

NAD⁺ Ameliorates Inflammation-induced Epithelial Barrier Dysfunction in Cultured Enterocytes and Mouse Ileal Mucosa*

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Abbreviations: iNOS, inducible nitric oxide synthase, LPS, lipopolysaccharide; cADPR, cyclic adenosine diphosphate-ribose; ADPR, adenosine diphosphate-ribose; NAD⁺, nicotinamide adenine dinucleotide; FD4, FITC-dextran mol wt ~4,000 Da; ZO, zonula occludens; TJ, tight junction

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Abstract

In the course of other experiments, we serendipitously observed that extracellular nicotinamide adenine dinucleotide (NAD⁺) ameliorated the development of epithelial hyperpermeability when monolayers of Caco-2 enterocyte-like cells were incubated with cytomix, a mixture containing IFN- γ , IL- β , and TNF- α . We sought to characterize the effects of NAD⁺ on inflammation-induced epithelial barrier dysfunction using Caco-2 monolayers that were exposed to cytomix in the absence or presence of NAD⁺ or other purine-containing molecules. Paracellular barrier function measured as the apical to basolateral passage of FITC-conjugated dextran (mol wt ~4,000 Da) was preserved in a concentration-dependent manner when immunostimulated Caco-2 cells were exposed to extracellular NAD⁺. Incubation with NAD⁺ prevented cytomix-induced derangements in the expression and localization of the tight junction proteins occludin and zonula occludens-1 in Caco-2 cells. Treatment of cytomix-stimulated cells with NAD⁺ also blocked NF- κ B activation, iNOS induction, and increased production of nitric oxide (NO \cdot). Ileal mucosal permeability to FD4 was increased in mice 18 h after lipopolysaccharide (LPS, endotoxin) injection, but treatment of endotoxemic mice with NAD⁺ ameliorated the development of gut mucosal hyperpermeability. Thus, extracellular NAD⁺ appears to ameliorate inflammation-induced intestinal epithelial barrier dysfunction by inhibiting NF- κ B activation and increased NO \cdot production.

A necessary function of the intestinal epithelium is the establishment of a selective barrier to allow the absorption of nutrients while restricting the uptake of toxic substances and microbes from the gut lumen. A major component of this epithelial barrier is the tight junction (TJ), a circumferential protein complex located at the apical/basolateral junction of opposing cells. The transmembrane proteins occludin and the claudin family are important TJ components that are thought to make homotypic contacts with proteins in the plasma membrane of the opposing cells (Tsukita *et al.*, 2001). The TJ complex is believed to be the point of cell-cell contact that presents the major barrier to paracellular transport (Anderson and Van Itallie, 1995).

Intestinal epithelial barrier function is compromised in a variety of inflammatory conditions including inflammatory bowel disease, cholestasis, hemorrhagic shock, and sepsis (Unno and Fink, 1998). Many mechanisms, including activation of myosin light chain kinase and excessive NO \cdot synthesis, have been proposed to explain the increase in paracellular permeability of intestinal epithelia following exposure to an inflammatory environment (Zolotarevsky *et al.*, 2002; Han *et al.*, 2003). NO \cdot interferes with the expression and subcellular localization of key TJ proteins including occludin, claudin-1, and peripheral membrane components of TJs including zonula occludens (ZO)-1, ZO-2, and ZO-3 (Han *et al.*, 2003).

During experiments designed to explore the biological effects of decreased intracellular NAD $^{+}$ content in immunostimulated human Caco-2 enterocyte-like cells (Khan *et al.*, 2002), we serendipitously observed that adding NAD $^{+}$ to the culture medium ameliorated derangements in epithelial barrier function caused by exposing these cells to a mixture of proinflammatory cytokines, called “cytomix.” This observation prompted us to review a growing literature regarding the role of NAD $^{+}$ and closely related compounds as signaling molecules. Several recent studies have shown that certain cell types secrete NAD $^{+}$ (Bruzzone *et al.*, 2001;

Romanello *et al.*, 2001) and/or respond to NAD⁺ in the extracellular milieu (Takasawa *et al.*, 1993a; Beers *et al.*, 1995; Guse *et al.*, 1995; Bruzzone *et al.*, 2001; Romanello *et al.*, 2001). Herein, we report that NAD⁺ is a signaling molecule that exhibited antiinflammatory properties when added at μ M concentrations to cultures of Caco-2 cells. Moreover, we present data showing that administration of NAD⁺ to mice ameliorated derangements in ileal mucosal barrier function induced by challenging the animals with LPS.

Materials and Methods

Reagents. All chemicals were from Sigma-Aldrich Chemical Co. (St. Louis, MO) unless otherwise noted. Recombinant human cytokines were obtained from Pierce-Endogen (Rockford, IN).

