

**Ductular reactions in the liver regeneration process with local inflammation after
physical partial hepatectomy**

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Abstract

Partial hepatectomy models in mice have been widely used for liver regeneration studies. A typical procedure removes approximately 2/3 of the liver by lobular ligation without tissue dissection. However, hepatectomy in humans involves physical damage (i.e., physical partial hepatectomy, PPHx). Therefore, the liver regeneration process after PPHx should involve reactions to acute local injury followed by systematic remodeling. To clarify the liver regeneration process after PPHx, we used a murine liver injury model that mimics the actual human surgical procedure. A 20–30% PPHx was performed by transection of the left lobe of the liver using an ultrasonically-activated scalpel in mice. Gene expression and morphological characteristics were analyzed during the liver regeneration process. Liver weight continuously increased by hypertrophic reaction of hepatocytes, while Ki67 staining showed hepatocyte proliferation. At the transected border, emergence of ductular reactions, a representative process of hepatic tissue remodeling that contain liver stem/progenitor cells, were observed. Gene expression of the transected border and non-damaged lobes revealed that inflammatory cytokine- and extracellular matrix-associated genes were significantly up-regulated at the transected border. Our PPHx model triggered local

extracellular matrix remodeling that resulted in ductular reactions. These processes occurred during the tissue repair process in local inflammatory responses as well as compensatory hepatocyte hypertrophy of the entire liver. These findings may provide insight for elucidating the mechanism of tissue repair and regeneration of the liver after PPHx.

Abbreviations

α -SMA	α -smooth muscle actin
CK19	Cytokeratin 19
DAB	3,3'-diaminobenzidine
DAPI	4',6-diamidino-2-phenylindole
DDC	3.5-diethoxycarbonyl-1.4-dihydrocollidine
DIC	Differential interference contrast
ECM	Extracellular matrix
EpCAM	Epithelial cell adhesion molecule
FFPE	Paraformaldehyde-fixed paraffin-embedded
HSC	Hepatic stellate cell
I κ B- α	IkappaB- α
LPC	Liver stem/progenitor cell
MMP	Matrix metalloproteinase
Muse	Multilineage-differentiating stress enduring

NF κ B Nuclear factor-kappaB

PPHx Physical partial hepatectomy

TNF- α Tumor necrosis factor- α

TUNEL Terminal deoxynucleotidyl transferase-mediated deoxyuridine triphosphate
nick-end labeling

Partial hepatectomy is performed on the premise that the remnant liver will regenerate by compensatory growth of the remaining segments.^{1,2} In rodents, a typical procedure removes approximately 2/3 of the liver by lobular ligation without tissue dissection (i.e., traditional partial hepatectomy).^{3,4} Human liver is not divided into multiple lobes, which makes segmental hepatectomy difficult. Therefore, the hepatectomy procedure is different in rodents and humans due to the anatomical difference. In general, hepatectomy in humans involves physical damage (i.e., physical partial hepatectomy, or PPHx), and therefore the liver regeneration process after PPHx should involve reactions as a result of the acute local injury followed by systematic remodeling.

The liver has a robust regenerative capacity with different modes of regeneration according to the type and extent of injury. In the situation of the traditional partial hepatectomy model, liver mass is restored by hypertrophic reaction and replication of

existing hepatocytes with or without minimal contribution of liver stem/progenitor cells (LPCs).^{2, 5} In comparison, when the proliferation of hepatocytes is impaired as a result of acute or chronic broad hepatic injury, LPCs emerge and expand forming duct-like structures, which are known as ductular reactions.^{6, 7} LPCs are thought to contain some population of cells that have the capacity to differentiate into both hepatocytes and biliary cells, which eventually contribute to tissue repair.^{8, 9} Ductular reactions are seen in a wide variety of human acute and chronic liver diseases.¹⁰ Although the mechanism of the ductular reaction is not fully understood, growth factors/cytokines released from the local microenvironment resulting in changes in the extracellular matrix (ECM) have been thought to be important components that elicit the ductular reaction.^{11, 12}

In this study, we used a murine liver injury model that mimics the actual human surgical procedure to clarify the liver regeneration process after PPHx. Using this model, we observed prompt ductular reactions localized to the damaged border during the wound healing process. Moreover, we found that the remodeling of ECM with an acute inflammation was a key molecular process associated with the emergence of ductular

reactions. These findings provide insight for elucidating the mechanisms of tissue repair and regeneration after PPHx.

MATERIALS AND METHODS

Animal Experiments

Eight to ten-week-old female C57BL/6J and SCID C.B-17/ICR-SCID/SCID mice (CLEA Japan Inc., Tokyo, Japan) were used for the experiments. All animal experiments in this study were approved by Iwate Medical University Ethical Committee for Animal Experiment Regulation (25-038). Mice were anesthetized by inhalation of isoflurane (2.5% v/v). After a 2 cm laparotomy, PPHx was performed by transection of the left lobe of the liver using a harmonic scalpel (Harmonic Focus[®], Ethicon Endo-Surgery, Blue Ash, OH, USA), which is an ultrasonically-activated surgical device for tissue dissection without lobular ligation, to remove approximately 20-30% of the entire liver. The average length of the liver transection line was approximately 10 mm. Regenerating liver samples were harvested at 24, 48, and 72 hours after the PPHx and then formalin fixed or embedded in O.C.T compound (Sakura

Finetek, Tokyo, Japan) for further analysis. Mice that underwent only laparotomy (sham operation) were used as a control. Sham-operated livers were harvested 24 hours after laparotomy for further analysis.

Histological Evaluation

Paraformaldehyde-fixed paraffin-embedded (FFPE) (4%) liver tissue blocks were cut into 3 μm sections and stained with hematoxylin and eosin.

Apoptotic Cell Detection

Apoptosis was evaluated by terminal deoxynucleotidyl transferase-mediated deoxyuridine triphosphate nick-end labeling (TUNEL) assay using the *In Situ* Cell Death Detection Kit, POD (Roche diagnostics, Mannheim, Germany). TUNEL staining was performed on the transected and remnant medial lobes at each time point after the PPHx.

Immunohistochemistry

The following primary antibodies were used for colorimetric immunohistochemical staining on paraffin embedded liver sections: rabbit anti-Ki67 antibody (1:300; RM-9106, Thermo Scientific, Fremont, CA, USA); mouse anti- α -smooth muscle actin (α -SMA) antibody (1:600; MS-113-P, Thermo Scientific); rabbit anti-matrix metalloproteinase (MMP)-9 antibody (1:75; ab124513, abcam, Cambridge, UK); and rabbit anti-cytokeratin 19 (CK19) antibody (1:50; ab15463, abcam). After primary antibody incubation, the FFPE sections were incubated with peroxidase-labeled anti-mouse or anti-rabbit secondary antibodies (Histofine Simple Stain Max-PO Kit, Nichirei, Tokyo, Japan). Then, 3,3'-diaminobenzine (DAB) was used for colorimetric signal development in the presence of peroxidase, which was conjugated with the secondary antibody (K3468, DAKO, Carpinteria, CA, USA). The sections were then counterstained with hematoxylin to visualize nuclei.

For immunofluorescent detection, the following primary antibodies were used on frozen sections (6 μ M): rabbit anti-CK19 antibody (1:40); rat anti-Thy1 antibody (1:20; 553011, BD Biosciences, San Diego, CA, USA); rat anti-epithelial cell adhesion molecule (EpCAM) antibody (1:150; 552370, BD Biosciences); rat anti-A6 antibody

(1:100; gift from Valentina Factor, NIH); and rabbit anti-Ki67 antibody (1:300). All primary antibodies, except the anti-Ki67 antibody, were detected with one of the following secondary antibodies: anti-rabbit Alexa Fluor 488 (1:300; A21206, Life Technologies, Carlsbad, CA, USA) or anti-rat Alexa Fluor 594 (1:300; A11007, Life Technologies). The anti-Ki67 primary antibodies were incubated with peroxidase-labeled anti-rabbit secondary antibody (Histofine Simple Stain Max-PO Kit, Nichirei). Bound secondary antibody was detected by treating with tyramide-Cy3 (NEL704A001KT, Perkin Elmer Life Sciences, Waltham, MA, USA). Nuclei were counterstained with 4'-6-diamidino-2-phenylindole (DAPI). The images were obtained by a laser confocal microscopy (Nikon Corp., Tokyo, Japan). To quantify the size of hepatocytes, Alexa Fluor 488 phalloidin (1:300; A12379, Life Technologies) was used to recognize the outline of hepatocytes. The average size of hepatocytes was quantified using ImageJ software (NIH Image; <http://rsb.info.nih.gov/ij/>).

cDNA Microarray Analysis

Total RNA was extracted using an RNeasy Mini kit (Qiagen, Tokyo, Japan) according to the manufacturer's protocol. RNA samples were extracted from 3 mm³ specimens of liver tissue from the transected border of SCID mice at each time point. The concentration and purity of the total RNA were determined using a SmartSpec Plus Spectrophotometer (Bio-Rad, Hercules, CA, USA). Gene expression profiling was performed according to the manufacturer's instructions (Agilent Whole Mouse Genome Microarray, 4×44K, Agilent Technologies, Santa Clara, CA, USA). The microarray data were analyzed by Genespring GX12 software (Agilent Technologies). Genes with > 3.0-fold up- or down-regulation compared with sham samples were considered differentially expressed. Differentially expressed genes at any single time point after the PPHx were analyzed by one-way hierarchical clustering with a Euclidean distance measurement and a Ward linkage in the Genespring GX12 software. Genes from major clusters were further analysed using the Database for Annotation, Visualization and Integrated Discovery (DAVID; <http://david.abcc.ncifcrf.gov>) for gene ontology classification.

RNA Preparation and Quantitative Reverse Transcription PCR

For this assay, C57BL/6J mice were used. Total RNA was extracted from 3 mm³ specimens of medial intact lobe and of the transected border of the damaged left lobe, respectively. First-strand cDNA was synthesized from one µg of total RNA using the ReverTra Ace qPCR RT Kit (Toyobo, Osaka, Japan). The synthesized cDNA was used for real-time PCR (LightCycler: Roche diagnostics, Mannheim, Germany) with FastStart SYBR Green Master (Roche diagnostics) as a fluorophore. RNA expression was normalized to the housekeeping gene 18s rRNA. Primer sequences are listed in Supplementary Table 1.

Proliferation of Mouse LPCs

An EpCAM positive liver progenitor cell line from the liver of a 3.5-diethoxycarbonyl-1.4-dihydrocollidine (DDC)-fed adult mouse was kindly provided by Professor Atsushi Miyajima and Dr. Minoru Tanaka of University of Tokyo.¹³ Serum-starved LPCs were treated with recombinant murine tumor necrosis factor- α (TNF- α) (rm TNF- α : Wako, Osaka, Japan) with or without BAY11-7082 (Wako),

which irreversibly inhibits I κ B- α (I κ B- α) phosphorylation resulting in the disruption of nuclear factor-kappaB (NF κ B) nuclear translocation and activation.¹⁴ Cells were stimulated with the reagents for 24 h to assess Cyclin D1 expression and for 15 min to assess p-I κ B- α expression.

Cell Viability Assay

Serum-starved LPCs (5.0×10^4 cells/mL) were treated with TNF- α , BAY11-7082, or their combination for 24 h. The number of viable cells was counted using a Cell Count Reagent SF (Nacalai Tesque, Tokyo, Japan).¹⁵ Light absorbance was measured at 450 nm with a microplate photometer (Immuno Mini NJ-2300: InterMed, Tokyo, Japan).

Western Blot

Total protein was extracted from the LPCs using a Total Protein Extraction Kit (BioChain Institute, Inc., Hayward, CA, USA). Five μ g of protein from each sample was separated using 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis and electrotransferred onto a polyvinylidene difluoride membrane. Immunoblotting was

performed using rabbit anti-p-I κ B- α (Ser32) (1:1000; #2859, Cell Signaling Technology, Danvers, MA, USA); mouse anti-Cyclin D1 (1:1000; #2926, Cell Signaling Technology); and mouse anti- β -actin (1:500; sc-47778, Santa Cruz Biotechnology, Dallas, TX, USA). Horseradish peroxidase-conjugated sheep anti-mouse IgG (1:5000; NA9310V, GE healthcare, Little Chalfont, Buckinghamshire, UK) or donkey anti-rabbit IgG (1:5000; N9340V, GE Healthcare) were used as secondary antibodies. Immunoreactive bands were detected by chemiluminescence using the ECL Prime Western Blotting Reagent (GE Healthcare).

Statistical Analysis

Unless otherwise noted, data were expressed as the mean \pm s.e.m. Statistical analyses including Student's unpaired, two-tailed *t*-test were conducted using GraphPad Prism, version 6 (GraphPad Software, San Diego, CA, USA). Differences were considered significant when the *P*-value was < 0.05 .

RESULTS

Mouse Model of PPHx

In contrast to a traditional partial hepatectomy that simply removes the medial and left lateral lobes by ligating the pedicle, our PPHx model employs a harmonic scalpel that remove approximately 20-30% of the entire liver (Figure 1a). The width of the ablation band of the liver resection was approximately 1 mm, which corresponded to the coagulated tissue (Figure 1b). Figure 1c shows histological characterization of the wound edge tissue at specific time points after the PPHx. At 24 h, massive hepatic parenchymal injury was observed in the damaged area. TUNEL assays exhibited a broad area of cells with DNA fragmentation in the damaged area in contrast to non-damaged areas (Figure 1c and Supplementary Figure 1). At 48 h, substantial infiltration of neutrophils and lymphocytes was observed, while small duct-like structures were seen along the damaged border. By 72 h, infiltrating cells further accumulated along the damaged border. The cells comprising the duct-like structures were CK19 positive, suggesting that ductular reactions had been initiated. We also confirmed that the negative control showed no staining (Supplementary Figure 2).

