Ubc9 overexpression and SUMO1 deficiency blunt inflammation after intestinal ischemia/reperfusion

Jörn Karhausen¹ · Joshua D. Bernstock² · Kory R. Johnson³ · Huaxin Sheng¹ · Qing Ma¹ · Yuntian Shen¹ · Wei Yang¹ · John M. Hallenbeck² · Wulf Paschen^{1,4}

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Abstract

The intestinal epithelium constitutes a crucial defense to the potentially life-threatening effects of gut microbiota. However, due to a complex underlying vasculature, hypoperfusion and resultant tissue ischemia pose a particular risk to function and integrity of the epithelium. The small ubiquitin-like modifier (SUMO) conjugation pathway critically regulates adaptive responses to metabolic stress and is of particular significance in the gut, as inducible knockout of the SUMO-conjugating enzyme Ubc9 results in rapid intestinal epithelial disintegration. Here we analyzed the pattern of individual SUMO isoforms in intestinal epithelium and investigated their roles in intestinal ischemia/reperfusion (I/R) damage. Immunostaining revealed that epithelial SUMO2/3 expression was almost exclusively limited to crypt epithelial nuclei in unchallenged mice. However, intestinal I/R or overexpression of Ubc9 caused a remarkable enhancement of epithelial SUMO2/3 staining along the crypt–villus axis. Unexpectedly, a similar pattern was found in SUMO1 knockout mice. Ubc9 transgenic mice, but also SUMO1 knockout mice were protected from I/R injury as evidenced by better preserved barrier function and blunted inflammatory responses. PCR array analysis of microdissected villus-tip epithelia revealed a specific epithelial contribution to reduced inflammatory responses in Ubc9 transgenic mice, as key chemotactic signaling molecules such as IL17A were significantly downregulated. Together, our data indicate a critical role particularly of the SUMO2/3 isoforms in modulating responses to I/R and provide the first evidence that SUMO1 deletion activates a compensatory process that protects from ischemic damage.

These authors contributed equally: Jörn Karhausen and Joshua D. Bernstock

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✉ Jörn Karhausen
jorn.karhausen@duke.edu

- ¹ Department of Anesthesiology, Duke University Medical Center, Durham, NC, USA
- ² Stroke Branch, National Institute of Neurological Disorders and Stroke, National Institutes of Health, Bethesda, MD, USA
- ³ Bioinformatics Section, Information Technology and Bioinformatics Program, Division of Intramural Research (DIR) NINDS/NIH, Bethesda, MD, USA
- ⁴ Department of Neurobiology, Duke University School of Medicine, Durham, NC, USA

Introduction

The integrity of the intestinal epithelium is crucial for preserving the body's homeostasis. Maintenance of this barrier generates substantial metabolic demands that arise in part from a need for rapid cell renewal, nutrient, electrolyte, fluid transport processes, and a state of 'physiologic' inflammation that ensures containment of bacteria and antigens [1]. Combined with an underlying vasculature, prone to shunting oxygen-rich blood away from the villus tip, these demands render the intestinal epithelium particularly sensitive to reductions of blood flow and resultant mucosal ischemia/hypoxia [2]. However, dramatic breakdown of intestinal integrity is generally prevented by extremely effective adaptive processes [2–4]. Further harnessing these protective responses bears great potential in clinical settings in which mucosal hypoperfusion has an important role, such as frank vascular occlusion, neonatal necrotizing enterocolitis [5], inflammatory bowel disease [2], and systemic inflammatory responses

associated with certain surgeries [6] and/or different forms of shock [7].

Posttranslational modification with small ubiquitin-like modifiers (SUMOs) is an effective way to dynamically influence responses to acute states of cellular stress through modification of transcription factors [8], enzyme and receptor activity [9], proteasomal degradation [10], cellular trafficking [11], protein aggregation [12], chromatin targeting [13], and protein–protein interactions [14]. Much of current literature has focused on SUMO modifications with respect to isolated proteins. This has led to conflicting reports with regard to global outcomes, as multiple components of a given pathway may be modified by SUMO in a distinct spatiotemporal sequence. Important new insights on the relevance of the SUMOylation system stem from studies manipulating Ubc9, the sole SUMO E2-conjugating enzyme. Here, rapid epithelial disintegration and death from intestinal failure after inducible global Ubc9 knockout (KO) draws attention to the unique dependency of the intestinal epithelium on a functioning SUMO machinery [15].

Vertebrates encode four different SUMO proteins: SUMO1, the highly homologous SUMO2 and 3 (often combined as SUMO2/3), and a still largely enigmatic SUMO4. Determining the specific contribution of these SUMO isoforms for health and disease has been extremely difficult, which is a major obstacle toward harnessing tissue protective effects of SUMOylation. For example, SUMO2/3-conjugated protein levels increase substantially after stimuli such as heat shock, oxidative stress, DNA damage, or hypoxia and ischemia [13, 16–19]; however, due to rapid cycles of SUMO2/3 conjugation and de-conjugation, these modifications are extremely dynamic. In addition, *in vivo* work is complicated by the redundancy between SUMO paralogs that compensates for the KO of SUMO1 or SUMO3, and by the fact that global KO of SUMO2 is embryonically lethal precluding functional studies of this particular isoform [20–22].

In the present work, we characterized the expression patterns of the major SUMO isoforms in the gut under physiological conditions and examined the effect of modulating SUMOylation on maintenance of gut integrity in a murine model of intestinal ischemia/reperfusion (I/R). To the best of our knowledge, this study provides the first evidence of differential regulation particularly of SUMO2/3 along the crypt villus axis, as well as of a significant induction of SUMO2/3 modification with I/R challenge. Work with mice overexpressing the sole E2 ligase, Ubc9, as well as from SUMO1-KO mice provide strong evidence that compensatory mechanisms targeting SUMO2/3 overexpression may be utilized to protect intestinal barrier integrity from I/R injury.