Animals. This research complied with regulations regarding animal care as published by the National Institutes of Health and was approved by the Institutional Animal Use and Care Committee of the University of Pittsburgh. Male 7-8 week old C57Bl/6J mice weighing 20-25 g were from Jackson Laboratories (Bar Harbor, ME). To induce a systemic inflammatory response, mice were injected i.p. with *Escherichia coli* (strain O111:B4) LPS (17 mg/kg) dissolved in 1.0 ml of PBS. Control animals (n=6) were injected with a similar volume of PBS. Mice receiving LPS were treated with an i.v. injection (100 μ l) of either PBS (n=6) or 132 mg/kg NAD⁺ in PBS (n=5) given 5 min before LPS injection, and this injection was repeated i.p. 12 h later. Groups of mice were anesthetized with 60–90 mg/kg sodium pentobarbital 18 h after the first injection with PBS or LPS. Segments of ileum were harvested to assess mucosal permeability to FD4 using an *ex vivo* everted gut sac method as previously described (Sappington *et al.*, 2002).

Cell Culture. Caco-2 human enterocyte-like cells were routinely maintained at 37 °C in DMEM with 10% FBS exactly as described (Han *et al.*, 2003). All cells were fed by changing the cell culture medium twice per week. For experiments, cells were grown on Transwell inserts (0.4 μ m pore size; Corning Costar Corp., Cambridge, MA) and used 14-17 days post-confluence. Paracellular permeability of Caco-2 monolayers was measured by determining the apical to basal clearance of FD4 (25 mg/ml) as described (Han *et al.*, 2003).

Immunoprecipitation and Western blotting. Western blots were performed as described (Han *et al.*, 2003). Briefly, Caco-2 cells were left unstimulated or exposed to various

agents. Cells were lysed in 1 ml of ice-cold NP-40-lysis buffer [25 mM 4-(2-hydroxyethyl)piperazine-1-ethanesulfonic acid (HEPES), pH 7.4, 150 mM NaCl, 4 mM ethylenediaminetetraacetic acid (EDTA), 25 mM NaF, 1% NP-40, 1 mM Na₃VO₄, 1 mM 4-amidinobenzylsulfonyl fluoride hydrochloride (APMSF), 10 µg/ml leupeptin, 10 µg/ml aprotinin] and insoluble material was collected by centrifugation and designated the NP-40 insoluble fraction. This pellet was dissolved in 100 µl of SDS-buffer (25 mM HEPES, pH 7.5, 4 mM EDTA, 25 mM NaF, 1% SDS, 1 mM Na₃VO₄) then diluted to 1 ml with NP-40-lysis buffer. The proteins were immunoprecipitated then subjected to Western blotting. Both steps were performed using anti-ZO-1 or anti-occludin polyclonal antibody from Zymed (South San Francisco, CA). Horseradish peroxidase-conjugated secondary antibodies were from Jackson ImmunoResearch Laboratories, Inc. (West Grove, PA). Blots were developed with the Enhanced Chemiluminescence detection system (Amersham, Piscataway, NJ), which contains a substrate that emits light following oxidation by the antibody conjugated peroxidase. The light emission is linear with respect to the amount of secondary antibody bound to the membrane, and the light emission is quantitated by exposing blue-light sensitive film (BioMax Film, Kodak, Rochester, NY). The images were captured using a Hewlett Packard (Palo Alto, CA) ScanJet 6300S.

Immunofluorescence. All procedures were performed at 0-4°C as described (Han *et al.*, 2003). Caco-2 cells growing on 24 well clear Transwell inserts were fixed and stained with a mouse anti-ZO-1 mAb (BD Translabs, Franklin Lakes, NJ) and a rabbit anti-occludin polyclonal antibody diluted 1:100. The secondary antibodies were 5 µg/ml FITC-conjugated affinity-purified donkey anti-mouse IgG and 15 µg/ml TRITC-conjugated affinity-purified goat anti-rabbit IgG (Jackson ImmunoResearch Laboratories).

Measurement of NO \cdot production and iNOS gene expression. NO \cdot production was measured and RT-PCR for iNOS mRNA and 18S RNA were performed as described (Sappington *et al.*, 2002). Relative expression of iNOS is reported after normalizing for 18S loading in Figure 4B.

Electrophoretic mobility shift assay (EMSA). Nuclear extracts were prepared and analyzed as described (Delude *et al.*, 1994). However, 38.3 μ l of 10% NP-40 was used, the nuclei were isolated by centrifugation at 310 g for 3 min, and nuclei were resuspended in 80 μ l of Buffer II [20 mM 2-amino-2-(hydroxymethyl)-1,3-propanediol (Tris)-HCl, pH 7.8, 5 mM MgCl, 320 mM KCl, 0.2 mM ethylene glycol-bis(2-aminoethylether)-N,N,N',N'-tetraacetic acid (EGTA), 0.5 mM dithiothreitol].

Statistical Analysis: Results are presented as means \pm SEM. Data were analyzed using analysis of variance (ANOVA). *P* values < 0.05 were considered significant.