PPHx Induces Ductular Reactions at the Damaged Border

To investigate the process of ductular reactions that emerged at the transected border, we next performed CK19 and Thy1 immunofluorescent staining. There were almost no CK19-positive cells at the transected border at 24 h (Figure 2a). At 48 h, CK19-positive cells (i.e., ductular reactions or LPCs) were observed in some areas of the transected border (Figure 2b). CK19-positive duct-like structures were also observed, especially in the periportal areas adjacent to the damaged area (Figure 2c). By 72 h, the number of CK19-positive cells had increased along the damaged border (Figure 2d). In addition, Thy1-positive cells, known as a niche for LPCs,¹⁶ also appeared in the vicinity of the ductular reactions (Figure 2e). The CK19-positive area at the damaged border was significantly increased over the time course (Figure 2f). Thy1-positive areas also appeared to be increased by 48 h, but the difference was not statistically significant. In intact tissue areas, only preexisting structured bile ducts stained positive for CK19 and apparent ductular reactions were not observed. We further examined the expression of A6 and EpCAM to characterize newly formed bile duct-like structures (i.e., ductular reactions) induced by PPHx. Ductular reactions showed A6 and EpCAM-positive cells

adjacent to the transected border (Supplementary Figures 3a and b). These findings suggest that physical liver damage by PPHx induces an LPC response that may form bile duct-like structures.

PPHx Does Not Affect Hypertrophic Reaction and Proliferation of Hepatocytes in the Remnant Liver

After traditional partial hepatectomy in mice, hypertrophy occurs throughout the entire liver followed by proliferation of hepatocytes.² To investigate whether hypertrophy and proliferation of hepatocytes are involved in the wound healing process, we monitored microscopic and macroscopic features after PPHx in the damaged front and intact remnant liver.

The liver weight increased in a time-dependent manner (Figure 3a). The transected site was identified grossly with no indication of adjacent tissue damage (Supplementary Figure 4). Microscopically, we examined the size of hepatocytes, which play a critical role in restoring liver volume after partial hepatectomy. The size of hepatocytes increased up to 24 h and slightly decreased from 48 to 72 h (Figure 3b). We

also examined the local proliferative activity of hepatocytes with Ki67 staining, but there was no significant difference between the transected left and remnant medial lobes at any time points (Figure 3c). This result was in agreement with a previous report that 50% of PPHx induced hepatocyte proliferation in the entire liver.¹⁷ Cyclin D1-encoding gene *Ccnd1*, which is an alternative cellular proliferation marker,¹⁸ was also significantly up-regulated at 72 h, but there was no significant difference between transected and intact lobes (Figure 3d). In contrast, Ki67-positive proliferating cells were significantly increased in the ductules at the transected border, suggesting that duct-like structures located at the transected border after PPHx have increased proliferative activity (Supplementary Figures 5a and b).

Gene Expression Profile in the Damaged Area Reveals an Inflammatory Response and Matrix Metalloproteinase Activities

We hypothesized that the wound region provides a microenvironment that mobilizes LPCs in response to injury. To determine the molecular profiles of the wound region, we performed microarray analysis for genes specifically expressed in the transected

border. A total of 3057 differentially expressed genes were subjected to one-way hierarchical clustering and represented in a heatmap of the time course (Supplementary Figure 6). Differentially-expressed genes were clustered by the time course and each cluster was related to functional annotation using DAVID. We selected up to the top four genes from ontology terms in each cluster based on the literature (Table 1).¹⁹⁻²¹ We noted that genes encoding pro-inflammatory cytokines (e.g., Tnf, Il6, and Il1b), CC and CXC chemokines and their receptors (e.g., Cxcl2, Cxcr2, Ccl2, and Ccr2) were up-regulated transiently at 24 h, and 32% of them returned to control levels at 48 to 72 h after the PPHx. Genes in ECM-associated genes, such as collagen transcripts (e.g., Col1a1, Col1a2, Col3a1, Col4a1, and Col4a2), S100a4 [which promotes liver fibrosis by activating hepatic stellate cells (HSCs)],²² MMP genes (e.g., Mmp2, 3, 7, 9, and 13), and tissue inhibitors of metalloproteinase genes (Timp1 and Timp2), were up-regulated at 24 h and remained at their highest levels for up to 72 h. From 48 h to 72 h, cell cycle-associated genes were up-regulated, which is consistent with previous reports indicating that genes in both cell cycle and DNA replication categories are up-regulated after a traditional partial hepatectomy.^{23, 24} In fact, Ki67 staining showed that

hepatocytes entered the cell cycle at 48 h. However, no difference was seen in Ki67 positivity of hepatocytes between transected and intact lobes (Figure 3c). In addition, no difference was seen in the level of *Ccnd1* mRNA expression between transected and intact lobes (Figure 3d). In contrast, inflammatory cytokine- and ECM-associated genes do not represent major categories for the traditional partial hepatectomy.²⁵ We confirmed that a subset of these genes were up-regulated at the transected border by quantitative reverse transcription PCR (Figures 4a and b). Taken together, these data suggest that remodeling of the ECM with an acute inflammation is a key molecular process of wound healing after PPHx. In addition, we confirmed that the expression levels of *Spp1* and *Afp* mRNA, which are a bile duct-associated gene and progenitor marker, respectively, were up-regulated at the transected border.

Among the genes of additional interest, *Krt19* mRNA, which encodes the CK19 protein, was significantly up-regulated at the transected border (Figure 4c). These data support the results of the immunofluorescence analysis indicating that ductular reactions only occurred at the transected border.

α -SMA-Positive Hepatic Stellate Cells Were Observed at the Transected Border

HSCs, which express MMPs,²⁶ are found in tight association with LPCs.¹¹ Therefore, we stained liver sections with an α -SMA antibody, which is a marker of HSC activation, to determine the location of HSCs in the liver. Immunohistochemical staining demonstrated that the number of α -SMA-positive cells was significantly increased at the transected border compared with an intact lobe from 48 h post-procedure (Figures 5a and b). In intact tissue areas, smooth muscle cells of hepatic artery and portal vein branches were mainly α -SMA-positive (Supplementary Figure 7). Although infiltrating Kupffer cells also influence the induction of LPCs, there was no significant change in the number of F4/80 positive Kupffer cells in the transected border or intact lobe (Supplementary Figures 8a and b). MMP-9 produced by infiltrating Kupffer cells is critical for LPC induction in the chronic liver injury mouse model.²⁷ Based on the lack of F4/80 positive cell infiltration, we examined localization of MMP-9 producing cells at the transected border. Immunohistochemical staining revealed that infiltrating cells expressed MMP-9 mainly at the transected border (Figure 5c), which is consistent with the area in which α -SMA-positive cells were located (Figure 5a), suggesting that a

substantial number of α -SMA-positive cells at the transected border may contribute to ECM remodeling.

TNF- α Triggers Proliferation of Mouse LPCs via IKK/NF κ B Signaling

Our experiments suggest that infiltrating cells at the transected border should contribute to ECM remodeling that induces LPCs to participate in ductular reactions.

Inflammatory responses associated with ECM remodeling are considered to be important components in the induction of LPCs.¹² To determine whether LPC proliferation is triggered by inflammatory cytokines, we assessed if the administration of TNF- α and IL-6 induces LPC proliferation in vitro. Administration of 50 ng/ml of TNF- α induced LPC proliferation by stimulating the activation of I κ B (Figures 6a and b). In contrast, however, administration of IL-6 did not induce LPC proliferation (data not shown). In a separate study, we found that CXCL2 induced mouse LPC proliferation in vitro, suggesting the involvement of an inflammatory response in LPC-associated liver regeneration (Kakisaka et al. unpublished data).

We next examined whether the proliferative effect of TNF- α was abolished by the NF κ B inhibitor BAY11-7082. It has been reported that BAY11-7082 suppresses hepatocyte proliferation at 10 μ M.²⁸ Incubation of LPCs with 5 μ M of BAY11-7082 for 24 h induced the suppression of proliferation, whereas lower concentrations (range 0.1~1 μ M) exhibited no suppressive effect (Figure 6a). Administration of 50 ng/ml of TNF- α stimulated the phosphorylation of I κ B- α (Figure 6b) and increased Cyclin D1 expression (Figure 6c). Moreover, the induction of these genes at the protein level was suppressed by co-treatment with as low as 0.1 μ M of BAY11-7082 (Figures 6b and c).

DISCUSSION

In this report, we evaluated a liver wound healing process after the simultaneous induction of liver volume loss and tissue damage. Using a murine liver injury model that mimics the actual human surgical procedure, we demonstrated that both hypertrophy and proliferation of hepatocytes were induced simultaneously in the entire liver by local severe tissue damage as a part of liver regeneration. At the transected border, ductular reactions were observed at 48-72 h after the PPHx along the damaged

border. Ductular reactions, which contain a population of LPCs, are thought to occur diffusely when the proliferation of hepatocytes is impaired by both acute and chronic liver damage.^{6, 10} Our results suggest that ductular reactions could be induced even if the damage is limited to a small area of the liver. Although the differentiation of LPCs into hepatocytes and cholangiocytes has been controversial,^{9, 29, 30} based on our results, it is conceivable that LPCs participate in the tissue repair process after PPHx.

We have recently shown that a unique stem cell population in adult human bone marrow mesenchymal stem cells, namely multilineage-differentiating stress enduring (Muse) cells, integrate near the transected border and subsequently differentiate into major liver components after PPHx.³¹ In that study, human muse cells were injected into the tail vein of SCID mice 24 h after PPHx, and we found that they engrafted preferentially into the damaged border and expressed liver progenitor markers 48 h after transplantation. From these previous observations, we hypothesize that the wound region provides a microenvironment that mobilizes LPCs.

Based on our microarray results, we found that the PPHx induced the expression of inflammatory cytokine genes, including Tnf, Il6, and Cxcl2, at the wound site

compared with intact lobes. The involvement of TNF- α , IL-6, and other inflammatory cytokines have been implicated in LPC proliferation when injury-induced proliferation of hepatocytes is impaired.³²⁻³⁴ Our in vitro experiment showed that TNF- α induced LPC proliferation. Indeed, the NF κ B inhibitor BAY11-7082 inhibited Cyclin D1 induction at a concentration that did not cause cytotoxicity. These results suggest that TNF- α signaling through NF κ B activation plays a role in the proliferation of LPCs.³³ A recent report demonstrated that TNF- α was responsible for inducing the morphological change of hepatocytes into biliary epithelial cells in vitro.²¹ Another recent study also demonstrated that mature hepatocytes could convert into LPCs to form bile duct-like structures in chronic injury models.³⁵ These observations suggest that LPCs induced by pro-inflammatory cytokines are important factors in the generation of ductular reactions.

We also observed that a significant up-regulation of ECM-associated genes, such as Mmp2, Mmp9, and S100a4, occurred simultaneously with the induction of ductular reactions. MMP-2 and MMP-9, which are involved in the degradation of the ECM, are closely associated with LPC mobilization.³⁶ In addition, S100a4 promotes

liver fibrosis by activating HSCs.²² Immunohistochemical staining of MMP-9 revealed that infiltrating cells expressed MMP-9 mainly at the transected border, which was consistent with the area in which both α -SMA-positive cells and ductular reactions were located. Our PPHx model induced severe hepatic parenchymal damage at the transected site. Pathological features of the transected site resemble acetaminophen-induced liver injury,^{37,38} whereby A6/CK-positive cells were observed as early as 3 h.³⁷ In the PPHx model, no CK19-positive cells were seen at the transected border up to 24 h. The CK19-positive cells were detected between 48 and 72 h after PPHx with infiltrating inflammatory cells at the transected border. Moreover, those CK19-positive ductular cells also expressed A6. These results are consistent with recent studies showing that ductular reactions appear to be tightly associated with HSCs, macrophages, and the ECM during the response to liver injury.^{11,39} Therefore, although our PPHx is a distinct model of liver injury using this procedure, the pathological features of severe injury by acetaminophen are not entirely dissimilar to those of PPHx. Taken together, we suggest that expansion of ductular reactions in the wound region is regulated by cell-cell and

cell-matrix interactions, which are triggered, at least in part, by TNF α /NF κ B

inflammatory pathway activation.

In conclusion, our PPHx model triggered local ECM remodeling that resulted in ductular reactions. These processes occurred during the tissue repair process in local inflammatory responses as well as compensatory hepatocyte hypertrophy of the entire liver. These findings may provide insight into elucidating the mechanism of hepatic tissue repair and regeneration of the liver after PPHx.

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DISCLOSURE/CONFLICT OF INTEREST

The authors declare no conflict of interest.

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TITLES AND LEGENDS TO FIGURES**Figure 1** Schematic of the PPHx and liver section histology after the PPHx. **(a)**

Conventional ligation point of traditional 2/3 partial hepatectomy (asterisk). The dashed

line indicates the transected line in the left lobe in this study. **(b)** Macroscopic view and

hematoxylin and eosin staining of the transected liver at 24 h after the PPHx. The right

side of the hematoxylin and eosin staining is the coagulated area. Scale bar, 500 μm . **(c)**

Histological examination of transected liver at the indicated time points. Scale bar

represents 200 μm except for the right lower panel. In the right lower panel, the arrows

indicate that CK19-positive duct-like structures. Scale bar, 50 μm .

Figure 2 CK19 and Thy1 immunofluorescent staining after a PPHx. **(a)**

CK19-immunostaining, DIC, and merged images at 24 h after the PPHx. Arrowheads

indicate the transected line. Scale bar, 300 μm . **(b)** CK19-immunostaining, DIC, and

merged images at 48 h after the PPHx. Scale bar, 300 μm . **(c)** CK19-positive duct-like

structures in periportal areas adjacent to the damaged area. Dashed line indicates the

transected border. Scale bar, 100 μm . **(d)** CK19-immunostaining, DIC, and merged

images at 72 h after PPHx. Scale bar, 300 μm . **(e)** Double immunostaining of CK19 (green) and Thy1 (red). Scale bar, 100 μm . **(f)** CK19- and Thy1-positive areas were quantified by immunostaining images. Student's *t*-test was performed between each time point and 24 h after the PPHx. Mean \pm s.e.m. (n=3), ** $P < 0.01$, *** $P < 0.001$, n.s., not significant. DIC, differential interference contrast. PV, portal vein.