Materials and methods

Animal experiments

Animal work was approved by the Duke University and the National Institutes of Health (NIH)/National Institute of Neurological Disorders and Stroke Animal Care and Use Committees, respectively. Initial studies were performed in C57BL/6 mice obtained from Jackson Laboratories (Bar Harbor, ME). Experiments in the previously established Ubc9 transgenic mice (Ubc9Tg) were performed in the “H3” and “N2” founder lines [23]. SUMO3-KO animals had previously been developed in our laboratory. [22] SUMO1-KO animals were graciously made available by Dr Michael R. Kuehn (NIH) [20] and were backcrossed with C57BL/6 mice.

For mesenteric I/R, mice were anesthetized with 1% isoflurane supplemented by a one-time administration of 80 mg/kg ketamine, 5 mg/kg xylazine intraperitoneally, and 0.05 mg/kg buprenorphine subcutaneously. A midline laparotomy was then made and a 2–3 cm ileal loop isolated and rendered ischemic by placing aneurysm clips (Kent Scientific, Torrington, CT) on the peripheral branches of the superior mesenteric artery and across the intestine itself, in order to block flow through collaterals. Ischemia was maintained for 15, 30, or 45 min, after which the clamps were removed and the incision closed. During the procedure, the body temperature was maintained at 37.0 ± 0.3 °C rectal temperature using a feedback-controlled heating pad. For the reperfusion phase of injury, mice were allowed to recover in a heated box with supplemental oxygen for a variable amount of time (0.5, 1, 3, or 6 h). After killing, tissue was rapidly removed and snap frozen in liquid nitrogen for RNA and protein isolation or transferred into 4% paraformaldehyde (PFA) for paraffin embedding. To obtain snapshot images representative of *in vivo* SUMO expression and distribution, a subset of mice was killed and perfused with 0.9% NaCl (10 ml) followed by 4% PFA (30 ml) via transcardiac puncture. The intestines were then removed, placed in 4% PFA overnight, transferred to phosphate-buffered saline, and embedded in paraffin.

A further subset of animals undergoing I/R ($n = 4$ in 3 h and 6 h reperfusion groups) received 0.6 mg/g body weight fluorescein isothiocyanate (FITC)-dextran (4000 MW at a concentration of 80 mg/ml; Sigma-Aldrich, St. Louis, MO) by gavage 3 h before killing [24]. At killing, plasma was collected and the FITC concentration quantitated by fluorometry against a FITC-dextran standard curve.

Epithelial fractions for western blotting

Intestinal epithelial protein for western blotting was obtained by two methods. We obtained an epithelial-

enriched fraction by mechanically scraping the superficial cell layer from the opened luminal surface of small intestines. Sequential fractionation of cells along the villus/crypt axis was performed using a modified version of the low-temperature method described previously [25]. Briefly, ~5 cm of distal small intestine was swiftly removed after killing and immediately flushed with ice-cold Hank's balanced salt solution containing 20 mM *N*-ethylmaleimide (NEM) and 0.5 mM dithiothreitol (DTT) (Sigma), opened longitudinally, and cut into 10 mm pieces. Gut pieces were then brought into ice-cold chelating buffer consisting of 27 mM trisodium citrate, 5 mM Na₂HPO₄, 96 mM NaCl, 8 mM KH₂PO₄, 1.5 mM KCl, 0.5 mM DTT, 55 mM D-sorbitol, and 44 mM sucrose supplemented with 20 mM NEM. Tissue was incubated under agitation and then transferred to a tube with fresh buffer, whereas the remaining lifted-off epithelia were spun down at 12 × G, and the cell pellet frozen in liquid nitrogen and stored at -80 °C. This was repeated six times to obtain epithelial fractions from most apical (F1) to most basal (F7). PCR for fatty acid-binding protein 2 (Fabp2), Leucine-rich repeat-containing G-protein-coupled receptor 5 (Lgr5), and lysozyme transcripts (Supplemental Fig. 1A), and hematoxylin and eosin (H&E) stain (Supplemental Fig. 1B,C) and western blotting for the crypt markers lysozyme and β-catenin (Supplemental Fig. 1D) were used to verify that this technique produced a fractionated epithelial cell harvest from villus to crypt.

Western blotting

Cells and tissue samples were lysed in western blot lysis buffer as described previously [26], sonicated, and heat inactivated by boiling for 10 min. Proteins were electrophoresed on SDS-polyacrylamide gel electrophoresis gels (4–15%; Bio-Rad, Hercules, CA) and then transferred to polyvinylidene difluoride membranes (Bio-Rad). Membranes were blocked using Tris/HCl-buffered salt solution supplemented with 0.1% Tween 20 and 5% skim milk powder, and incubated with primary antibody overnight at 4 °C. Primary antibodies were as follows: polyclonal anti-SUMO1 (Novus Biologicals, Littleton, CO), polyclonal anti-SUMO2/3 (Covance, Denver, PA), monoclonal anti-Ubc9 (BD Bioscience, San Jose, CA), monoclonal anti-β-catenin (Thermo Fisher Scientific, Rockford, IL), polyclonal anti-lysozyme (Thermo), and polyclonal anti-IL17A antibody (Novus). After washing, membranes were incubated with anti-rabbit or mouse horseradish peroxidase conjugates (Bio-Rad) and targets visualized using the Supersignal West Pico Luminol Enhancer solution (Thermo). Membranes were stripped and re-probed with a mouse monoclonal anti-β-actin antibody (Sigma) as loading control. Quantification of band intensity

was performed using ImageJ software (NIH, Bethesda, MD).