Results

Extracellular NAD⁺ preserves intestinal epithelial barrier function. Caco-2 monolayers growing on permeable supports were left untreated or exposed to cytomix (1000 U/ml IFN- γ , 10 ng/ml TNF- α , and 1 ng/ml IL-1 β) for 24 or 48 h in the absence or presence of graded concentrations of NAD⁺ or adenosine. Paracellular permeability was determined by measuring the apical-to-basolateral flux of FD4. Epithelial permeability was unaffected when monolayers were incubated with NAD⁺ alone. Exposing Caco-2 cells to cytomix increased the clearance of FD4 across the monolayers (Fig. 1A). Addition of NAD⁺ ameliorated cytomix-induced epithelial hyperpermeability in a concentration-dependent manner. The protection afforded by 10 and 100 μ M NAD⁺ was statistically significant.

NAD⁺ is structurally related to a nucleoside, adenosine, that is known to exert antiinflammatory effects by binding to certain purinergic receptors (Sajjadi *et al.*, 1996); accordingly, we considered the possibility that activation of a receptor for adenosine was responsible for the amelioration of cytokine-induced hyperpermeability by NAD⁺. To investigate this idea, we exposed Caco-2 monolayers to cytomix in the absence or presence of 1-100 μ M adenosine (A in Fig. 1A). Adenosine had no effect on the induction of hyperpermeability by cytomix, suggesting that NAD⁺ was not simply an adenosine mimetic.

We next sought to determine whether treatment with NAD⁺ preserves intestinal mucosal barrier function *in vivo*. Mice were injected i.p. with LPS (n=11; 17 mg/kg) suspended in PBS (1 ml) or a similar volume of the PBS vehicle alone (n=6). Ileal segments were harvested 18 h later, and an *ex vivo* everted gut sac technique was used to measure mucosal permeability to FD4. Some mice that were injected with LPS were also treated with two 132 mg/kg doses of NAD⁺ (n=5) or—Other mice—were injected with similar volumes of the PBS vehicle (n=6). As expected

(Sappington *et al.*, 2003), injecting mice with LPS caused a significant increase in mucosal permeability to FD4 (Fig. 1 B). Treatment with two doses of NAD⁺ significantly ameliorated the increase in mucosal permeability induced by LPS in mice.

Extracellular NAD⁺ ameliorates derangements in TJ protein expression and cellular localization caused by cytomix. Proinflammatory cytokines are known to alter the expression and localization of several TJ proteins in cultured intestinal epithelial cells, including occludin, claudin-1, and ZO-1 (Youakim and Ahdieh, 1999; Han *et al.*, 2003). We tested whether NAD⁺ preserves the expression and targeting of TJ proteins in cells that were exposed to cytomix. Caco-2 cells were grown in the absence or presence of cytomix or NAD⁺ and NP-40-insoluble protein complexes were isolated after 48 h. The insoluble complexes, which represent insoluble cytoskeletal proteins and associated TJ proteins (Sakakibara *et al.*, 1997), were solubilized with SDS and subjected to immunoprecipitation followed by immunoblotting with anti-occludin or anti-ZO-1 antibodies. Exposure to cytomix for 48 h markedly decreased the amounts of occludin and ZO-1 present in TJs (Fig. 2A). When cells were simultaneously incubated with cytomix and 10 or 100 μM NAD⁺, nearly normal levels of these proteins were preserved in NP-40-insoluble complexes.

To again exclude the possibility that NAD⁺ was acting as an adenosine mimetic, we prepared NP-40-insoluble extracts from Caco-2 cells that were untreated or exposed to cytomix for 48 h in the absence or presence of 10 μM adenosine. Adenosine failed to prevent the decrease in NP-40-insoluble occludin or ZO-1 in Caco-2 cells exposed to cytomix (Fig. 2B). In other systems, the signaling effects of NAD⁺ are dependent on its conversion to cyclic ADP-ribose (cADPR) (Deaglio *et al.*, 2001; Okamoto and Takasawa, 2002). Therefore, we evaluated the effects of extracellular cADPR on cytomix-induced alterations in occludin expression. Co-

incubating cells with 10 or 100 μM cADPR preserved expression of occludin in Caco-2 cells exposed to cytomix (Fig. 2C). Remarkably, these concentrations of cADPR increased the expression of occludin to greater than baseline levels in immunostimulated Caco-2 cells. This effect of cADPR was specific for the cyclic nucleoside, since similar concentrations of (non-cyclic) ADP-ribose failed to preserve occludin expression in immunostimulated Caco-2 cells (data not shown).