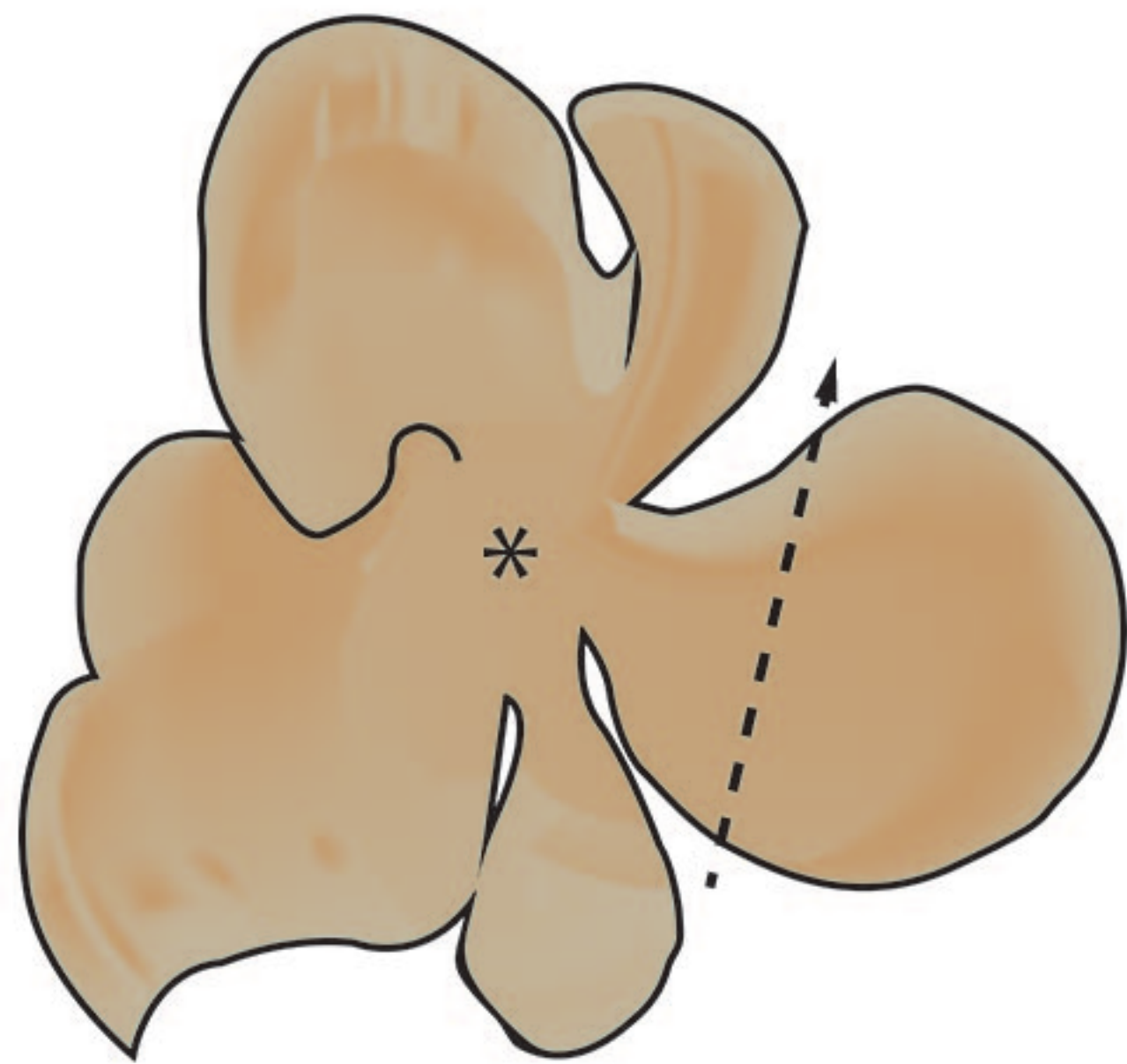
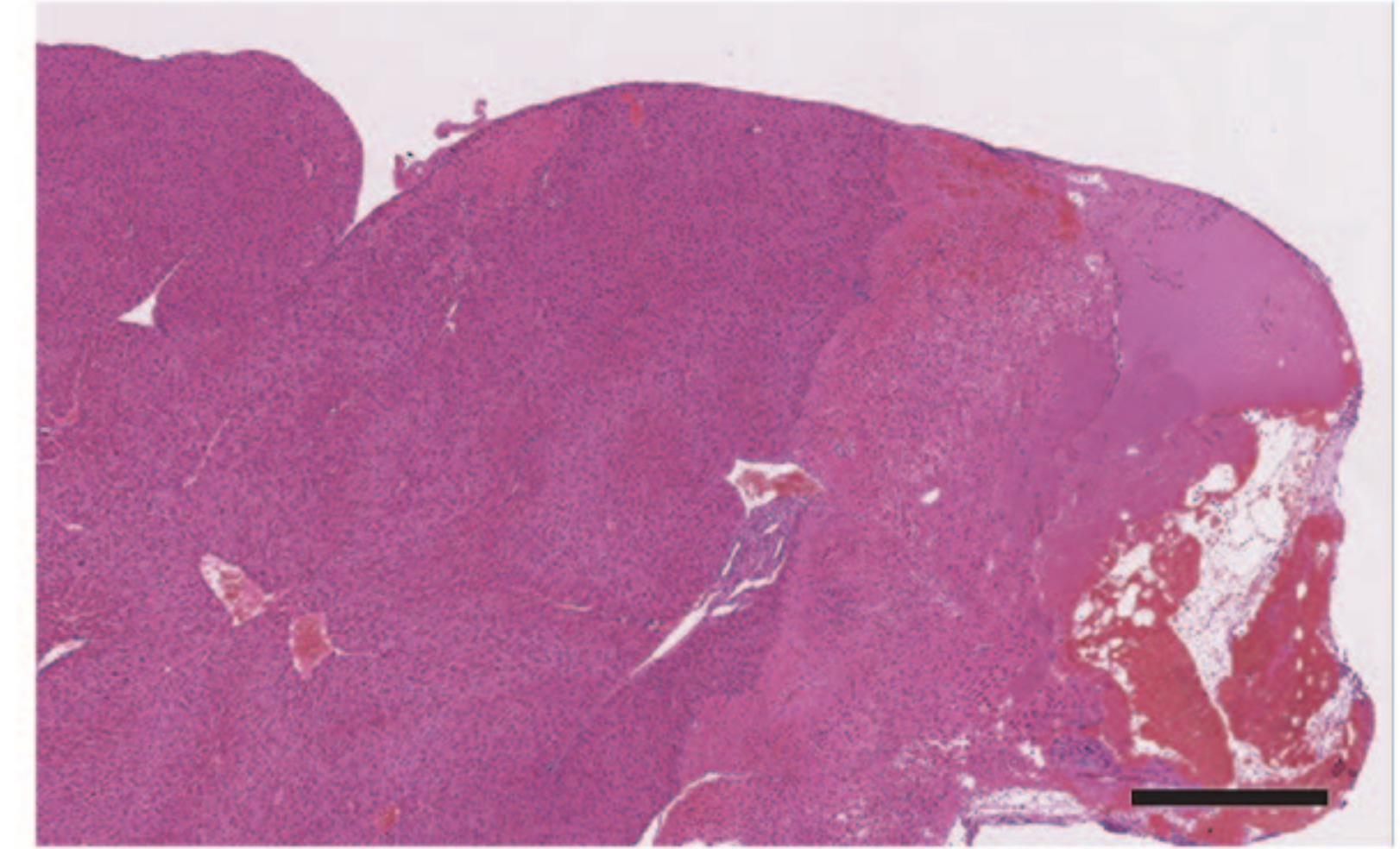
Figure 3 Hypertrophic reaction and proliferation of hepatocytes in the remnant liver after the PPHx. **(a)** Changes of liver to body weight ratios after the PPHx. Student's *t*-test was performed between each time point and 24 h after PPHx. Mean \pm s.e.m. (n=3), * $P < 0.05$, ** $P < 0.01$. **(b)** Size of hepatocytes after PPHx. Student's *t*-test was performed between sham-operated liver and each time point. Mean \pm s.e.m. (n=3), ** $P < 0.01$, *** $P < 0.001$. **(c)** Ki67 staining on liver sections of transected left lobe and intact medial lobe. Scar bars, 100 μm . Mean \pm s.e.m. (n=3), *** $P < 0.001$. **(d)** Ccnd1 mRNA expression levels in transected lobe and intact lobe. Fold induction was estimated on sham-operated liver. Mean \pm s.e.m. (n=3), * $P < 0.05$.

Figure 4 Gene expression analysis of samples taken from the transected border and intact lobe. **(a)** Expression of *Tnf*, *Il6*, and *Cxcl2* mRNA. **(b)** Expression of *Mmp2*, *Mmp9*, and *S100a4* mRNA. **(c)** Expression of *Spp1*, *Afp*, and *Krt19* mRNA. Fold induction was estimated for the values of intact lobe at each time point after normalization to 18s mRNA levels. Student's *t*-test was performed. Results are represented as mean \pm s.e.m. (n=3), * P <0.05, ** P <0.01, *** P <0.001.

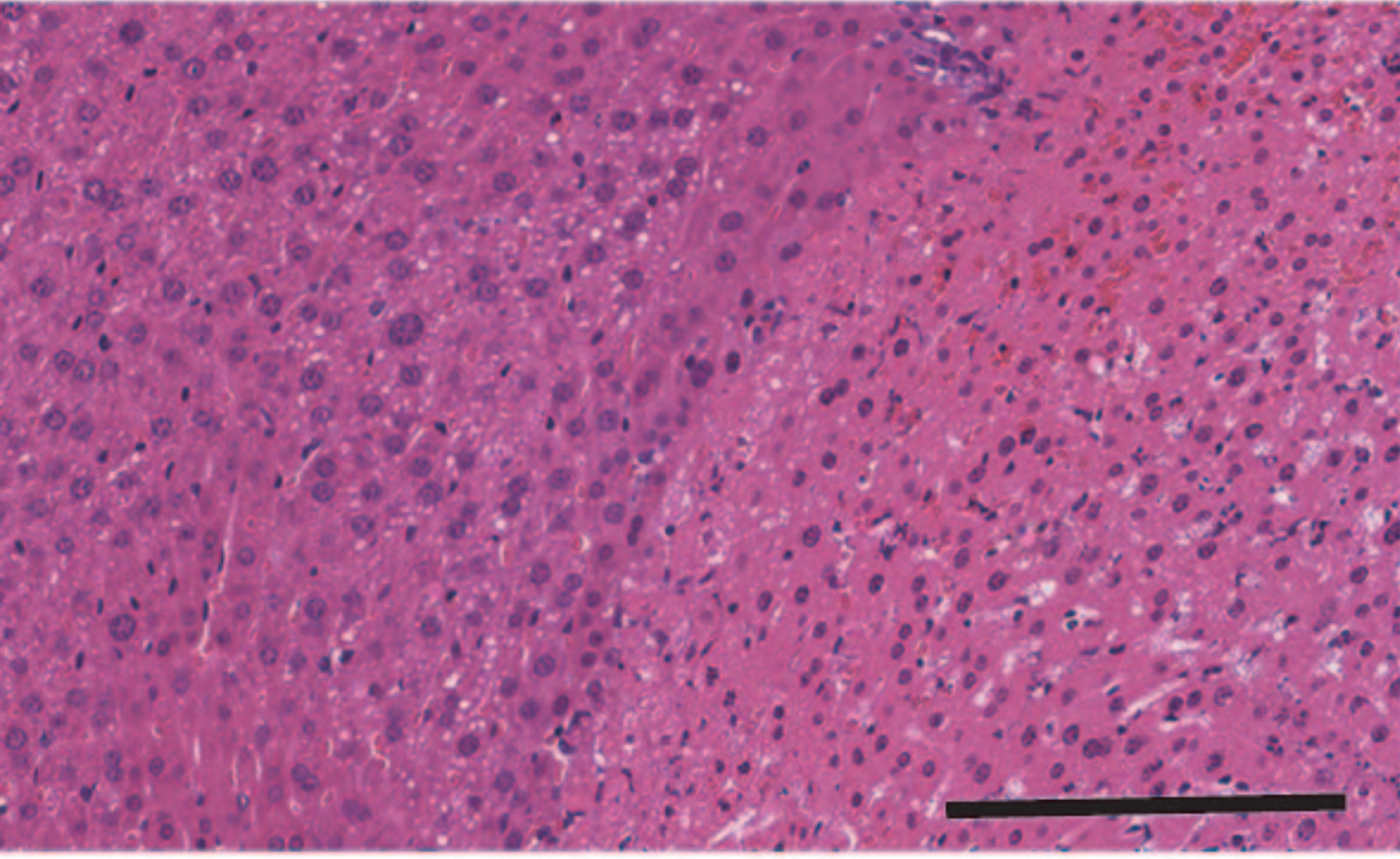
Figure 5 Immunohistochemical analysis of α -SMA and MMP-9. **(a)** Histological examination of α -SMA-positive area in the transected area at the indicated time points. **(b)** Quantitative analysis of α -SMA-positive area using ImageJ program. Ten randomly selected high-power fields were quantified. Scale bar, 200 μ m. Student's *t*-test was performed. Mean \pm s.e.m., * P <0.05, ** P <0.01, *** P <0.001, n.s., not significant. **(c)** Histological examination of MMP-9 in the transected area at the indicated time points. Scale bar, 200 μ m.