Laser microdissection and capture of intestinal villus-tip epithelia

Microdissection was performed from 8 μm paraffin sections after deparaffination with *CitraSolv* (Fisher Scientific, Hanover Park, IL) and sequential ethanol baths. Microdissection was performed on an Axio Observer Z1 microscope equipped with PALM MicroBeam UV cutting beam (Carl Zeiss MicroImaging, München, Germany).

Intestinal samples from Ubc9Tg and wild-type (WT) littermates after 45 min of ischemia and 3 h of reperfusion were used for these studies. To obtain consistent samples, we limited tissue collected to villi with no more than an injury score of 2 according to the scoring system of Chiu et al. [27], i.e., to the villi with intact epithelial covering but some evidence of a developing sub-epithelial space. Samples were cut using a ×20/0.40, Korr LD Plan-Neofluar objective, yielding samples that consisted almost exclusively of villus-tip epithelia (Supplemental Fig. 2A–C). A total of 2,000,000 μm² per sample was collected.

RNA isolation from microdissected tissue and PCR array analysis

RNA from collected tissue sections was isolated using the RNeasy FFPE kit (Qiagen, Redwood City, CA) according to the manufacturer's specifications. Isolated RNA was then pre-amplified using the PreAMP cDNA synthesis kit in combination with PreAMP primer mix specific for the mouse cytokines and chemokines RT2 Profiler PCR array (catalog number: PAMM-150Z) and RT2 SYBR green quantitative PCR (qPCR) mastermix (all products Qiagen). PCR was performed on a LightCycler 480 (Roche Molecular Systems, Pleasanton, CA) following the RT2 Profiler program recommendations. Data analysis was based on the crossing point (C_p) determined by the second derivative maximum analysis method and was performed on the RT2 Profiler PCR array webportal.

Myeloperoxidase assay

Myeloperoxidase (MPO) activity was measured in snap-frozen tissue as described previously [28]. Briefly, tissue was homogenized in potassium phosphate buffer containing 0.5% hexadecyltrimethylammonium bromide (Sigma) and MPO activity was determined by measuring the oxidation of o-dianisidine in the presence of hydrogen peroxide (Sigma).

Gene transcription analysis (microarray and RT-qPCR analysis)

RNA was isolated using a commercially available kit (Macherey-Nagel, Bethlehem, PA). Microarray analysis was performed on four independent samples for each of six groups: sham (WT and Ubc9Tg), 45 min ischemia and 3 h reperfusion (WT and Ubc9Tg), and 45 min ischemia and 6 h of reperfusion (WT and Ubc9Tg). Total RNA was labeled on a sample-by-sample basis according to the manufacturer's guidelines for use with the Mouse Clariom S GeneChip (Affymetrix, Santa Clara, CA). Labeled cRNA were hybridized to these arrays in a blinded, interleaved manner. The Scanner 3000 (Affymetrix) was used in conjunction with GeneChip Operation Software (Affymetrix) to generate one.CEL file per hybridized cRNA. Next, the Expression Console (Affymetrix) was used to summarize the data contained across all.CEL files and generate 22,206 SST-RMA normalized transcript cluster expression values. Quality of this expression was challenged and assured via Tukey's box plot, covariance-based principal component analysis (PCA) scatter plot, and correlation-based heat map using functions supported in "R" (<http://www.cran.r-project.org>). To remove noise-biased expression values, lowess modeling was performed using expression values by class (coefficient of variation [CV] ~ mean expression). Lowess fits were then overplotted to identify the common low-end expression value in which the relationship between mean expression (signal) and CV (noise) deviated from linearity (mean expression value = 3.75). Subsequent flooring of measurements to this value reduced the overall number of gene-level measurements to 21,768. For transcript clusters not discarded, analysis of variance testing of the expression values across samples using class as the factor was performed under Benjamin-Hochberg false discovery rate multiple comparison condition, resulting in 6387 genes with a corrected $p < 0.05$. These genes were consecutively post hoc challenged via Tukey's honest significance difference test. The final number of 5487 differentially expressed genes were selected based on a p -value smaller than 0.05 and an absolute difference of means of greater than 1.5 fold. Next, annotations for these subset transcript clusters were obtained from NetAffx (Affymetrix), whereas enriched pathways and functions for the subset transcript clusters by class comparison were identified using Ingenuity Pathway Analysis (IPA) (Qiagen). For reverse transcriptase-qPCR (RT-qPCR) analysis, equal amounts of RNA were reversed transcribed using the High Capacity cDNA Reverse Transcription Kit (Applied Biosystems, Foster City, CA). qPCR was performed with the TaqMan Universal PCR Master Mix (Applied Biosystems) and FAM-labeled TaqMan Gene Expression Assays (Thermo)

for fucosyltransferase 2 (*Fut2*; Mm00490152_s1), suppressor of cytokine signaling 3 (*Socs3*; Mm00545913_s1), interleukin 6 (*Il6*; Mm0046190_m1), chemokine (C-X-C motif) ligand 2 (*Cxcl2*; Mm00436450_m1), endothelial cell selectin (*Sele*; Mm00441278_m1), platelet selecting (*Selp*; Mm01295931_m1), a disintegrin-like and metallo-peptidase (reprolysin type) with thrombospondin type 1 motif 1 (*Adamts1*; Mm01344169_m1), matrix metalloproteinase 8 (*Mmp8*; Mm00439509_m1), or tyrosine aminotransferase (*Tat*; Mm01244282_m1), each with a VIC-labeled β -actin primer/probe set (*Actb*; Mm02619580_g1) for internal calibration. All samples were tested in triplicate on a 7500 Fast Real-Time PCR System (Applied Biosystems) and analysed via the $2^{-\Delta\Delta CT}$ method.