Immunofluorescence microscopy was used to assess the structure of TJs in Caco-2 cells grown on Transwells. The cells were either left untreated (Fig. 3; Panel CON) or exposed to cytomix for 48 h in the absence (Panel C48) or presence (Panel N10) of extracellular NAD^+ (10 μM added at 0 and 24 h). The cells were permeabilized, fixed, and stained using FITC- and TRITC- conjugated antibodies directed against ZO-1 and occludin, respectively. In control confluent Caco-2 monolayers, these proteins both were predominantly localized to regions near cell-cell boundaries. Staining was even and continuous. Following treatment with cytomix, dispersed lesions appeared throughout the monolayer, in which immunostaining for ZO-1 at cell boundaries became faint, diffuse, and discontinuous. Exposure to cytomix also was associated with "ruffling" of ZO-1 and occludin immunostaining. Monolayers that were exposed to cytomix in the presence of 10 μM NAD^+ showed continuous staining of ZO-1 and occludin at cell-cell boundaries. However, some ruffling of the staining pattern was still evident.

Extracellular NAD^+ decreases iNOS expression in immunostimulated Caco-2 cells.

Changes in epithelial permeability caused by exposure to proinflammatory cytokines are due, at least in part, to excessive production of $\text{NO}\cdot$ (Unno *et al.*, 1995; Han *et al.*, 2003). Accordingly, we sought to determine whether NAD^+ decreased the production of $\text{NO}\cdot$ (as assessed by measuring NO_2^- and NO_3^- in culture supernatants) by Caco-2 cells exposed to cytomix. As

expected, cytomix increased the production of $\text{NO}\cdot$, but this effect was inhibited in a concentration-dependent manner by the addition of extracellular NAD^+ (Fig. 4A). Maximal inhibition was observed using $100\ \mu\text{M}\ \text{NAD}^+$; inhibition was decreased when NAD^+ was increased to $1000\ \mu\text{M}$. Cytomix increased expression of iNOS mRNA (Fig. 4B), and NAD^+ blocked this effect. Again, there was a biphasic response to NAD^+ treatment; $10\ \mu\text{M}$ provided maximal inhibition and higher doses inhibited less well.

Extracellular NAD^+ inhibits the activation of $\text{NF-}\kappa\text{B}$ in immunostimulated Caco-2 cells.

iNOS induction is dependent, in part, on activation of the transcription factor $\text{NF-}\kappa\text{B}$. Therefore, we tested the ability of NAD^+ to inhibit DNA binding by $\text{NF-}\kappa\text{B}$ in Caco-2 cells exposed to cytomix. Treatment of immunostimulated cells with $1\ \mu\text{M}\ \text{NAD}^+$ had no effect on cytomix-induced $\text{NF-}\kappa\text{B}$ activation, but co-incubation of cells with 10 or $100\ \mu\text{M}\ \text{NAD}^+$ clearly decreased $\text{NF-}\kappa\text{B}$ DNA binding (Fig. 4C) in nuclear extracts. NAD^+ alone had no effect on the nuclear levels of $\text{NF-}\kappa\text{B}$.

Discussion

NAD⁺ is a ubiquitous cellular constituent that is used by cells as an electron acceptor (or, in its reduced form, NADH, as an electron donor) in a wide variety of enzyme-catalyzed redox reactions. Accumulating data suggest that NAD⁺ also functions as a signaling molecule. Certain cell types secrete NAD⁺ (Bruzzone *et al.*, 2001; Romanello *et al.*, 2001) and/or respond to NAD⁺ in the extracellular milieu (Takasawa *et al.*, 1993a; Beers *et al.*, 1995; Guse *et al.*, 1995; Bruzzone *et al.*, 2001; Romanello *et al.*, 2001). cADPR can enter cells through either a CD38-dependent (Franco *et al.*, 1998) or a CD38-independent mechanism (Franco *et al.*, 2001). The resulting increase in intracellular cADPR concentration activates ryanodine-sensitive calcium channels on endoplasmic reticulum membranes causing the release of intracellular calcium stores to the cytoplasm (reviewed in Higashida *et al.*, 2001).

Herein, we showed that extracellular NAD⁺ preserves intestinal epithelial barrier function when enterocyte-like Caco-2 cells are exposed to an inflammatory milieu. NAD⁺ modulated the decreased expression and localization of the TJ proteins ZO-1 and occludin in immunostimulated Caco-2 cells. The salutary effects of NAD⁺ on TJ protein expression probably explain the ability of the compound to ameliorate cytomix-induced hyperpermeability. Treatment with NAD⁺ decreased the nuclear translocation of NF-κB, prevented accumulation of iNOS mRNA, and decreased the production of NO[•] in cytomix-stimulated Caco-2 cells. Interestingly, the concentration-response relationship for NAD⁺ was biphasic; increasing NAD⁺ concentrations up to 100 μM increased the protection afforded by the compound, but 1 mM NAD⁺ was less effective than 100 μM. Although we can only speculate about the basis for this complex dose-response relationship, it is worth noting that 5 mM NAD⁺ induces apoptosis in osteoblasts (Romanello *et al.*, 2001). Another possible explanation for this effect could be that 1mM NAD⁺