Figure 6 Effects of TNF- α and I κ B- α inhibitor BAY11-7082 on mouse LPC

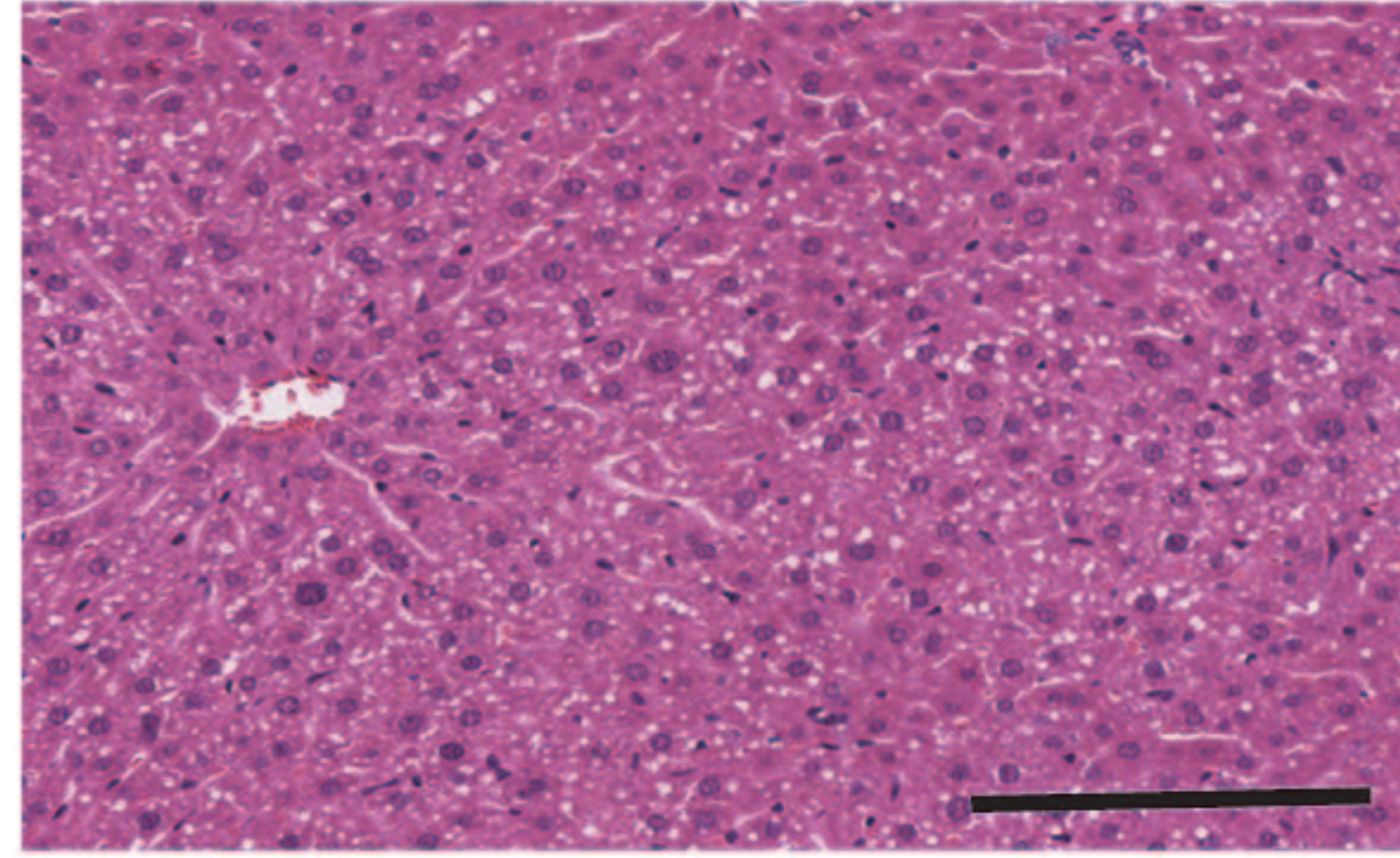
proliferation. **(a)** Cell viability of LPCs was estimated in response to treatment with 50 ng/ml TNF- α , BAY11-7082, and their combination. Student's *t*-test was performed to compare each treated sample to control sample. Results are represented as mean \pm s.e.m. (n=3), **P*<0.05, ***P*<0.01, ****P*<0.001. **(b)** and **(c)** Western blot analysis was performed using cell lysates of LPCs and antibodies specific for p-I κ B- α and Cyclin D1 as indicated. LPCs were treated with 10 ng/ml or 50 ng/ml TNF- α with or without 0.1 μ M, 1 μ M, or 5 μ M BAY11-7082. β -actin was used as the internal control.