Tissue staining and injury scoring

Sections were deparaffinized, H&E stained, and coded to conceal both genotype and experimental conditions before evaluation. Intestinal injury was quantified using the scoring system established by Chiu et al. [27]. The highest degree of injury in each field of view using a $\times 200$ magnification was noted to calculate average values for the entire length of the specimen. The final injury score resulted from two independently performed examinations of each sample. Naphthol AS-D chloroacetate esterase staining for tissue neutrophils was performed following the manufacturer's instructions (Sigma). Positively stained cells were quantified microscopically at $\times 200$ magnification in six unrelated regions of interest per section and reported in relation to the number of villi included in these regions.

Immunofluorescence

Sections were deparaffinized and heat-mediated antigen retrieval was performed in sodium citrate buffer. After blocking at room temperature, sections were incubated with the SUMO1, SUMO2/3, or Ubc9 antibodies specified above plus a monoclonal anti-e-cadherin antibody (BD Bioscience) for anatomical orientation. Labeling was achieved with goat anti-rabbit IgG or goat anti-mouse IgG conjugated to Alexa 488 or Cy3 (Thermo), and slides were mounted with ProLong anti-fade with 4',6-diamidino-2-phenylindole as counterstain (Thermo).

Statistical analysis

Statistical significance of differences among groups was determined by the two-sample t -test. Included bar graphs depict means $\pm$ SEM. Results were considered statistically significant if $p \leq 0.05$.

Fig. 1 SUMO1 and SUMO2/3 distribution in murine small intestine. Immunostaining of small intestinal sections from animals that were paraformaldehyde-perfused via transcardiac puncture. Sections were

stained as indicated with e-cadherin (green), DAPI (blue), and either SUMO1 (red: **a–c**) or SUMO2/3 (red: **d–f**). **a,d** Small intestinal section, $\times 200$; **b,e** villus tip, $\times 600$; **c,f** crypt, $\times 600$

Results

SUMO1 and SUMO2/3 are differentially expressed along crypt villus axis of small and large intestine

Immunofluorescence staining of intestinal samples revealed a distinct epithelial gradient of SUMO2/3 staining, with very low signal in villus tip but strong nuclear staining outlining the crypt epithelia (Fig. 1d–f). In contrast, SUMO1 exhibited some degree of gradient in epithelial cells, but this was less pronounced and SUMO1-positive epithelial cell nuclei were evident throughout the villus crypt axis (Fig. 1a–c and Supplemental Fig. 1D). Furthermore, submucosal cells both in the villus core as well as in the intestinal wall were robustly positive for both SUMO1 and SUMO2/3. This general staining pattern for SUMO1 and SUMO2/3 was found also in colonic samples, which indicates that a crypt-pronounced SUMO2/3 distribution exists in other regions of the intestine too (Supplemental Fig. 3).

We had previously shown that active SUMOylation processes typically increase particularly the nuclear SUMO signal vs. a more diffuse cytoplasmic stain of the free SUMO pool. [19] To verify that the observed, increased

nuclear staining pattern indeed signified increased SUMO conjugation of proteins, we performed western blot analysis of epithelial fractions from the small intestine (Supplemental Fig. 1D). Here, the high molecular smear representative of SUMOylated proteins was largely limited to crypt epithelial fractions when probing for SUMO2/3 and, to a lesser extent, for SUMO1. Interestingly, the sole SUMO ligase Ubc9 was expressed at higher levels in epithelial crypt cells too. Together, these data suggest that SUMO conjugation is highly regulated under baseline conditions in intestinal epithelia where it defines the crypt epithelial compartment.

Enhanced SUMO2/3 signal in the small intestinal villus tip area after I/R injury

We have previously shown that SUMO2/3 conjugation is rapidly induced in states of ischemia. [17, 19] To investigate whether intestinal epithelial SUMO2/3 signal was responsive to ischemic stress despite its tight restriction under baseline conditions, we subjected mice to mesenteric I/R injury. For this, we examined ischemic challenges that ranged from 15 to 45 min followed by a consistent reperfusion time of 30 min. As outlined in Fig. 2, I/R caused a

Fig. 2 SUMO2/3 staining expands into the villus-tip region after I/R. Sections were stained as indicated with e-cadherin (green: **a–d**) and SUMO2/3 (red: **a–h**) in animals subjected to sham procedure (**a,e**) or 15 (**b,f**), 30 (**c,g**), and 45 min (**d,h**) of ischemia. Additional sections were stained for e-cadherin (green: **i,k**) and SUMO1 (red: **i–m**) after sham procedure (**i,j**) or 45 min of ischemia (**k,m**). Reperfusion period was 30 min in all I/R animals. Magnification $\times 200$, size bar denotes 50 μm

rapid expansion of SUMO2/3 staining, initially as a spotty nuclear signal in more apical villus regions but with ischemic times longer than 30 min, as a robust epithelial SUMO2/3 staining within the villus tip. In contrast, differences in SUMO1 staining were much subtler due to the relatively high baseline villus tip signal (Fig. 2*i–l*).

Ubc9 overexpression leads to enhanced nuclear SUMO2/3 staining in intestinal villus epithelia of Ubc9Tg animals

Our previous work had suggested the Ubc9 may be a rate-limiting step in the SUMO conjugation process [29]. In agreement, elevated Ubc9 levels in Ubc9 transgenic (Ubc9Tg) animals translated into elevated baseline levels of SUMO1- and SUMO2/3-modified proteins in epithelial-enriched fractions taken from the small intestine

(Fig. 3*a–d*). In accordance, immunostaining of small intestinal sections documented a noticeable extension of both Ubc9 (Fig. 3*e,g*) and nuclear epithelial SUMO2/3 (Fig. 3*f,h*) staining into the villus tip in Ubc9Tg mice as compared with WT controls.