inhibits extracellular NADase activity, resulting in decreased production of the second messenger cADPR (Balducci and Micossi, 2002). Another observation worth noting is that 10 μM NAD^+ afforded maximal protection against decreased TJ protein expression and iNOS expression, the 100 μM NAD^+ was the most effective dose for preventing $\text{NO}\cdot$ production and decreased barrier function. Additional experiments will be necessary to better elucidate the dose-response relationships for the various pharmacological effects of extracellular NAD^+ . Taken together, our results suggest that NAD^+ acts as an antiinflammatory agent in cell culture, decreasing the responses of Caco-2 cells to exogenous proinflammatory cytokines. The biological relevance of these *in vitro* observations is emphasized by our results from *in vivo* studies, wherein we documented that NAD^+ can act as a pharmacological agent to ameliorate alterations in gut barrier function induced by injecting mice with LPS.

When Caco-2 cells were exposed to NAD^+ alone, we observed an increase in the expression of ZO-1 and occludin compared to control cells (Fig. 2A). We hypothesize that NAD^+ treatment of resting cells may decrease the basal production of $\text{NO}\cdot$. We have previously shown that pharmacologic inhibition or genetic ablation of iNOS causes increased expression of ZO-1 and occludin in ileum (Xiaonan Han, Mitchell P. Fink, Russell L. Delude, unpublished observations) and liver (Xiaonan Han, Mitchell P. Fink, Russell L. Delude, unpublished observations) of mice. Together these results suggest that basal $\text{NO}\cdot$ production decreases TJ proteins in epithelial cells.

The ecto-NADases CD38 and CD157 cyclize extracellular NAD^+ to form cyclic ADP-ribose (cADPR) and nicotinamide (Deaglio *et al.*, 2001; Okamoto and Takasawa, 2002). Both proteins are expressed by a variety of cells and may exist in soluble form in the plasma of normal humans (Deaglio *et al.*, 2001). Whether cleavage and cyclization of NAD^+ is necessary for its

antiinflammatory effects in Caco-2 cells *in vitro* or endotoxemic mice is unknown. This notion seems plausible, however, because we observed that extracellular cADPR, but not ADPR, preserved the expression of occludin in immunostimulated Caco-2 cells. This observation suggests, but does not prove, that extracellular NAD^+ acts via the formation of cADPR as a second messenger.

Intracellular and extracellular NAD^+ and/or cADPR may be important in regulating intracellular calcium concentration, $[\text{Ca}^{2+}]_i$, in a variety of cells (Takasawa *et al.*, 1993a; Beers *et al.*, 1995; Guse *et al.*, 1995; Bruzzone *et al.*, 2001; Romanello *et al.*, 2001). For example, cADPR-dependent changes in $[\text{Ca}^{2+}]_i$ may be important for regulating insulin release from pancreatic β -cells in response to alterations in exogenous glucose concentration (Hedeskov, 1980). Glucose increases the intracellular concentration of ATP in β -cells. ATP, in turn, inhibits the activity of an enzyme, cADPR-glycohydrolase, that converts cADPR to ADPR (Takasawa *et al.*, 1993b; Kato *et al.*, 1995; Tohgo *et al.*, 1997). Thus, according to this model, elevated glucose concentration increases intracellular cADPR concentration, leading to activation of ryanodine channels allowing the mobilization of calcium from intracellular stores. Interestingly, diabetogenic substances (*e.g.*, streptozotocin and alloxan) are known to damage DNA, which promotes activation of the enzyme, poly(ADP ribose) polymerase (PARP). Because PARP consumes NAD^+ as a substrate for poly-ADP-ribosylation, activation of PARP depletes cells of NAD^+ (Takasawa *et al.*, 1993a). Depletion of NAD^+ , the precursor of cADPR, inhibits signaling via this pathway. Interestingly, we have reported that cytomix induces intracellular NAD^+ depletion through an NO--dependent process in Caco-2 cells (Khan *et al.*, 2002). Whether cytomix-induced intracellular NAD^+ depletion affects intracellular cADPR levels and $[\text{Ca}^{2+}]_i$ in Caco-2 cells remains to be investigated.

Human osteoblast-like HOBIT cells differentiate into an osteoclast-like phenotype following exposure to extracellular NAD^+ (Romanello *et al.*, 2001). HOBIT cells express CD38 ADP-ribosyl cyclase activity, and NAD^+ treatment increases $[\text{Ca}^{2+}]_i$, presumably after conversion to cADPR. Resting HOBIT cells release NAD^+ into the culture medium, especially following mechanical perturbation. Since mechanical stimulation is an important factor driving osteoblast differentiation (Duncan and Turner, 1995), mechanical activation of NAD^+ release by osteoblasts may be an important paracrine signal driving differentiation of these cells (Romanello *et al.*, 2001).