a**b****c**

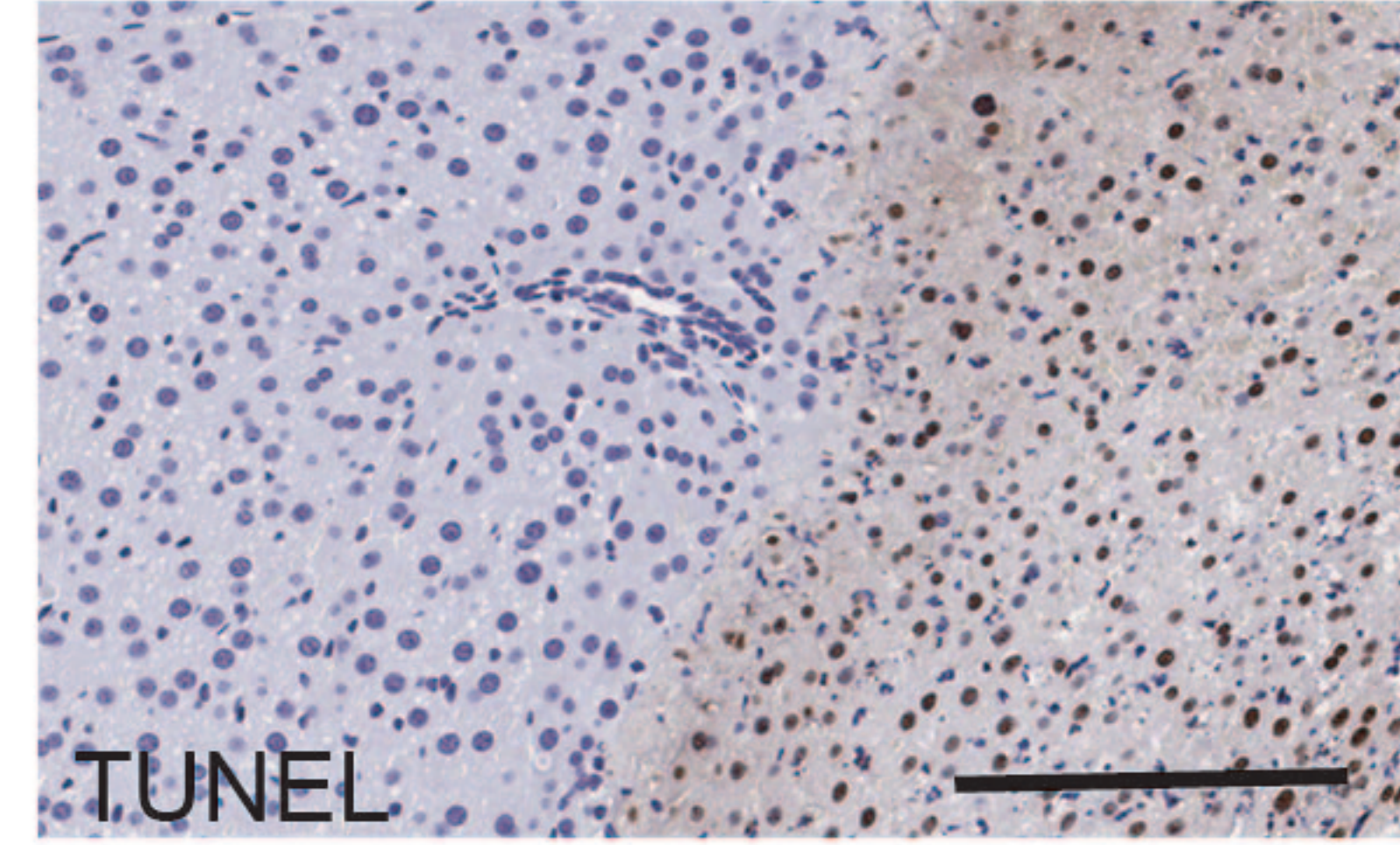
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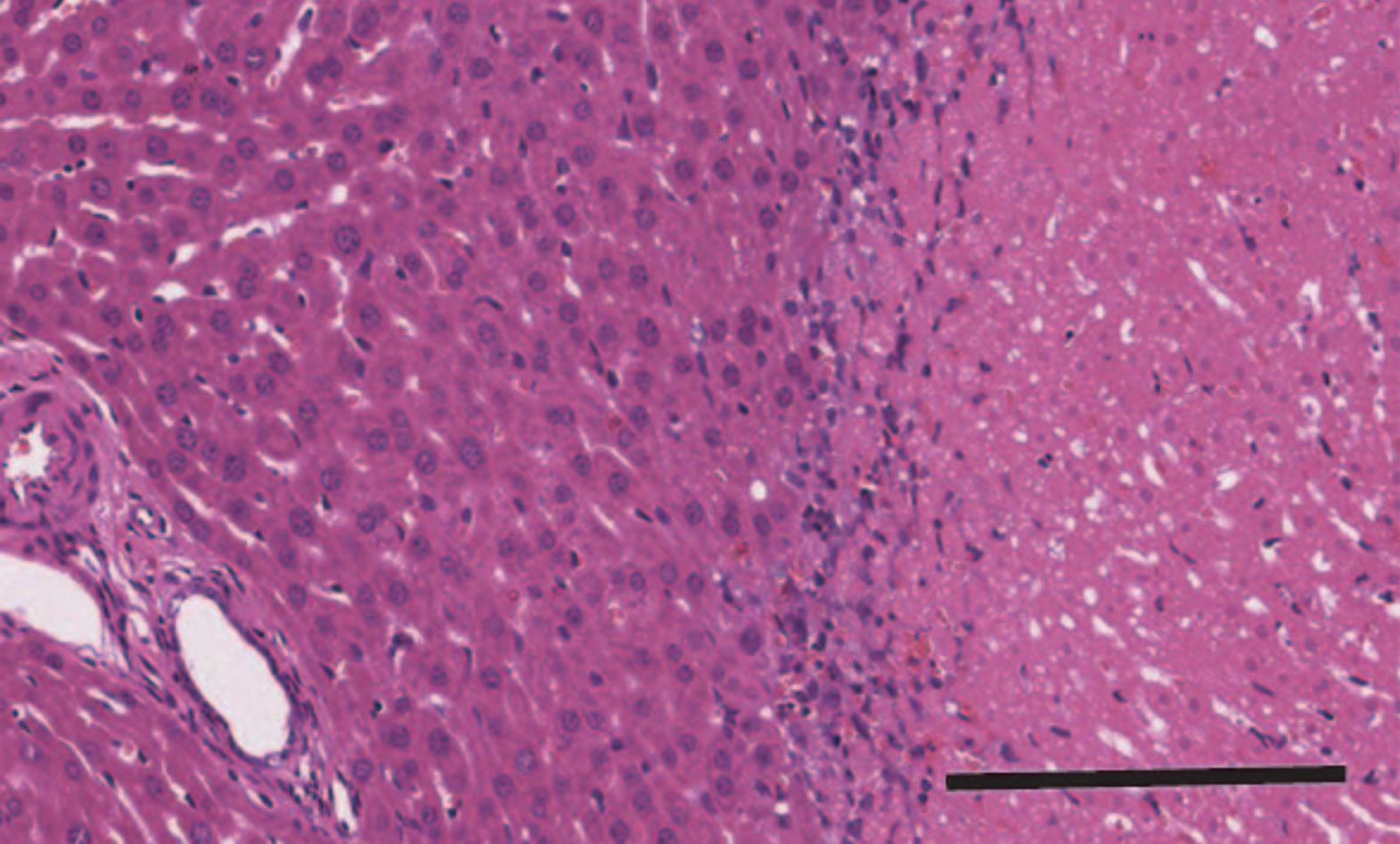
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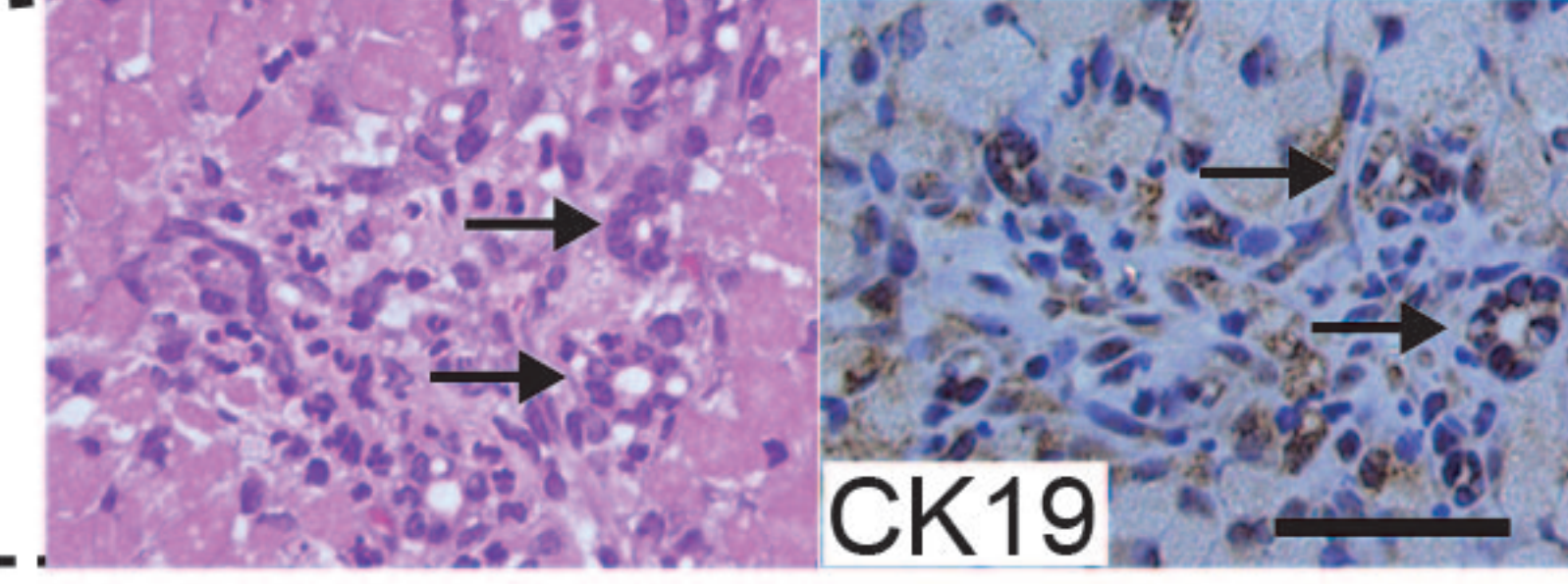
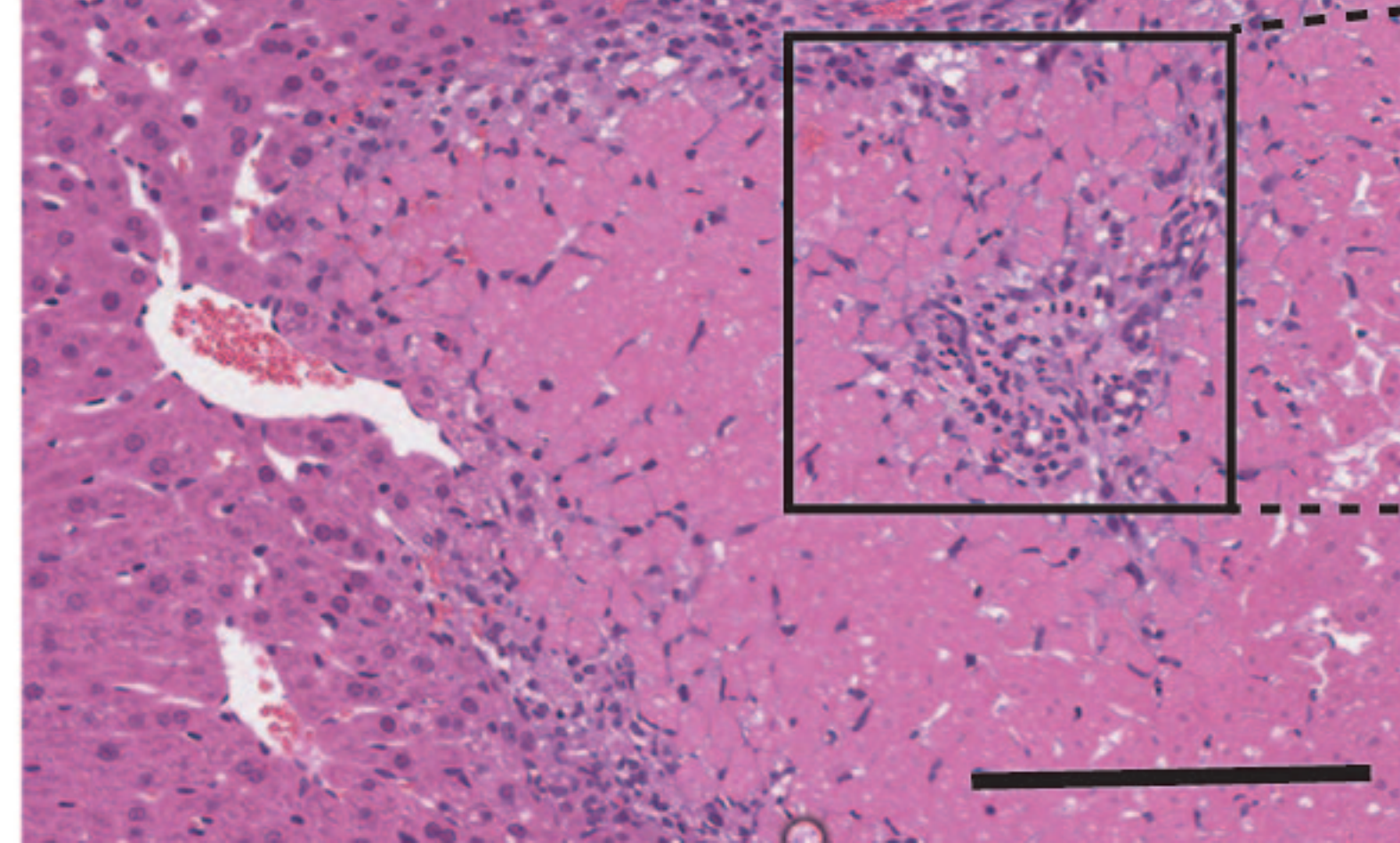