Ubc9Tg animals are protected from intestinal I/R injury

Next, we investigated whether enhanced SUMO conjugation was a protective response by using Ubc9Tg animals and their WT littermates in the mesenteric I/R model. We examined a consistent ischemic stimulus (45 min) with multiple reperfusion times to have endpoints at the height of injury (1 h reperfusion) and different stages of inflammation and healing (3 and 6 h reperfusion). Histological scoring indicated that the severity of damage inflicted after 45 min

Fig. 3 Overexpression of Ubc9 enhances baseline small intestinal epithelial SUMO2/3 expression. Mucosal scrapings from small intestines of WT or Ubc9Tg animals were analyzed by western blotting for **a** SUMO1, **b** SUMO2/3, **c** Ubc9, and **d** β -Actin protein levels. Markers denote molecular weight in kilo Dalton. Small intestinal sections

from **e.g** wild-type or **h.f** Ubc9Tg animals were stained for Ubc9 (red: **e,f**) or SUMO2/3 (red: **g,h**) as well as e-cadherin (green: **e-h**). Magnification $\times 200$. Inserts detail the villus tip staining for Ubc9 or SUMO2/3, respectively

of ischemia was comparable between WT and Ubc9Tg animals (Chiu score after 1 h reperfusion; Ubc9Tg 2.9 ± 1.5 vs. WT 2.7 ± 1.0). However, Ubc9Tg animals exhibited a significantly enhanced recovery 3 h (Chiu score 1.9 ± 0.6 vs. WT 2.7 ± 0.4) and 6 h (Chiu score 1.7 ± 0.8 vs. WT 2.7 ± 0.6) after reperfusion (Fig. 4a). In agreement, we observed significantly less leakage of gavaged FITC dextran into the serum in Ubc9Tg animals both 3 h (11 ± 4 μ g/ml vs. WT 49 ± 30 μ g/ml) and 6 h (9 ± 5 μ g/ml vs. WT 39 ± 23 μ g/ml) after reperfusion. (Fig. 4b). In addition, a marked reduction in the infiltration of inflammatory cells was seen in Ubc9Tg as compared with WT animals as evidenced by reduced MPO activity in Ubc9Tg 3 h (Ubc9Tg 1.9 ± 1.9 mU/mg tissue vs. WT 9.4 ± 3.5 mU/mg tissue) and 6 h (Ubc9Tg 0.5 ± 0.4 mU/mg tissue vs. WT 7.8 ± 3.5 mU/mg tissue) after reperfusion (Fig. 4c). A similar trend was observed when counting Naphthol AS-D Chloroacetate esterase-stained neutrophils. Few positive cells were encountered 1 h after reperfusion in either mouse strain, but in Ubc9Tg animals

these numbers remained relatively low (3 h reperfusion: 0.6 ± 0.3 neutrophils/crypt; 6 h reperfusion: 0.5 ± 0.4 neutrophils/crypt), whereas we observed significant inflammatory infiltration in WT animals (3 h reperfusion: 7.4 ± 3.0 neutrophils/crypt; 6 h reperfusion: 6.2 ± 1.7 neutrophils/crypt (**data not shown**). Therefore, although initial injury appeared comparable, Ubc9Tg recovered faster than their littermate controls, likely to be due to a blunted inflammatory response in the reperfusion period.

Reduced inflammatory gene expression in Ubc9Tg vs. WT animals after intestinal I/R

As a significant portion of SUMO target proteins are transcription factors, we next aimed to define the protective phenotype seen in Ubc9 animals using RNA microarray analysis. In this analysis, we tested whole-tissue intestinal samples of Ubc9Tg vs. WT animals undergoing sham procedure or 45 min of ischemia with either 3 or 6 h of

Fig. 4 Ubc9 Tg animals are protected from injury due to blunted inflammatory responses after intestinal I/R. **a** Sections from wild-type or Ubc9Tg animals undergoing 45 min of ischemia, and either 1, 3, or 6 h of reperfusion were stained by H&E and scored using the Chiu intestinal I/R score. $n = 6/\text{condition}$. **b** Plasma levels of gavage FITC dextran (in μg FITC dextran/ml plasma) as a measure of intestinal permeability from wild-type or Ubc9Tg animals after 45 min of ischemia and either 3 or 6 h of reperfusion. $n = 4/\text{condition}$. **c** Myeloperoxidase assay of intestinal samples collected after 45 min and 3 or 6 h of reperfusion. $n = 6/\text{condition}$, $*p \leq 0.05$. **d** Microarray analysis of transcriptional responses in Ubc9Tg (Tg) vs. wild-type littermates after

I/R. **e** Covariance-based principal component analysis (PCA) scatter plot depicts the Ubc9Tg vs. wild-type sample-to-sample relationships by time using the union set of 5487 genes identified differentially expressed for at least one possible sample type comparison. **f** Venn diagram summarizes the number of differential expressed genes observed between Ubc9Tg vs wild type that are either exclusive by time or shared across time. The top 10 significantly enriched **f** functions and **g** pathways returned from the Ingenuity Pathway Analysis (IPA) tool when provided the list of 474 genes differentially expressed between Ubc9Tg and wild type subjected to 45 min of ischemia and 6 h of reperfusion. $n = 4/\text{condition}$

reperfusion (raw data have been deposited in NCBI's Gene Expression Omnibus [<http://www.ncbi.nlm.nih.gov/geo/>], GEO Series accession number GSE96733). A total of 22,206 gene-level expression measurements were generated per sample from which our analysis identified a final set of 5487 as being differentially expressed (for approach and selection criteria, see Materials and Methods section of the manuscript). PCA analysis (Fig. 4d) demonstrated separation between sham and the different I/R treatment groups but differences between respective WT and Ubc9Tg animals only in the 6 h reperfusion cohort (distribution of differentially regulated genes is shown in the Venn diagram, Fig. 4e). From this group, we selected nine differentially regulated genes for validation by qPCR (Table 1) and found

the same overall trend, although differences in fold changes were noted. IPA pathway and function analysis of differentially regulated genes in this group revealed a strong enrichment in functions associated with inflammatory cell recruitment such as “migration of cells,” “cell movement of myeloid cells/leukocytes/granulocytes/phagocytes,” or “inflammatory response” (Fig. 4f). Consequently, the most significantly enriched pathways when comparing Ubc9Tg and WT animals after 6 h of reperfusion were “granulocyte adhesion and diapedesis,” as well as pathways associated with cytokines such as IL15, IL6, and IL17A (Fig. 4g).

