent the IL-1 β -induced increase in MLCK mRNA levels. * $P < 0.01$ versus control. **F:** NIK siRNA transfection did not affect the IL-1 β -induced increase in MLCK protein expression.

increase in MLCK promoter activity or MLCK expression (Figure 9, D–F). These results suggested that MEKK-1 regulation of Caco-2 TJ permeability was mediated in part by up-regulation of MLCK gene activity and increase in MLCK protein expression.

Discussion

MAP3 kinases play a crucial role in the regulation of wide array of biological processes in the intestinal epithelial cells.^{47,48} However, their involvement in the regulation of intestinal epithelial TJ barrier remains unknown. The major aim of this study was to investigate the role of MAP3 kinases NIK and MEKK-1 in the regulation of intestinal epithelial TJ permeability, using filter-grown Caco-2 intestinal epithelial cells as an *in vitro* model of intestinal epithelium. Previous studies have shown IL-1 β to cause an increase in intestinal epithelial TJ permeability *in vitro* and *in vivo*^{21,49,50}; however, the specific regulatory pathways involved remain unknown. In this study, we examined the involvement of MAP3 kinases in the regulation of IL-1 β induced increase in Caco-2 intestinal epithelial TJ permeability. IL-1 β is a prototypical, proinflammatory cytokine that plays an integral role in the inflammatory process of the gut.^{11,12} The IL-1 β stimulation of biological actions is initiated by IL-1 β binding to the IL-1 receptor on the cell membrane surface, and subsequent IL-1 receptor complex recruitment of adaptor proteins including

IL-1 RacP, MyD88, Tollip, and tumor-necrosis factor receptor-associated factors 2 and 6.^{9,10} The recruitment of adaptor proteins to the IL-1 β receptor complex leads to the recruitment and activation of MAP3 kinases. MAP3 kinases play a crucial regulatory role in the activation of protein kinase pathways that lead to specific cellular responses.^{37–39} IL-1 β has been shown to induce activation of MAP3 kinases NIK and MEKK-1 in various cell types.^{40–43} Our data show that IL-1 β causes a rapid activation of both MEKK-1 and NIK in Caco-2 cells. The siRNA-induced silencing of MEKK1 expression abolishes the IL-1 β -induced increase in Caco-2 TJ permeability while NIK silencing does not have any effect. Thus, our results suggested that the IL-1 β -induced increase in Caco-2 TJ permeability was mediated in part by the activation of MEKK-1 pathway.

The MAP-3 kinases are important regulators of NF- κ B pathways. Two distinct signaling pathways have been described that lead to the activation of specific NF- κ B dimers: the canonical (or classical) and the noncanonical (or alternative) pathways.^{45,46} In the canonical pathway, ligand binding to the cell surface receptor leads to the ligand–receptor complex recruitment of membrane shuttle kinases that lead to the phosphorylation and degradation of inhibitory I κ B protein and activation of NF- κ B dimer p50/p65.⁴⁵ The ligand-induced activation of non-canonical pathway results in the phosphorylation of p100 subunit, leading to the generation and activation of

p52/Rel B dimer.^{45,46} Previous studies have shown NF- κ B to play an important role in the cytokine (TNF- α , IL-1 β , and IFN- γ)-induced increase in intestinal epithelial TJ permeability.^{21,23} In these studies, the inhibition of NF- κ B activity prevented the cytokine-induced increase in Caco-2 TJ permeability.^{21,23} In the present study, the role of NF- κ B pathways in MAP3 kinase regulation of Caco-2 TJ permeability was examined. Our data show that IL-1 β induces activation of both canonical and non-canonical NF- κ B pathways in Caco-2 cells. The IL-1 β activation of canonical pathway appeared to be mediated by MEKK-1 pathway as MEKK-1 knockdown inhibited the activation of p65 in Caco-2 cells without affecting p52 activation. Conversely, the NIK knockdown prevented the IL-1 β activation of p52 without having an effect on p65 activation. Together, these results suggested that MEKK-1 mediated the IL-1 β -induced activation of canonical pathway in Caco-2 cells and that NIK regulated the activation of noncanonical pathway. Consistent with the regulatory role of MEKK-1 in Caco-2 TJ permeability, the knockdown of p65 prevented the IL-1 β -induced increase in Caco-2 TJ permeability while p100 silencing did not have any effect. Thus, our data suggested that the IL-1 β -induced increase in Caco-2 TJ permeability was mediated by MEKK-1-induced activation of canonical pathway. Although NIK is primarily involved in the activation of noncanonical pathway, in certain conditions NIK has been shown to promote canonical pathway activation.⁵¹ In our studies, NIK did not play a role in canonical pathway activation or in Caco-2 TJ barrier regulation.