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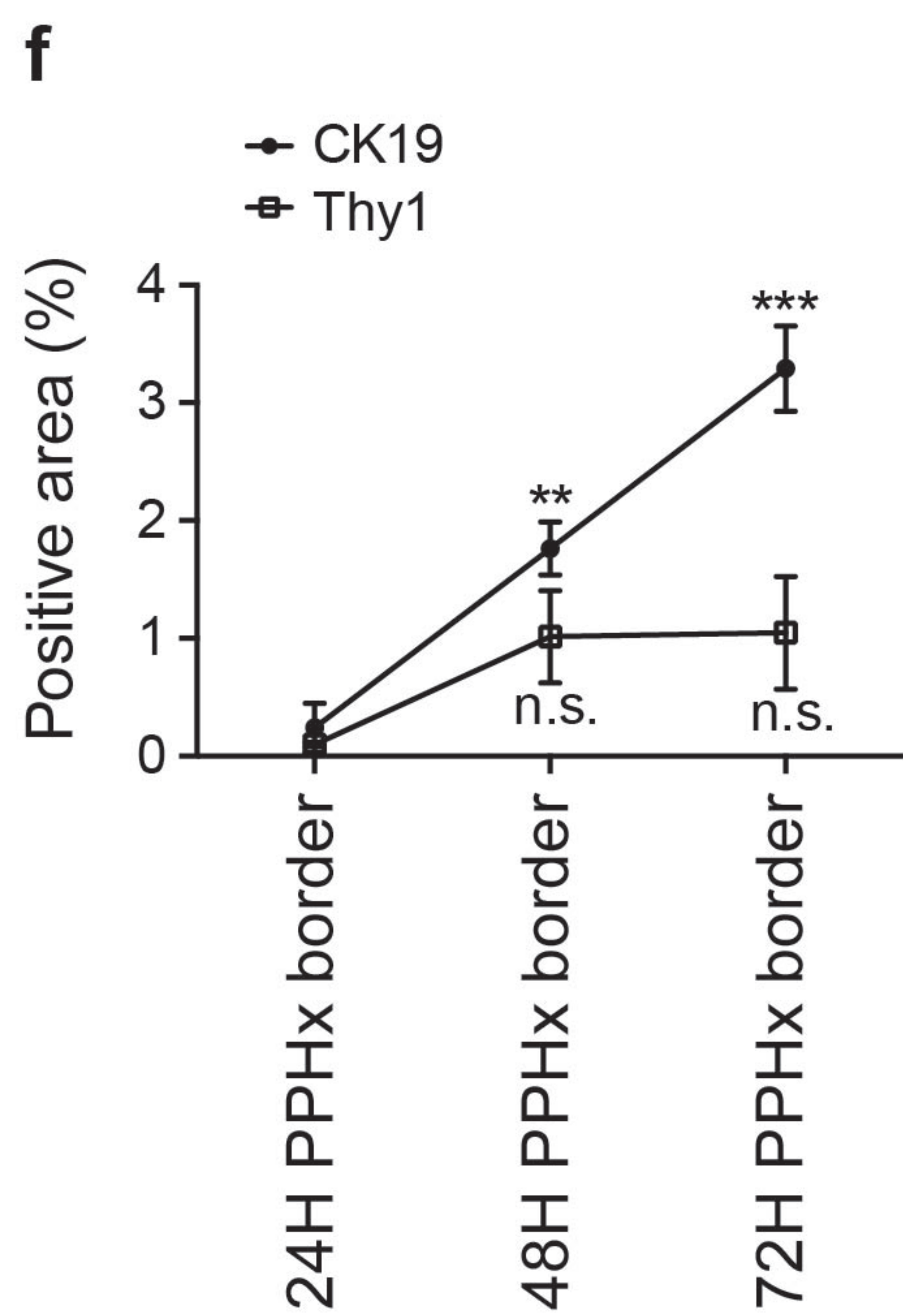
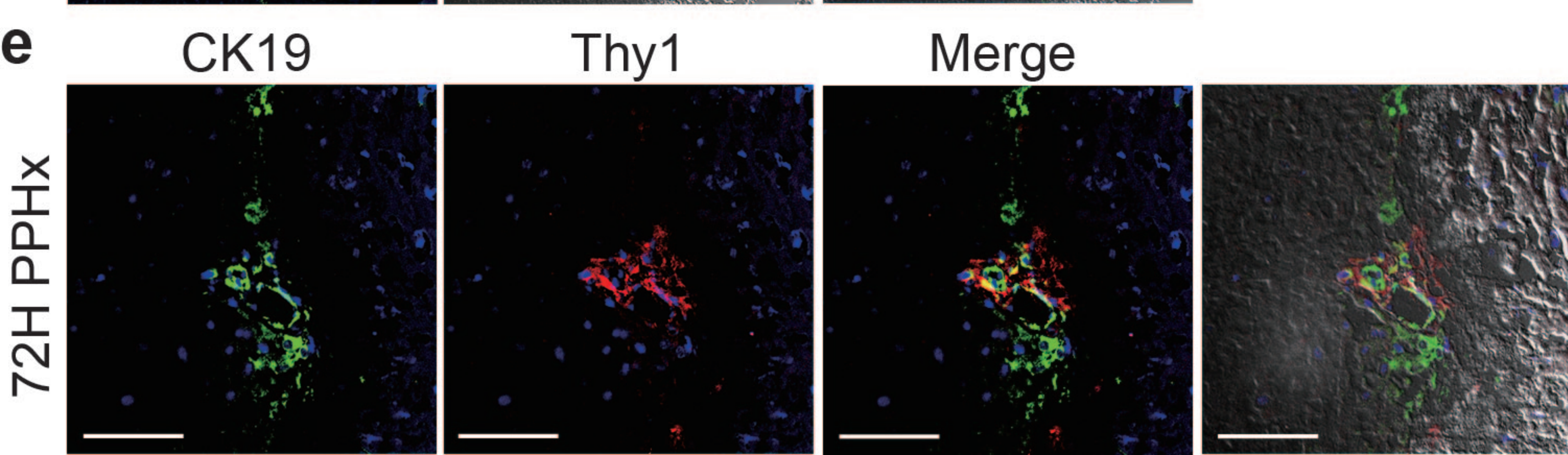
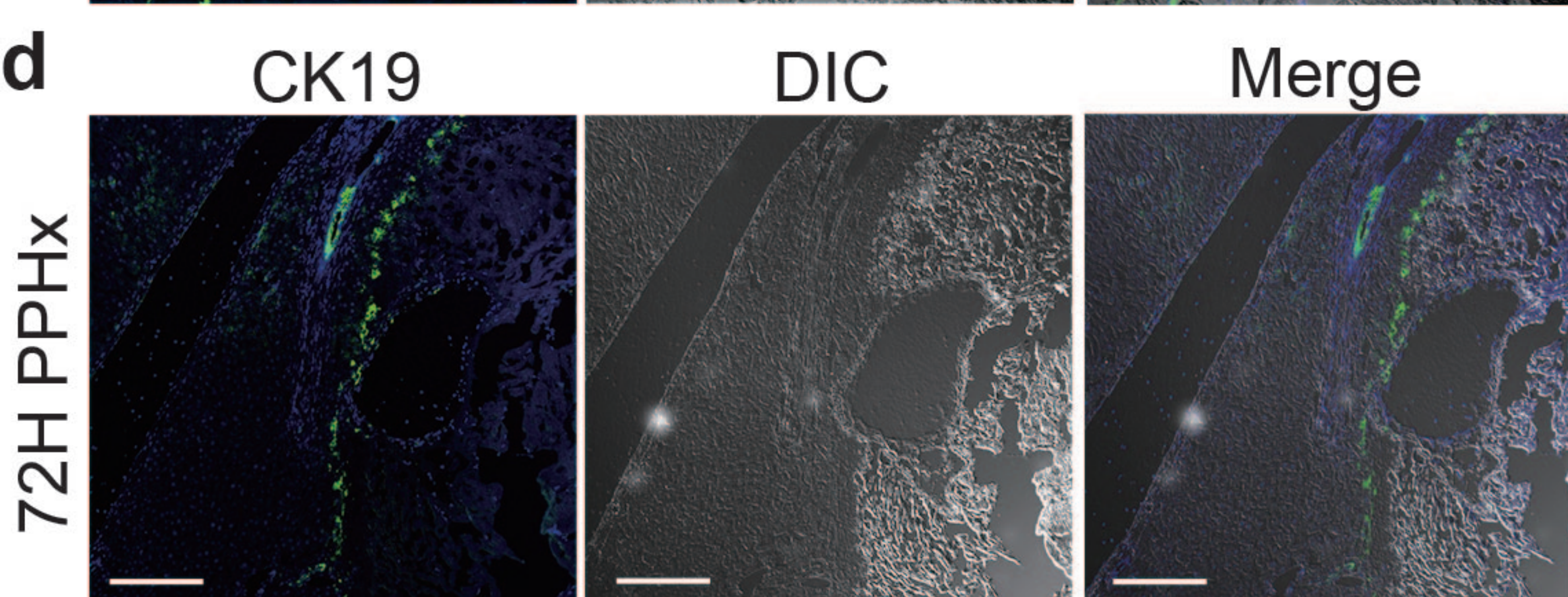
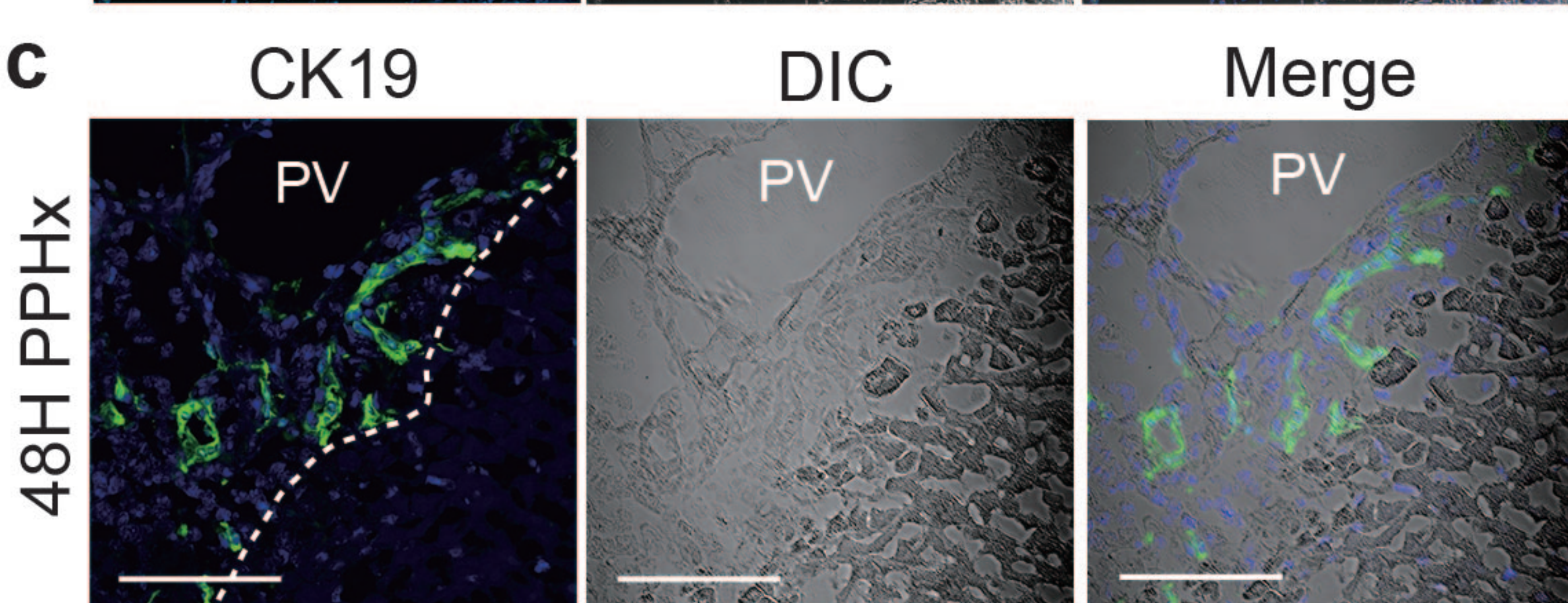
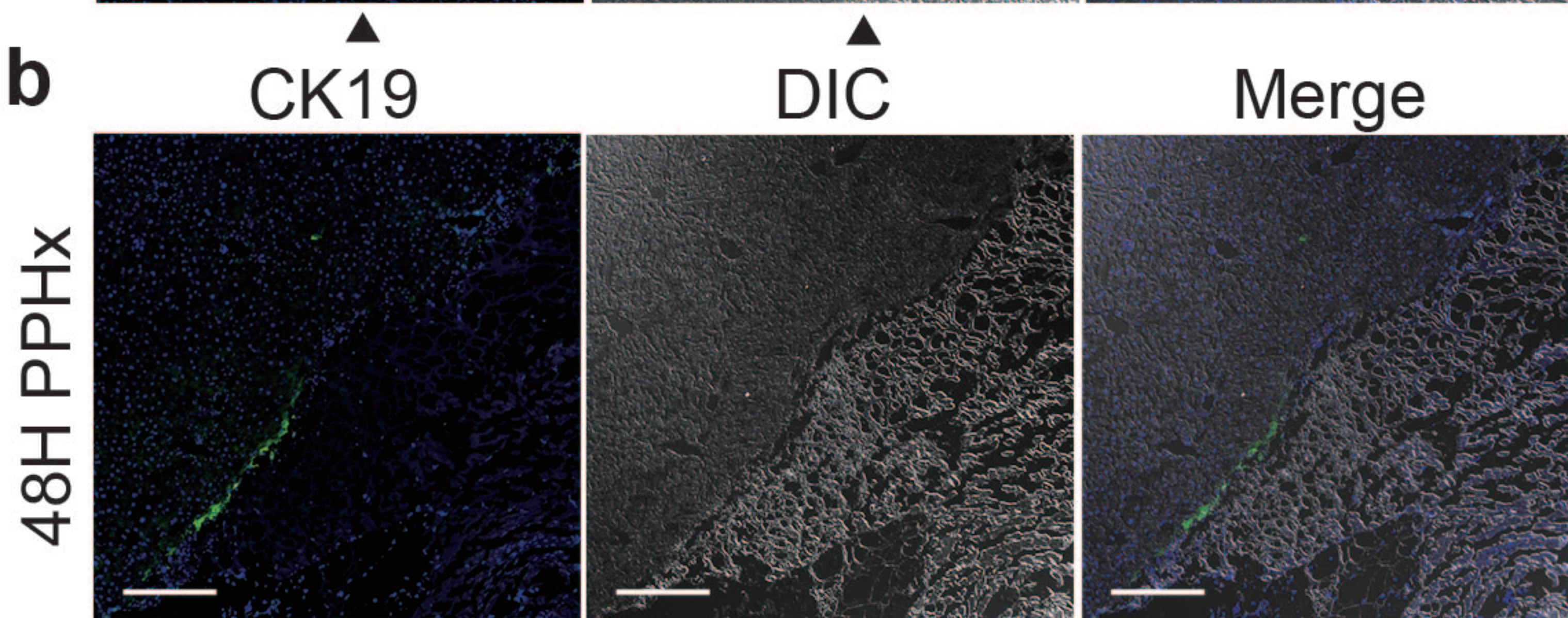
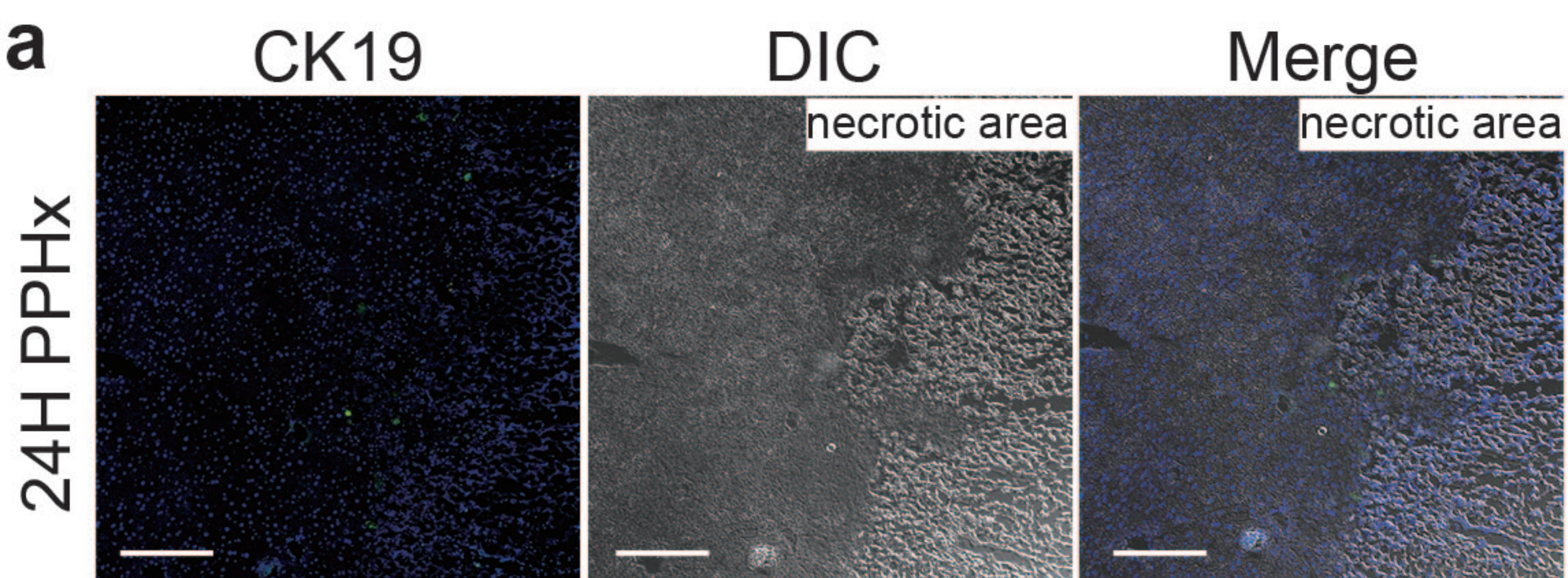