These results demonstrate that Ubc9 overexpression is associated with a complex reprogramming of the inflammatory response after I/R.

Table 1 Verification of microarray data by qPCR analysis

Gene symbol	Gene ID	qPCR	Microarray
Fut2	14,344	− 3.30	− 18.31
Socs3	12,702	− 3.89	− 4.04
Il6	16,193	− 3.64	− 7.15
Cxcl2	20,310	− 3.16	− 3.33
Selp	20,344	− 3.89	− 4.99
Sele	20,339	− 2.88	− 3.26
Adamts1	11,504	− 2.01	− 2.88
Mmp8	17,394	− 3.82	− 2.57
Tat	234,724	− 5.04	− 16.27

Relative expression of nine selected genes in Ubc9Tg vs. WT animals. $n = 6$ for qPCR and $n = 4$ for microarray.

Laser microdissection and capture reveals modulated chemokine responses in villus-tip epithelial cells of Ubc9Tg animals

Given the fact that histological changes were evident 3 h after reperfusion, whereas our microarray from whole tissue identified the most relevant changes only 6 h after reperfusion, we hypothesized that earlier responses in villus-tip epithelia may have been under-represented in the whole tissue expression analysis. The villus tip not only constitutes the location of the most pronounced injury but also that of the most apparent increase in SUMO expression with either I/R or Ubc9 overexpression. Therefore, we obtained tissue highly enriched for villus tip epithelia by laser microdissection and capture, and we examined the expression of major chemokine and cytokines via PCR array. Figure 5a,b show scatter and volcano plots, which represent significantly regulated genes in villus-tip samples from Ubc9Tg animals vs. WT littermates 3 h after reperfusion. Five genes (out of 84) were induced at least 2-fold (p -value < 0.05), whereas 6 genes were downregulated at least 2-fold (p -value < 0.05) (Fig. 5c). These results show that SUMO overexpression modulates early epithelial inflammatory signaling with a marked suppression of chemotactic factors such as Cxcl9, Cxcl16, Ccl5, Ccl20, Il17a, and Il27 in line with the reduced inflammatory infiltration observed in these animals. As IL17A constitutes a major determinant of chemotactic responses, we verified IL17A protein levels in these animals. Notably, we found that 6 h after reperfusion, IL17A protein was significantly induced in ischemic vs. non-ischemic small intestinal loops, and confirmed that this response was blunted in Ubc9Tg animals (Fig. 5d). Together, these findings further support that SUMOylation controls inflammation after I/R injury and identify a specific epithelial contribution to inflammatory signaling in this context.

SUMO1-KO animals display compensatory overexpression of SUMO2/3 in intestinal epithelia and are partially protected from I/R injury

The specific contribution of SUMO2/3 vs. SUMO1 in adaptive responses to injury has never been examined in vivo. To address this, we first defined the relative distribution of the major SUMO isoforms in the intestines of SUMO1 and SUMO3-KO animals. No phenotypical abnormalities have been described in these animals [20–22] and, indeed, when we stained SUMO3-KO animals under baseline conditions, we found no differences in either SUMO1 or SUMO2/3 distribution compared with WT littermates (Supplemental Fig. 4). However, unchallenged SUMO1-KO animals unexpectedly displayed an enhanced nuclear SUMO2/3 expression that clearly extended into the epithelium of the villus tip and thus mirrored findings from Ubc9Tg animals or after I/R injury (Supplemental Fig. 5). When exposed to the mesenteric I/R model (45 min of ischemia and 6 h of reperfusion), minor differences in overall injury severity were observed between SUMO1-KO vs. WT littermates (Fig. 6a). However, SUMO1-KO animals displayed better intestinal barrier preservation after FITC dextran gavage (SUMO1-KO $9.2 \pm 3.2 \mu\text{g}$ FITC dextran/ml plasma vs. WT $39.1 \mu \pm 18.1 \mu\text{g}$ FITC dextran/ml plasma) (Fig. 6b), together with a blunted inflammatory response as evidenced by the significantly reduced MPO activity (SUMO1-KO $1.35 \pm 2.91 \text{ mU/mg}$ vs. WT $16.26 \pm 5.97 \text{ mU/mg}$) (Fig. 6c). In agreement, staining of cells from the granulocyte lineage with Naphthol AS-D Chloroacetate revealed a numerous infiltration of polymorphnuclear cells in seemingly well-preserved villi adjacent to sites of epithelial injury in WT mice (Fig. 6d,e) but only isolated neutrophils in SUMO1-KO mice (Fig. 6f). Lastly, and consistent with the transcriptional pattern observed in Ubc9Tg animals, we observed a blunted response in the selected group of genes involved in inflammatory cell recruitment (Fig. 6g). Together, this indicates that compensatory SUMO2/3 overexpression in SUMO1-KO is sufficient to regulate important inflammatory responses in I/R injury. However, the more complete protective phenotype in Ubc9Tg animals suggests that SUMO1 and/or SUMO-independent effects of Ubc9 may also contribute to adaptive responses.