An important question that is not addressed by our current studies or other publications concerns the availability of extracellular NAD^+ for signaling *in vivo*. Normal circulating levels of NAD^+ are 250 ± 30 nM and 140 ± 10 nM in humans and mice, respectively (Kim *et al.*, 1993). Thus, the normal circulating level of NAD^+ is substantially less than the concentration required to preserve barrier function in immunostimulated Caco-2 monolayers. Thus, the effects of NAD^+ reported here could be exclusively pharmacological. Alternatively, high local concentrations of NAD^+ could occur *in vivo* when the compound is released by cells in response to normal or pathological events. It is known, for example, that fibroblasts secrete NAD^+ (Bruzzone *et al.*, 2001), and fibroblasts are intimately associated with the basal side of the intestinal epithelial sheet (Parker *et al.*, 1974).

In conclusion, NAD^+ is capable of ameliorating structural and functional changes in intestinal epithelial cells induced by a proinflammatory milieu. Although NAD^+ *per se* is unlikely to be useful as a pharmacological agent, the potent anti-inflammatory effects of this ubiquitous endogenous molecule suggest the existence of a receptor on the surface of enterocytes that could serve as a useful drug target, leading to the development of novel agents for the

treatment of inflammatory conditions that affect gut epithelial function. Furthermore, our studies provide strong support for continued research into the function of NAD⁺ and, possibly, NAD⁺-derived cADPR in the function of intestinal epithelial cells during normal and pathologic conditions.

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Footnotes:

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Figure Legends

Figure 1. *Effect of extracellular NAD⁺ on the permeability of Caco-2 enterocytic monolayers and mouse ileal mucosa.* A) Caco-2 cells were incubated for 24 or 48 h under control conditions or with graded concentrations (μM) of either NAD⁺ (N) or adenosine (A). Fresh NAD⁺ was added to the wells after 24 h. CON indicates untreated monolayers and CYM indicates cytomix was added ($n = 6-12$ per condition). B) Mice were injected at T=0 h with PBS or LPS. Mice receiving LPS also received two injections of either PBS or NAD⁺ (132 mg/kg) dissolved in PBS. Ileal segments were harvested at T=18 h and permeability to FD4 measured. * indicates $p < 0.05$ versus control; ** indicates $p < 0.01$.

Figure 2. *Effect of extracellular NAD⁺ on the presence of ZO-1 and occludin in TJ complexes of Caco-2 cells.* A) Cells were either left untreated (CON) or exposed to cytomix (CYM) in the absence or presence of 10 (N10) or 100 (N100) μM NAD⁺ for 48 h. NP-40-insoluble protein was subjected to immunoprecipitation and Western blot analysis with an anti-ZO-1 or anti-occludin polyclonal antibody. B) Caco-2 cells were exposed to cytomix (CYM) in the presence or absence of 10 μM adenosine (ADN) C) Caco-2 cells were exposed to cytomix in the presence or absence of the indicated concentrations of cADPR. Western blots were repeated at least three times with reproducible results.

Figure 3. *Effect of NAD⁺ on ZO-1 (green) and occludin (red) localization in Caco-2 cells exposed to cytomix.* Caco-2 cells were either left untreated (CON) or exposed to cytomix for 48 h (C48). Some cells were exposed to cytomix for 48 h and treated at 0 and 24 h with 10 μM NAD⁺ (N10). Distribution and colocalization of ZO-1 (green) and occludin (red) was disrupted,

and this was blocked by NAD^+ . Magnification is 1000 \times in the left three columns. The images in the far right column show detail of the merged images that were increased to approximately 2000 \times using Adobe Photoshop. The bar is 10 μm .

Figure 4. *Effect of NAD^+ on cytomix-induced $\text{NO}\cdot$ production, iNOS expression and NF- κB activation.* A,B). Increased production of $\text{NO}\cdot$ and iNOS expression caused by cytomix was inhibited in a concentration-dependent manner by the addition of extracellular NAD^+ . C). Nuclear extracts were prepared from the cells 4 h after exposure to the agents (cytomix and NAD^+), and these extracts were used in an electrophoretic mobility shift assay using the murine NF- κB site from the κ light chain promoter. [All](#) figures are representative of 4 different experiments.