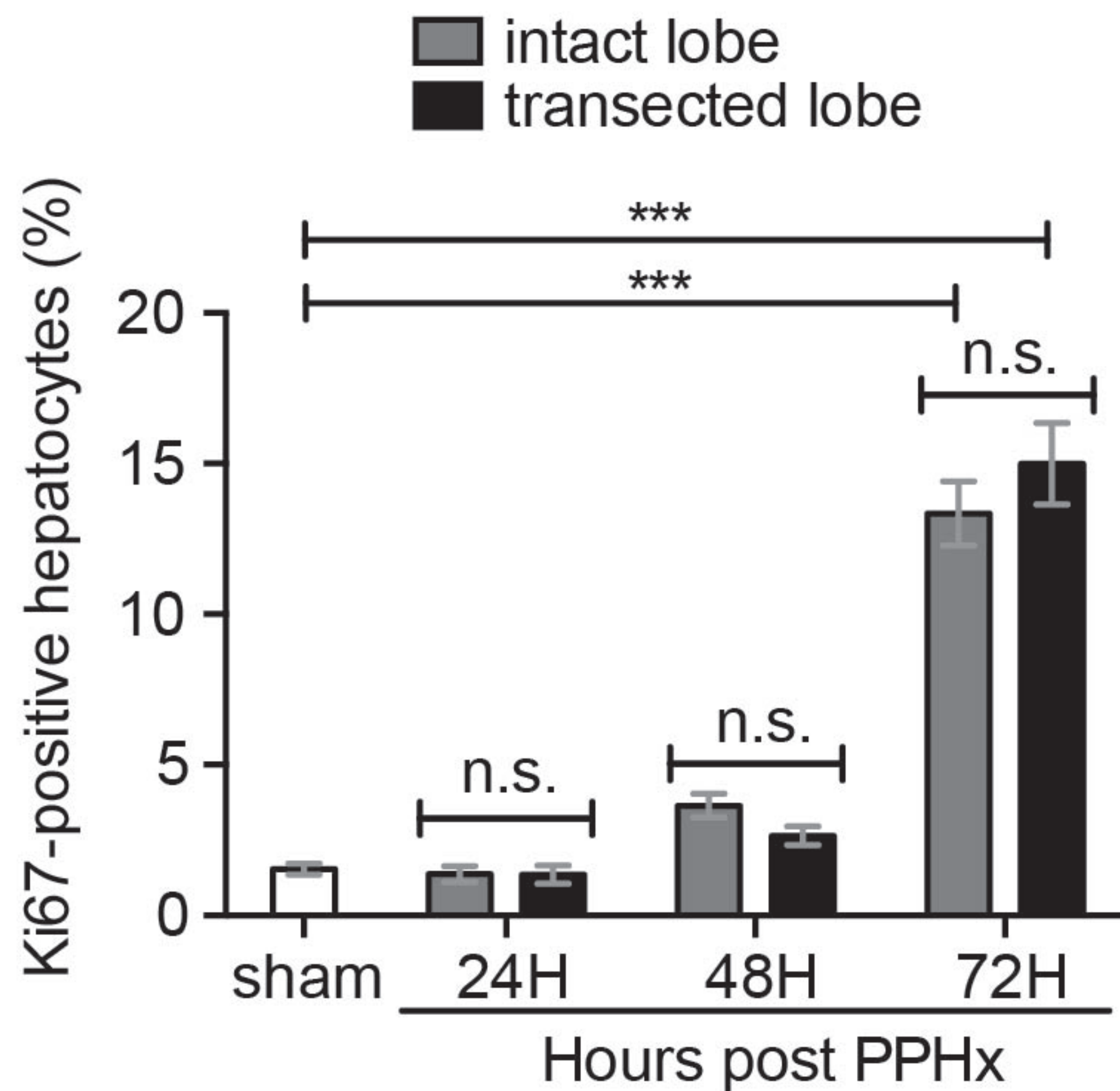
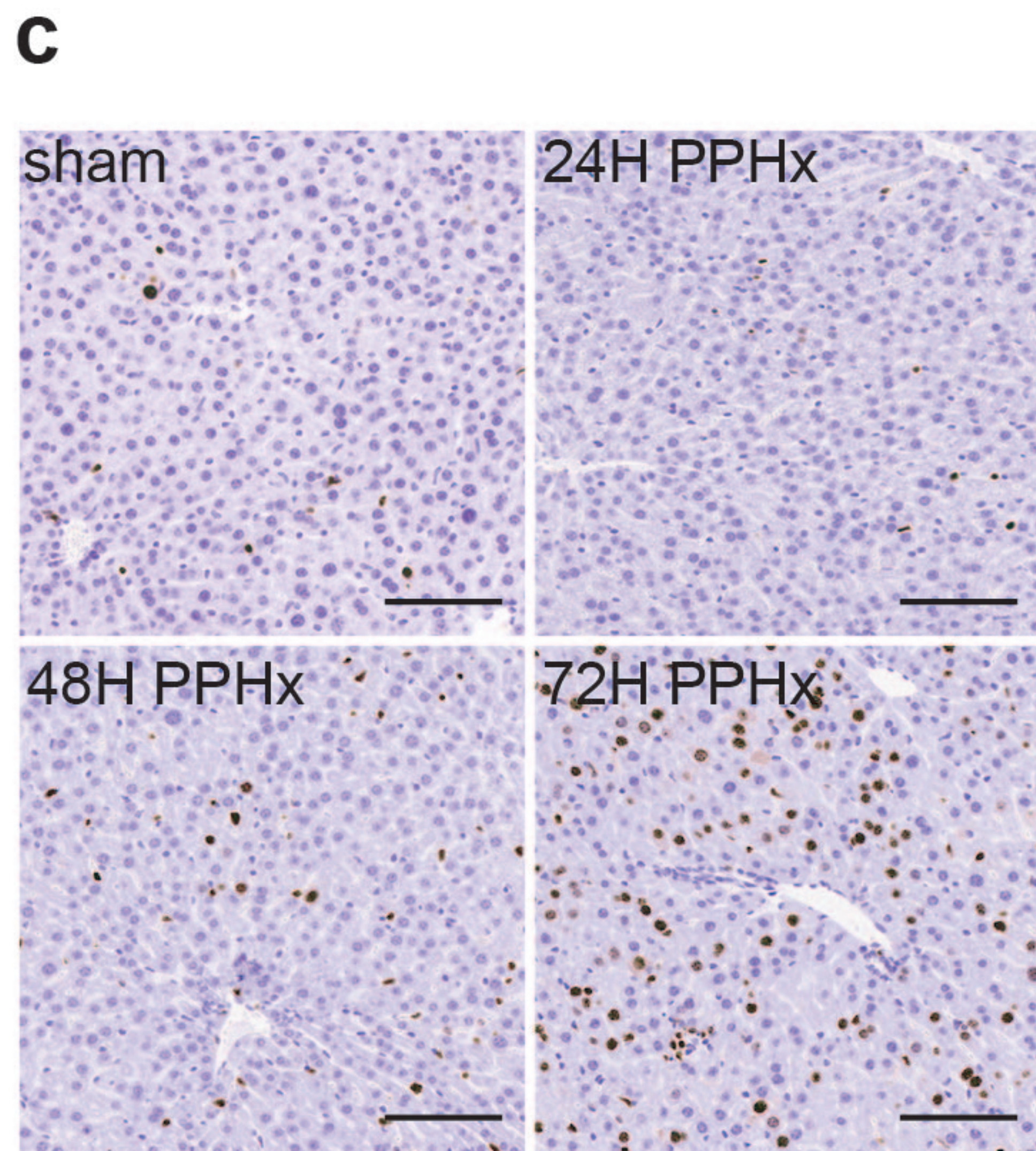
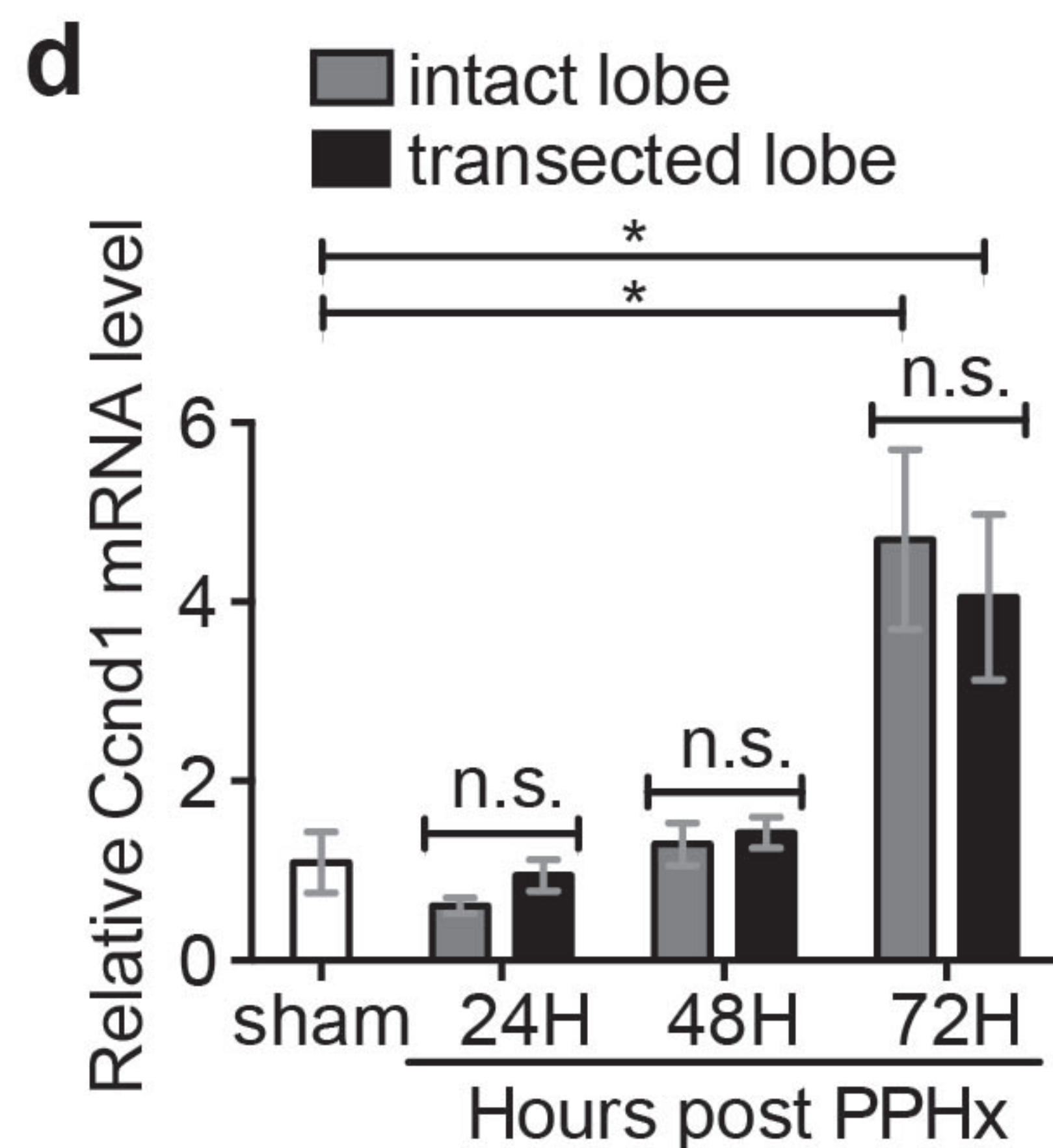
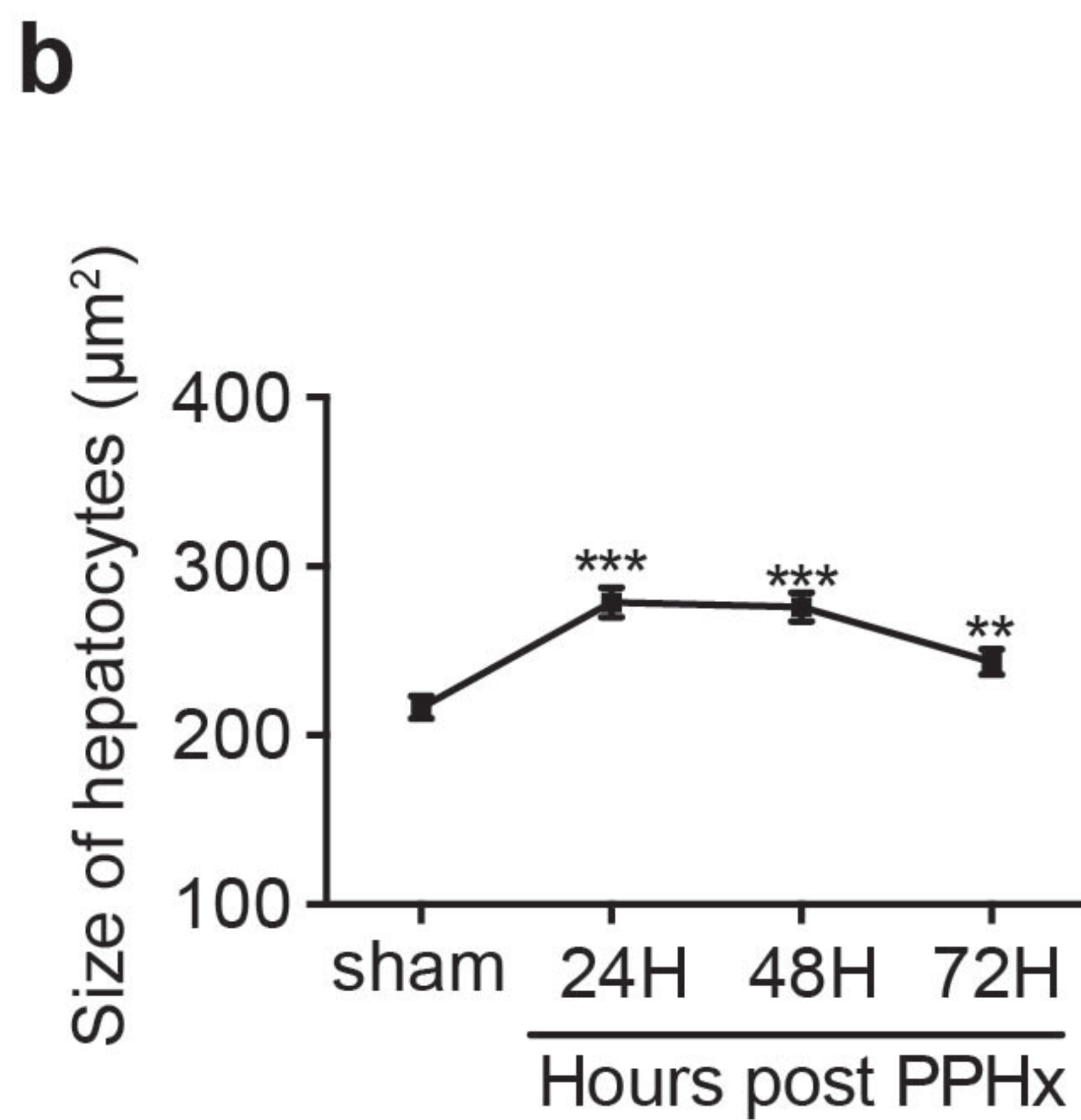
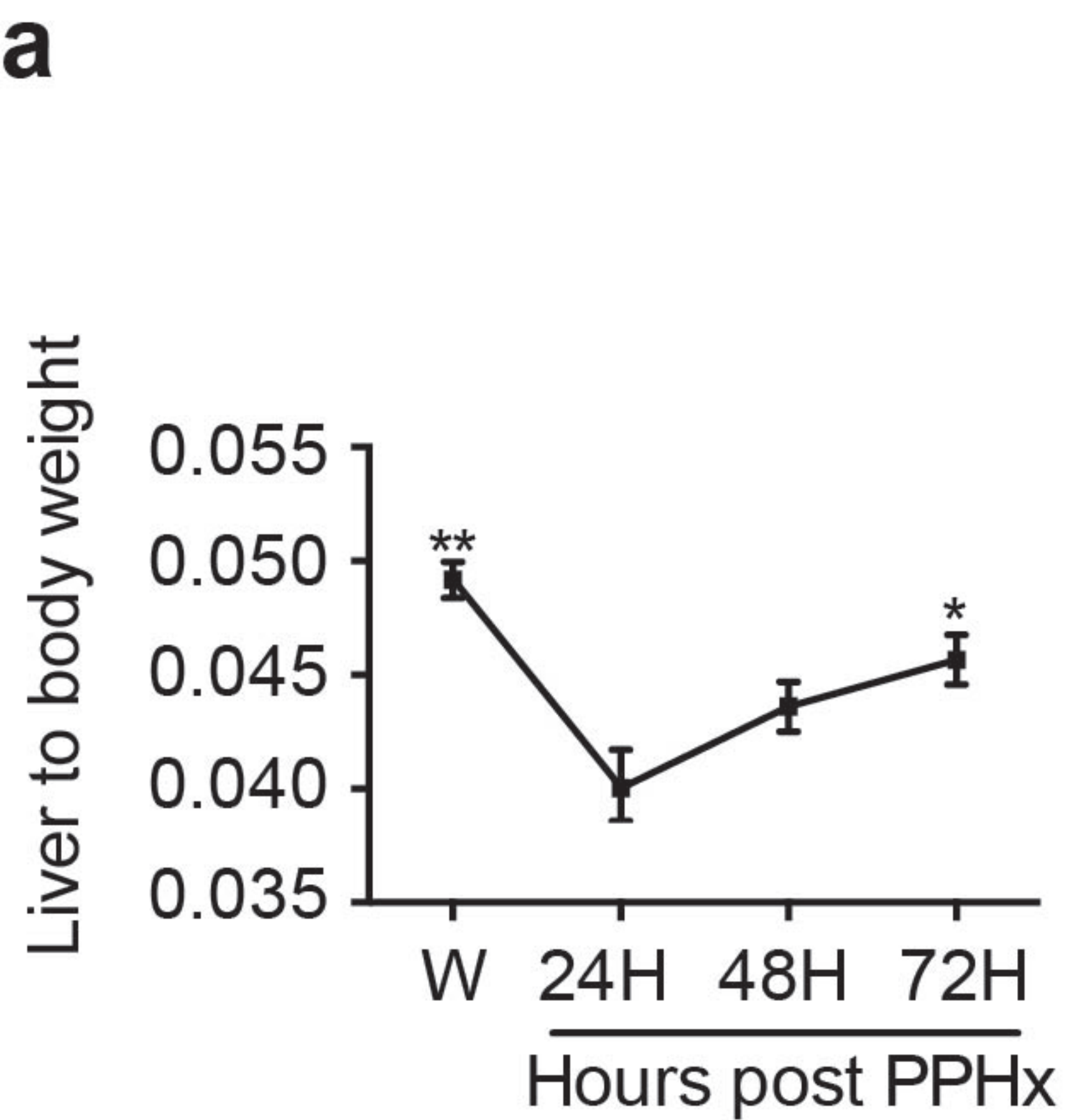