The inhibitory κ B kinase (IKK) complex plays an integral role in the regulation of NF- κ B activity.^{45,52} IKK complex consists of two catalytic subunits IKK- α and IKK- β and a regulatory subunit IKK- γ /NEMO. IKK- α and IKK- β are protein kinases and have enzymatic function.⁴⁹ IKK- γ is a scaffolding protein and does not have catalytic activity, but nevertheless plays a critical role in the formation of IKK complex.^{45,52} The activation of IKK complex leads to the phosphorylation of serine residues within the activation loop of IKK catalytic subunits. The activated IKK catalytic subunits phosphorylate I κ B proteins at Ser32 and Ser36 in I κ B α , leading to the eventual degradation of I κ B α by 26S proteasome and activation and nuclear translocation of NF- κ B p65/p50 dimer.⁴⁹ The involvement of IKK catalytic subunits in the regulation of TJ barrier function has not been previously reported. In this study, the role of specific IKK catalytic subunits in IL-1 β modulation of Caco-2 TJ barrier function was also examined. Our data indicated that both IKK- α and IKK- β were activated in response to IL-1 β stimulation. IKK- β appeared to be the catalytic subunit primarily responsible for the IL-1 β -induced increase in Caco-2 TJ permeability as IKK- β silencing completely inhibited, while IKK- α silencing only partially inhibited, the increase in Caco-2 TJ permeability. The involvement of IKK catalytic subunits in TJ barrier regulation correlated with their ability to induce I κ B α degradation and induce activation of p65 subunit. These findings are consistent with the involvement of canonical pathways in the regulation of Caco-2 TJ barrier. Our results also confirmed that MEKK-1 was responsible for

Figure 10. Proposed scheme of the intracellular pathways involved in IL-1 β -induced increase in intestinal epithelial tight junction (TJ) permeability. IL-1 β treatment resulted in activation of the MEKK-1 and NIK signaling cascades. MEKK-1 activation resulted in a step-wise activation of IKK and the canonical NF- κ B pathway and activation of the MLCK gene, culminating in the opening of the TJ barrier.

the IL-1 β -induced activation of IKK catalytic subunits. Together, our results suggested that the IL-1 β -induced increase in Caco-2 TJ permeability was regulated by MEKK-1-induced activation of IKK catalytic subunits; the activated IKK catalytic subunits presumably catalyzed the phosphorylation and subsequent degradation of I κ B α and activation of NF- κ B p65 subunits, leading to the opening of the Caco-2 TJ barrier (Figure 10).

As for the downstream mechanisms involved, MLCK has been shown to be a key effector protein regulating the cytokine induced opening of the intestinal epithelial TJ barrier.^{2,22,28} Previous studies have shown that the IL-1 β -induced increase in MLCK gene and protein expression was required for the increase in Caco-2 TJ permeability.²² Our present data show that MEKK-1 plays an important regulatory role in mediating the IL-1 β -induced activation of MLCK gene and increase in MLCK protein expression. Our results suggested that the IL-1 β -induced increase in MLCK gene and protein expression was due to MEKK-1 pathway up-regulation of MLCK gene activity (Figure 10). Presumably, MEKK-1-induced activation of IKK and NF- κ B resulted in NF- κ B-dependent activation of MLCK gene.⁴⁴ As shown previously, the increase in MLCK protein expression leads to MLCK-dependent opening of the TJ barrier (Figure 10).^{21,53} It has been shown that MLCK catalyzes the phosphorylation of myosin light chain (MLC); which in turn activates Mg²⁺-myosin ATPase leading to the contraction of perijunctional acto-myosin filaments and mechanical tension induced opening of the TJ barrier.^{54,55}

In conclusion, our data show for the first time that MAP3 kinase MEKK-1 (but not NIK) plays an integral role in IL-1 β modulation of Caco-2 intestinal epithelial TJ bar-

rier. Our results indicated that the IL-1 β -induced activation of MEKK-1 leads to the activation of IKK catalytic subunits IKK- β and IKK- α (Figure 10). The activated IKK catalytic subunits induce phosphorylation and degradation of I κ B, leading to the activation of canonical NF- κ B pathway and canonical pathway-dependent opening of the Caco-2 TJ barrier. Our data also indicate that MEKK-1 regulates the activation of MLCK gene activity, leading to the increase in MLCK protein expression. Together, these data provide important new insight into the role of MEKK-1 in regulating the IL-1 β modulation of intestinal epithelial TJ barrier (Figure 10).

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