Fig 1. Han *et al.*

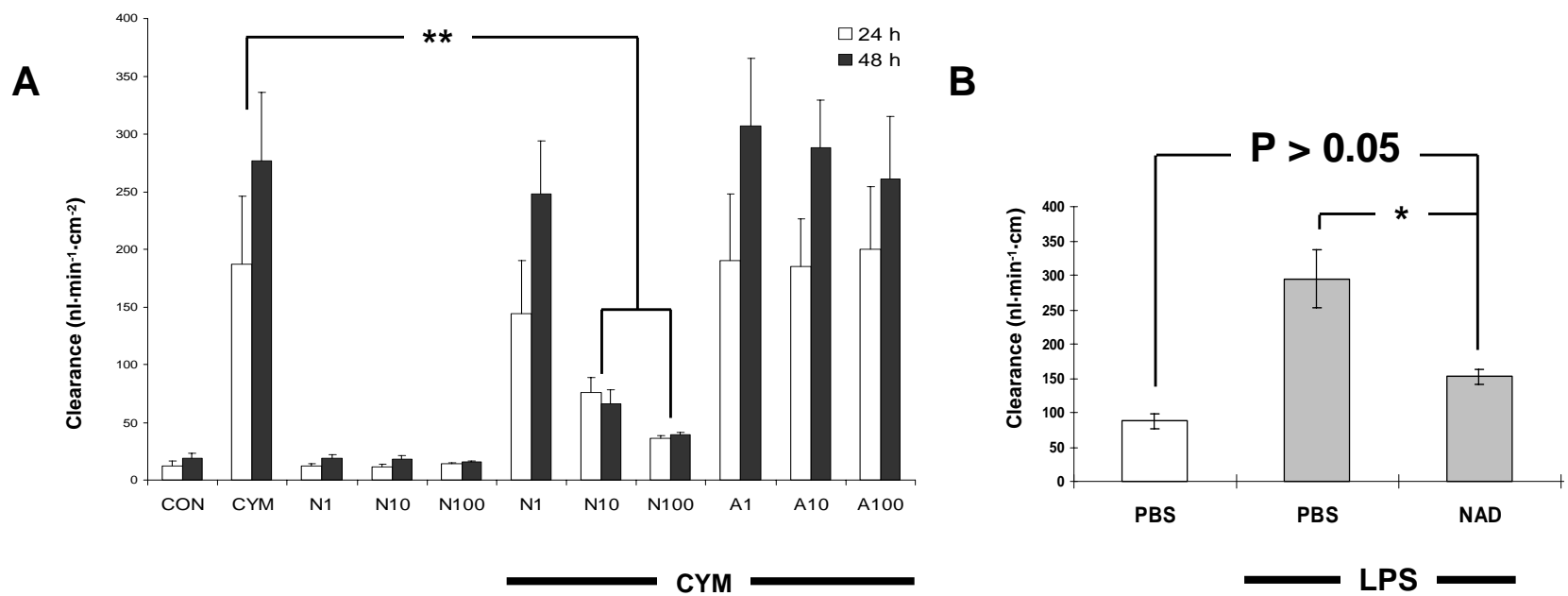


Fig 2. Han *et al.*

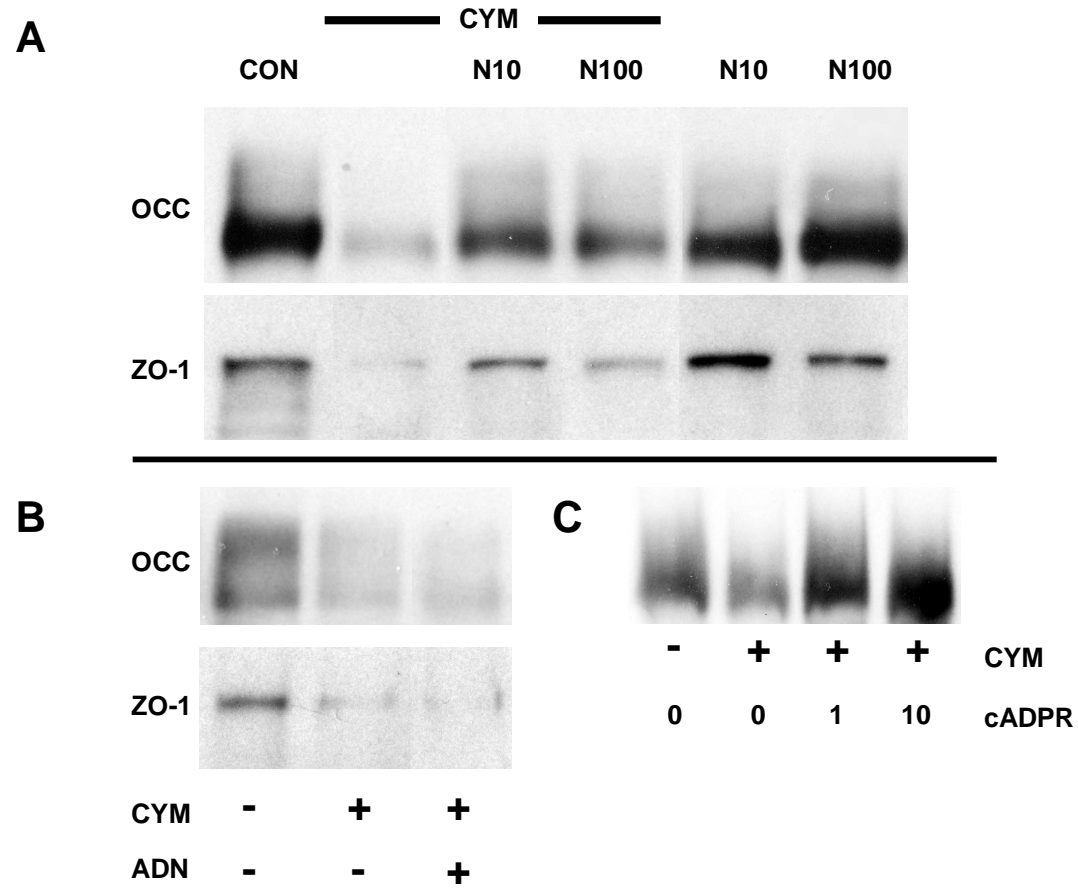