Discussion

SUMO conjugation allows for rapid cellular reprogramming in states of severe metabolic stress such as I/R. Under such conditions, especially SUMO2/3-conjugation rapidly increases [29] and evidence suggests an important adaptive and cell-protective role through regulation of metabolism,

Fig. 5 Modulation of villus-tip epithelial responses in Ubc9Tg animals after I/R. Intestinal epithelial villus-tip epithelia from ischemic regions of wild type and Ubc9Tg (45 minutes of ischemia and 3 h of reperfusion) were isolated by laser microdissection and capture and RNA was isolated to perform a cytokine targeted PCR array. **a** Scatter and **b** Volcano blot outline the distribution of regulated genes; significantly up- and downregulated genes are listed in **c**. **d** IL17A protein levels in non-ischemic small intestinal loops (NI) and loops subjected to ischemia/reperfusion (IR; 45 min ischemia and 6 h or reperfusion). Quantification was performed by densitometry and is represented as the ratio of IL17A to β -Actin band intensity. $n = 5$; $*p \leq 0.05$

hypoxia adaptive responses, inflammation, RNA processing, and SUMOylation-dependent ubiquitin conjugation [10, 30–38]. Our present study adds important knowledge to the field by showing a distinct SUMO2/3, and to a lesser degree SUMO1, crypt-to-villus gradient under physiologic conditions and, importantly, a marked expansion of epithelial SUMOylation in an I/R injury model. Mice over-expressing Ubc9, and—unexpectedly—mice lacking SUMO-1, demonstrated enhanced nuclear SUMO2/3 signal in intestinal epithelial cells and improved outcomes after mesenteric I/R, primarily through blunting of inflammatory responses. This data provides powerful support that SUMOylation, in particular with the SUMO2/3 isoforms, is a crucial protective response after intestinal I/R injury.

Although SUMOylation is clearly essential for intestinal homeostasis [15] as well as for barrier integrity and inflammatory control during bacterial invasion [39], the intestinal expression pattern of the SUMO paralogs has never been studied in detail. Hypothesizing that SUMOylation events may perform a protective role in this

metabolically challenged tissue, we expected a strong SUMO signal in the apical epithelium in parallel with the steep oxygen gradient that exists along the crypt/villus axis [2]. However, the exact opposite was the case, with strong SUMO2/3 signal in the connective tissue and the epithelial crypt but little staining of villus-tip epithelia. Results from Demarque et al. [15] bear impressive evidence of the functional importance of our findings. Here, the induced loss of Ubc9 in adult animals led to the depletion of the small intestinal proliferative compartment, to massive disturbances of intestinal epithelial architecture, and ultimately to rapid demise from intestinal failure [15]. Therefore, SUMOylation is an indispensable and, as shown by our data, a highly regulated mechanism for small intestinal epithelial organization. What remains unknown is how SUMOylation is regulated in this setting and what its specific functions are. It will be extremely interesting to integrate the observed SUMO2/3 conjugation pattern with existing knowledge on crypt-villus gradients of regulators of intestinal epithelial renewal and differentiation [40].

Fig. 6 SUMO1 knockout (SUMO1-KO) animals display a blunted inflammatory response to intestinal I/R injury. **a** H&E-stained section from wild-type and SUMO1- KO animals were scored using the Chiu intestinal I/R score. **b** Plasma levels of gavaged FITC dextran (in μg FITC dextran/ml plasma) from wild type or SUMO1-KO animals. **c** Myeloperoxidase assay. **d–f** Representative images from Naphthol AS-D Chloroacetate esterase staining of intestinal sections from wild-type (**d,e**) or SUMO1-KO animals (**f**) after I/R injury. Images were

taken from relatively intact sections adjacent to areas of significant injury. Arrows denote infiltrating neutrophils. Magnification $\times 400$. **g** qPCR from post-ischemic small intestines in SUMO-1 KO and WT mice. Graphs represent fold change over sham-treated wild-type animals and notes the fold difference of response between SUMO1-KO and WT, consistent with Table 1. Assays were performed on wild-type ($n = 4$) or SUMO1-KO ($n = 5$) animals subjected 45 min of ischemia and 6 h of reperfusion. $*p \leq 0.05$

Interestingly, the physiologic restriction of SUMO2/3 signal was rapidly lifted after I/R, which led to a severity-dependent expansion of SUMOylation into the apical epithelium. The villus tip is particularly exposed to metabolic challenges such as hypoxia [2] and free radical generation [41], and therefore injury following I/R is typically most pronounced here [27]. Increased global SUMOylation, particularly of SUMO2/3 has been reported in different organs after I/R including the kidney, heart, and brain [17, 19, 29, 42, 43], and is thought to be part of an overall cytoprotective response. However, tools capable of defining the specific contribution of SUMOylation and of the individual SUMO isoforms to protective responses in this context have been lacking. The SUMO E2 conjugase Ubc9 is considered a gatekeeper within the SUMOylation

process, as evidenced by the fact that SUMO1/2/3 overexpression alters free SUMO levels but not those of SUMO-modified proteins [29]. In our current work, we now observed that Ubc9 overexpression induced SUMO1 and SUMO2/3 signal, particularly in the villus tip, and that it conveyed significant protection in a mesenteric I/R model. Together with previous work that correlated expression levels of Ubc9 with SUMO conjugation levels in the brain and, importantly, with increased tolerance to focal cerebral ischemia [23], this data strongly support the idea that global SUMOylation levels may be harnessed to protect against I/R injury and related inflammation.