Fig 3. Han *et al.*

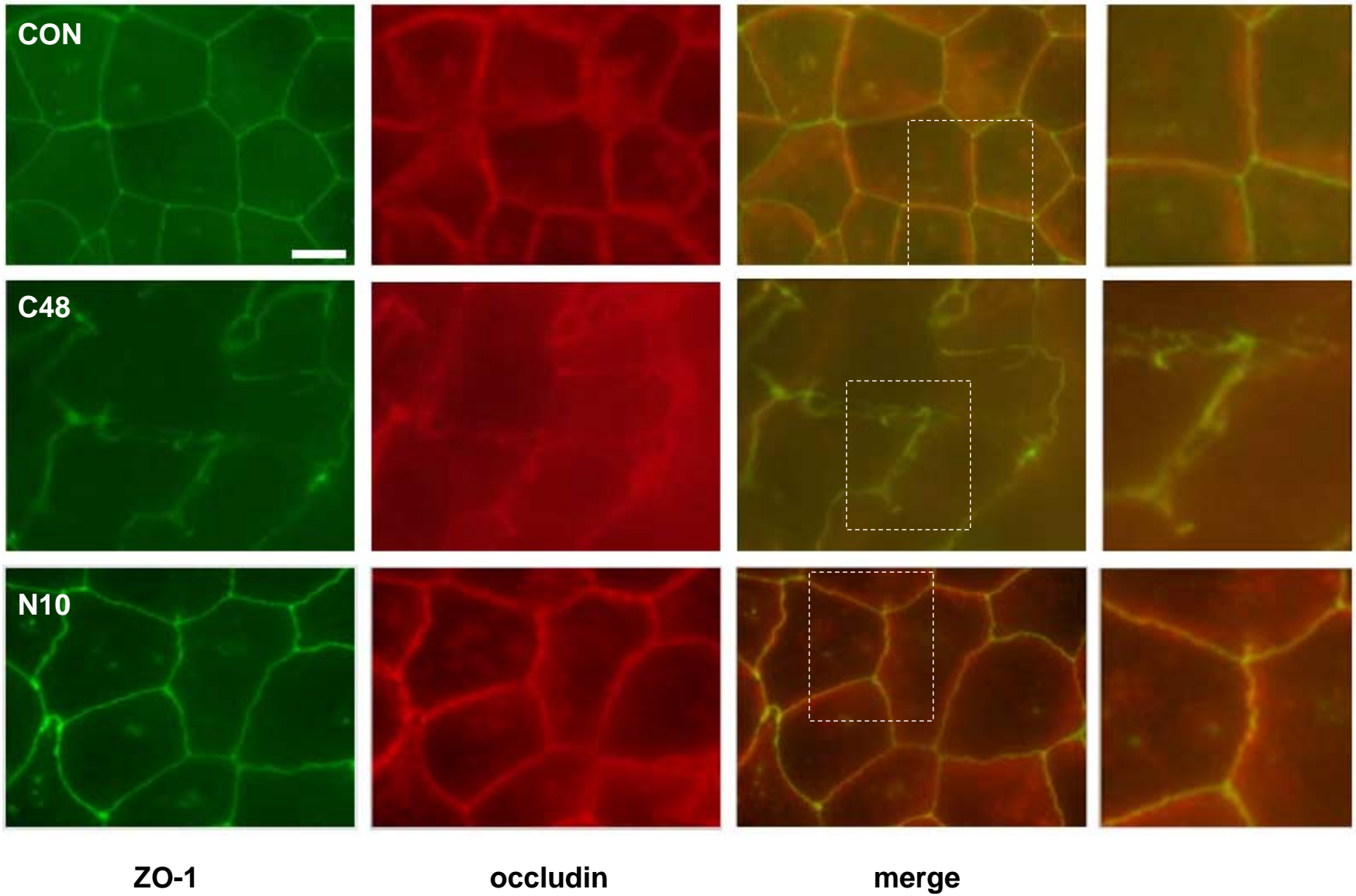


Fig 4. Han *et al.*