SUMOylation prominently modifies components of the transcriptional apparatus (recently reviewed in ref. [44]). However, so far only one work has attempted to define the

impact of altered SUMOylation on gene expression *in vivo*. In a neuron-specific SUMO1/2/3 knockdown model [45], SUMO served as a transcriptional repressor under physiologic conditions but had decidedly gene-activating effects following transient cerebral ischemia. Unlike our study, pro-growth and anti-apoptotic pathways were prominently enhanced. Although direct comparisons between the very distinct approaches are of course problematic, the divergent findings may highlight that SUMOylation has substantially different roles in a largely differentiated, non-replicating tissue displaying very low SUMO conjugation under physiologic conditions (brain) vs. a rapidly regenerating tissue with relatively high baseline SUMO signal (intestine). Importantly, the gut is exposed to trillions of bacteria and milligrams of lipopolysaccharide within its lumen. Following intestinal injury, it is therefore imperative to prevent dysregulated inflammatory activation and to rapidly re-institute barrier integrity. In Ubc9Tg mice, this appears to be achieved at least in part by regulating influx of inflammatory cells as evidenced both by our pathological findings and by the prominent regulation of genes involved in inflammatory cell infiltration, such as that of adhesion proteins (p-selectin [46] and e-selectin [47]), of chemotactic factors (Cxc11 [48], 2 [49], 3, and 5 [50]), of inflammatory regulators (Il6 [51], Il15, and Il17A [52]), and proteases (e.g., AdamTs 1, 4, 14, Adam8, and Mmp8) [53]. Upstream analysis of the differentially expressed genes provided for a candidate network of eight major regulators of stress responses (Supplemental Fig. 6), most of which are known to be modulated by SUMOylation [31, 32, 54–58]. It is noteworthy that these findings show significant overlap with the work of Fritah et al. [39], which identifies SUMOylation as an important host-protective mechanism against gut-invasive pathogens through reduction of inflammatory epithelial destruction and which implicates, by PCR array analysis and stable isotope labeling with amino acids in cell culture (SILAC)-based quantitative proteomics, many of the same responses (Il6, Cxcl3, Cxcl1, Nfkb1a, and Xbp1).

Similar to our results in Ubc9Tg animals, Fritah et al. [39] observed that pro-inflammatory gene expression was not significantly altered under baseline conditions in mice, haploinsufficient for Ubc9. However, the lack of a signal in early postischemic timepoints also raised the question of whether responses of the villus-tip epithelium, i.e., the tissue with most dramatic changes of SUMOylation, were adequately represented in our whole-tissue microarray study. An important limitation of published genetic approaches [15, 39], including our own, is the inability to dissect cell type-specific responses to changes in SUMOylation. We aimed to address this by a focused analysis of apical epithelial transcriptional responses from samples obtained through laser microdissection and capture.

Ensuing cytokine and chemokine-targeted analysis supported observations from our microarray data that SUMOylation modulates Il17a expression, a key cytokine linking

T-cell activation to neutrophil mobilization and activation [59], and one of great importance in intestinal inflammation [60, 61]. The significance of these results is supported by the parallel decrease of the Il17a target gene Ccl20 [59, 61] and the induction of Il22, a member of the IL10 family known to inhibit production of Il17a [62]. Interestingly, in addition to Il17a, other factors downregulated in Ubc9Tg animals also shared T-cell chemotactic and regulatory functions (Cxc19 [63], Il27 [64], Ccl20 [65], Cxcl16 [66], and Ccl5 [67]) in parallel with a growing appreciation both of T-cell responses during early ischemia-reperfusion events [68] and of the role of SUMO in regulating these [69].

Our observation that global SUMO1 deletion results in better outcome following I/R injury was a completely unexpected finding, because it is generally believed that in organs under stress, increased SUMOylation improves outcome and decreased SUMOylation impairs it. For example, increased SUMOylation by overexpression of Ubc9 improves outcome following small intestine I/R (this study) or in experimental stroke [23], whereas silencing SUMO2/3 or SUMO1,2,3 expression worsens outcomes. [45, 70] As overexpression of SUMO1 is also reported to support protective responses in a heart failure model [71], we therefore expected outcomes following small intestine I/R injury to be worse in SUMO1-KO mice. SUMO1-deficient animals have been studied and apart from abnormalities in RanGAP1 localization and in formation of promyelocytic leukemia (PML) nuclear bodies [20], no phenotype has been attested in these animals [20, 21]. It was also noted that in SUMO1-KO mice most, if not all, SUMO-1 functions are compensated for by SUMO2/3, albeit with no evidence of increased SUMO2 or SUMO3 mRNA expression. [21] Our study confirmed this compensation of SUMO1 by SUMO2/3 overexpression and, most importantly, showed that such induction of SUMO2/3 blunts intestinal inflammatory responses after I/R. However, this protection from I/R injury is not as effective as in Ubc9Tg animals, which indicates independent effects of SUMO1 and/or of Ubc9 itself. Here we have to acknowledge that conclusions from our work may be limited by the difficulties we encountered in the quantification of SUMO isoform levels. On the whole-tissue level, high background signal in the connective tissue made it difficult to quantify changes that occurred in epithelial cells and methods to isolate epithelial cells were too complex and lengthy to produce faithful snapshot images of the extremely dynamic SUMOylation process. Therefore, although data from SUMO1-deficient mice underline the importance of

SUMO2/3 in the regulation of intestinal protective programs, our approach may have underappreciated changes of SUMO1 levels after I/R and/or in Ubc9 animals. Together, results reported here describe a unique pattern of nuclear SUMO2/3 staining in epithelial cells of the small intestine along the crypt-villus axis; they demonstrate that improved outcome following I/R in Ubc9-overexpressing mice is associated with preserved barrier function and blunted inflammatory responses, and they identify a previously undocumented process, whereby in SUMO1-KO mice, the compensatory overexpression of SUMO2/3 increases tissue I/R resistance. The tremendous protective power of SUMO conjugation reported here further highlights the importance of our ongoing work aimed at boosting the SUMOylation pathway to protect organs from damage induced by metabolic stress. [72–75]

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Compliance with Ethical Standards

Conflict of interest The authors declare that they have no conflict of interest.

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