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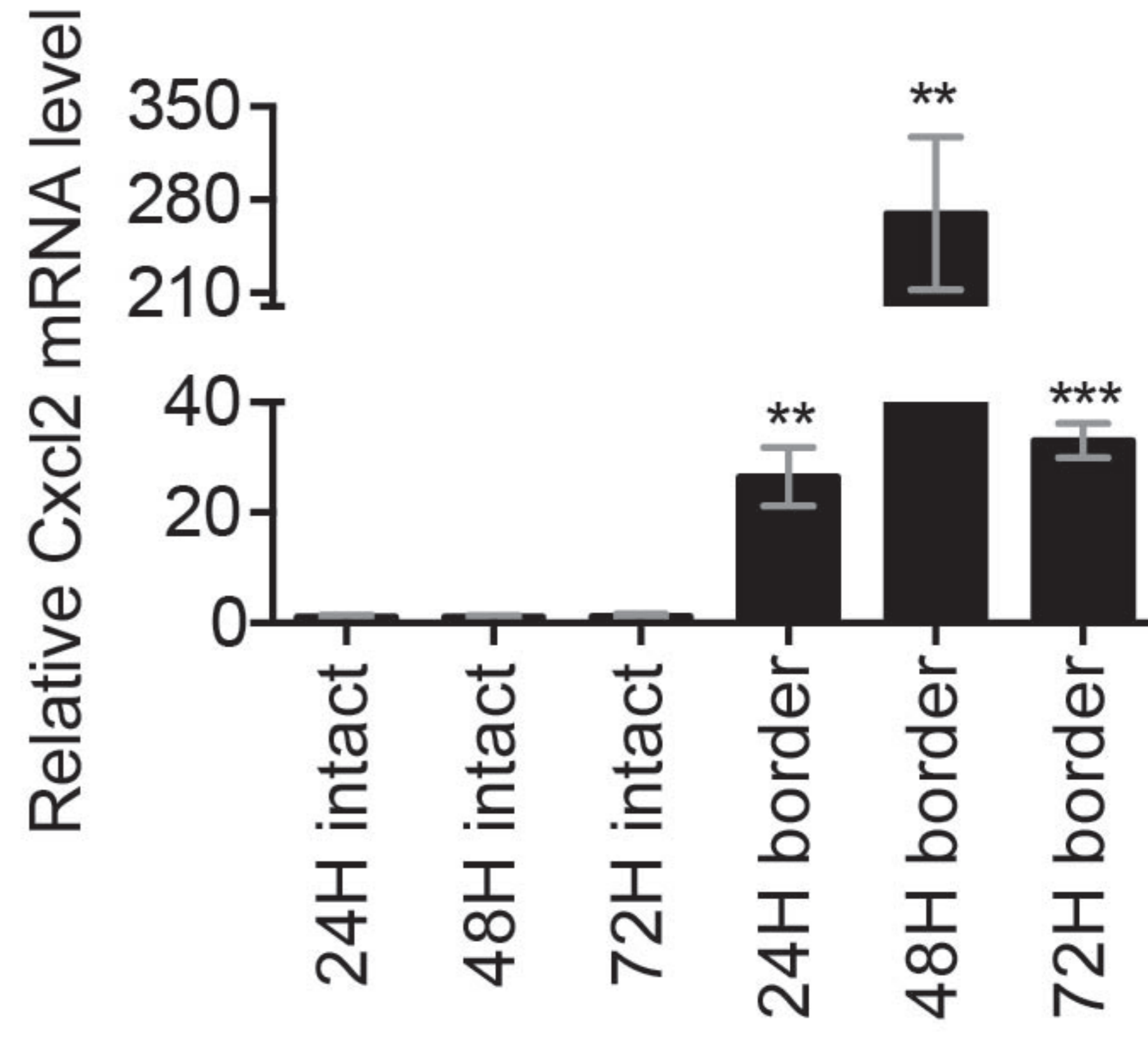
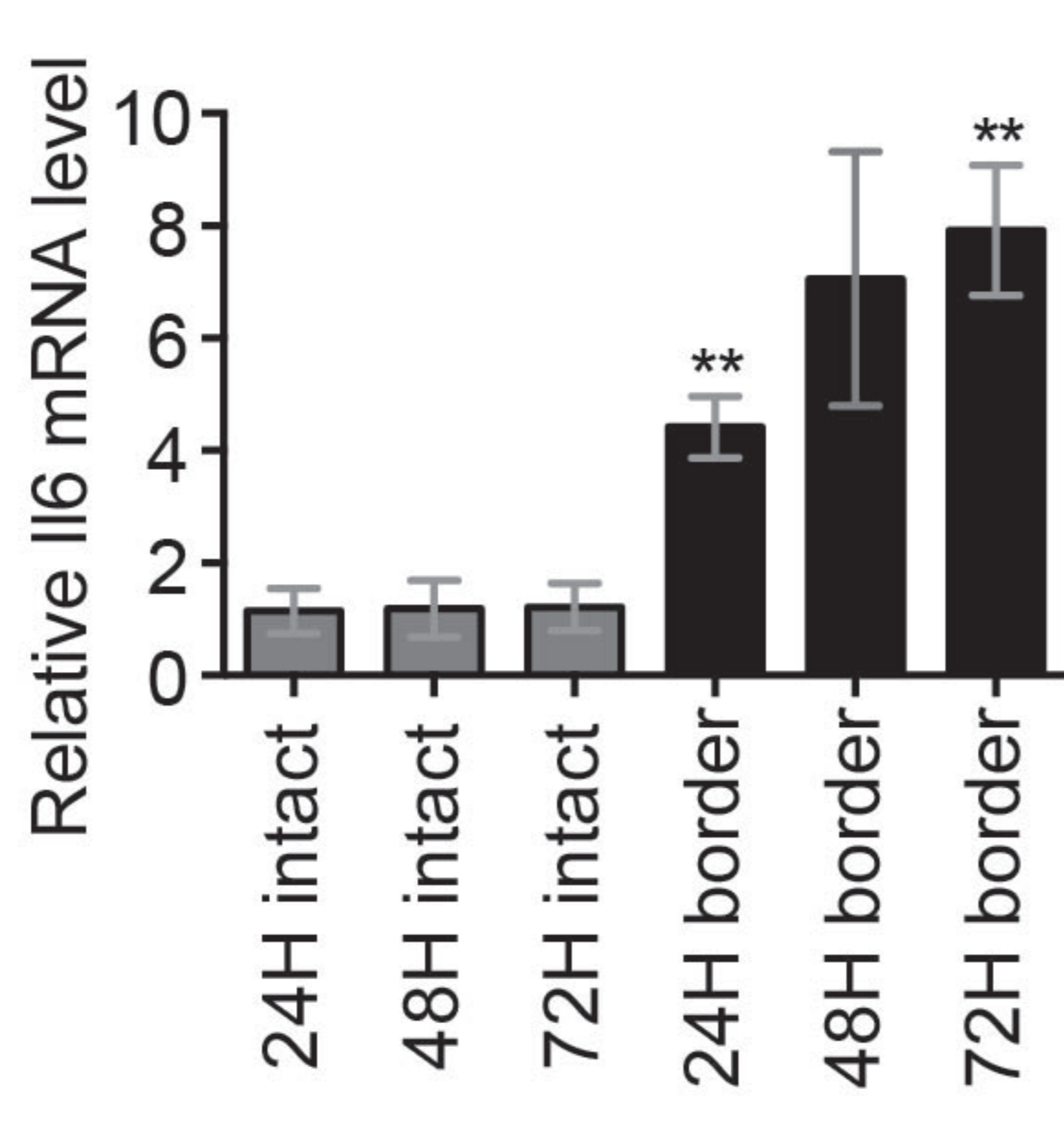
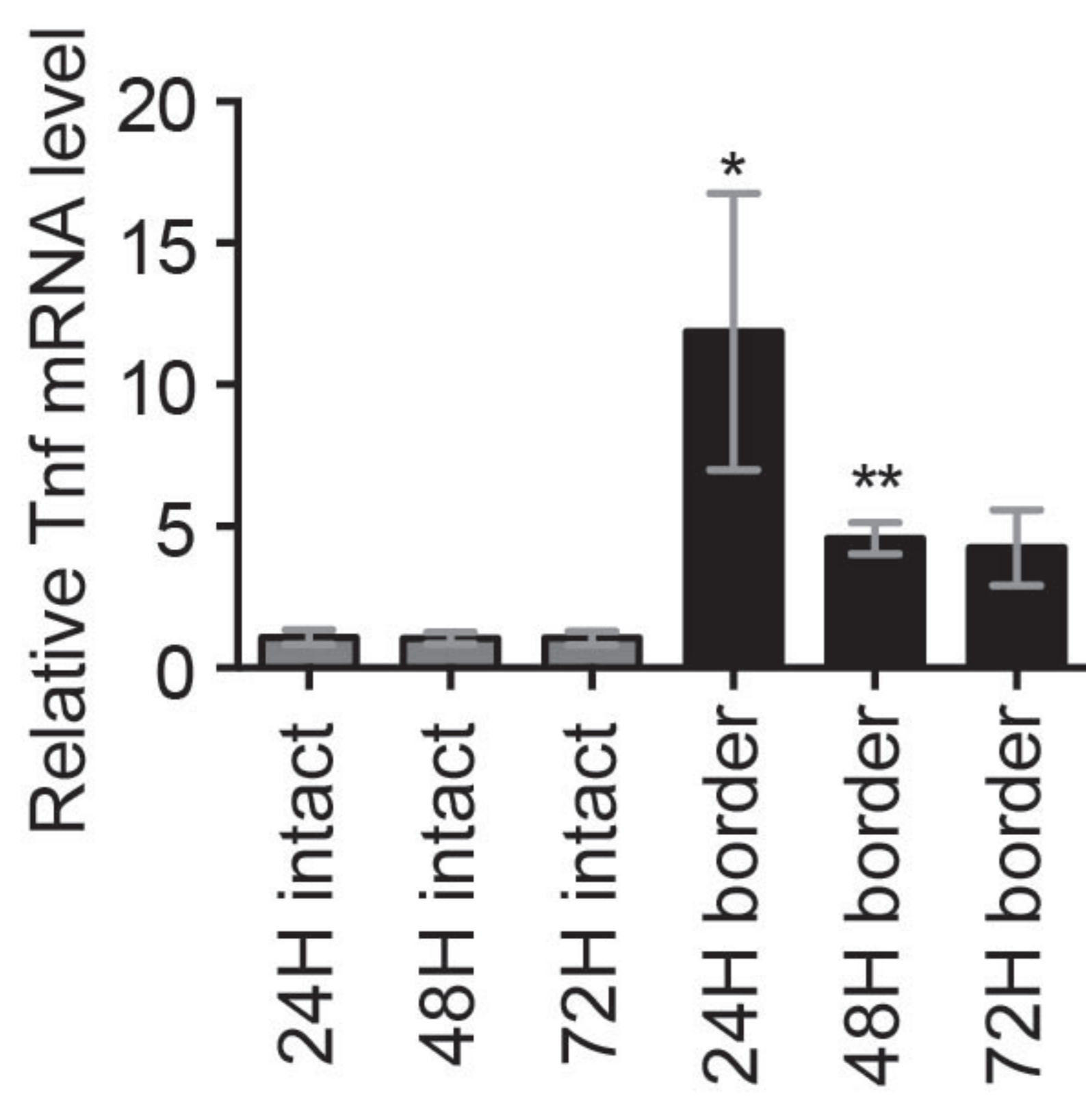
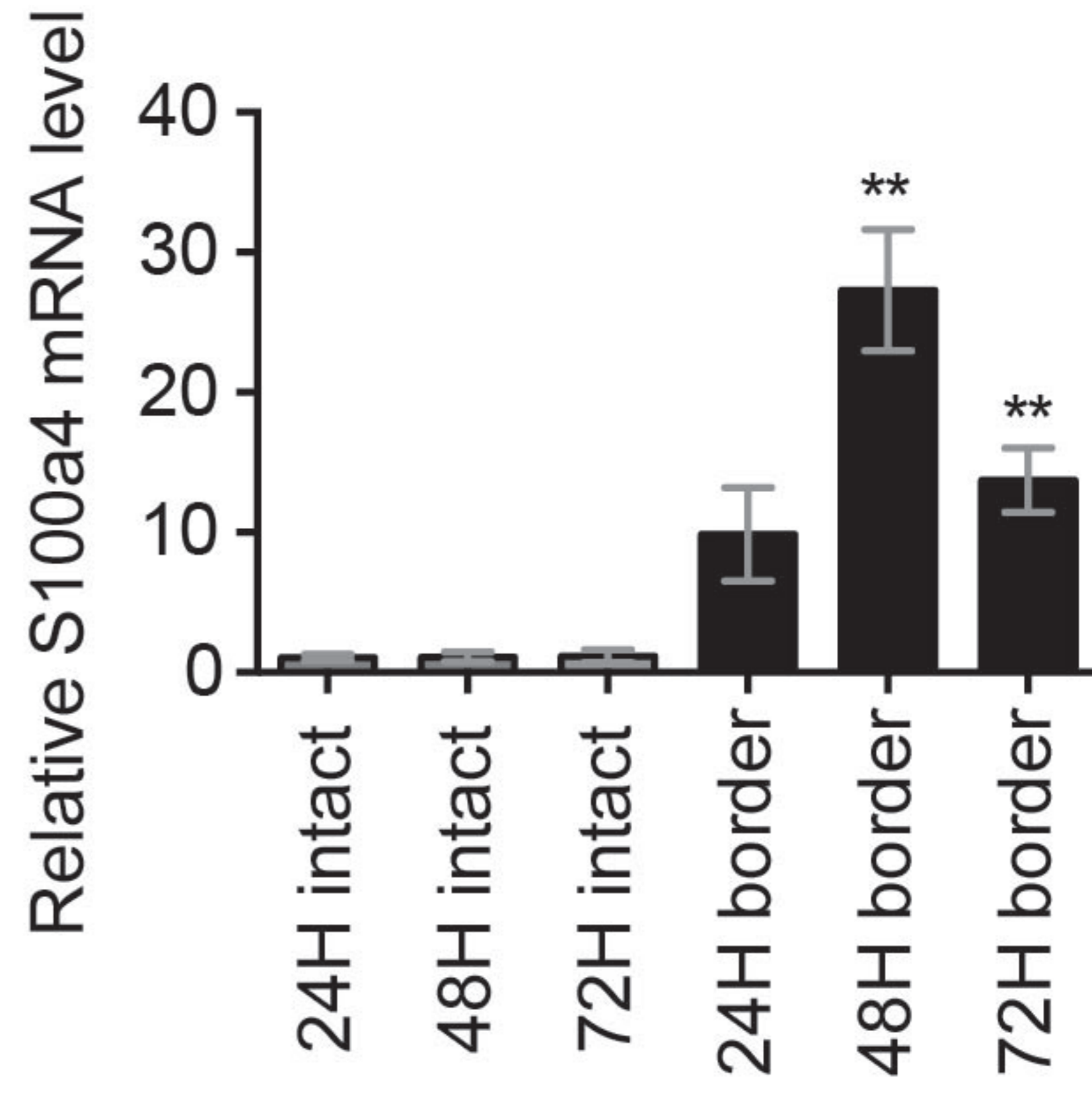
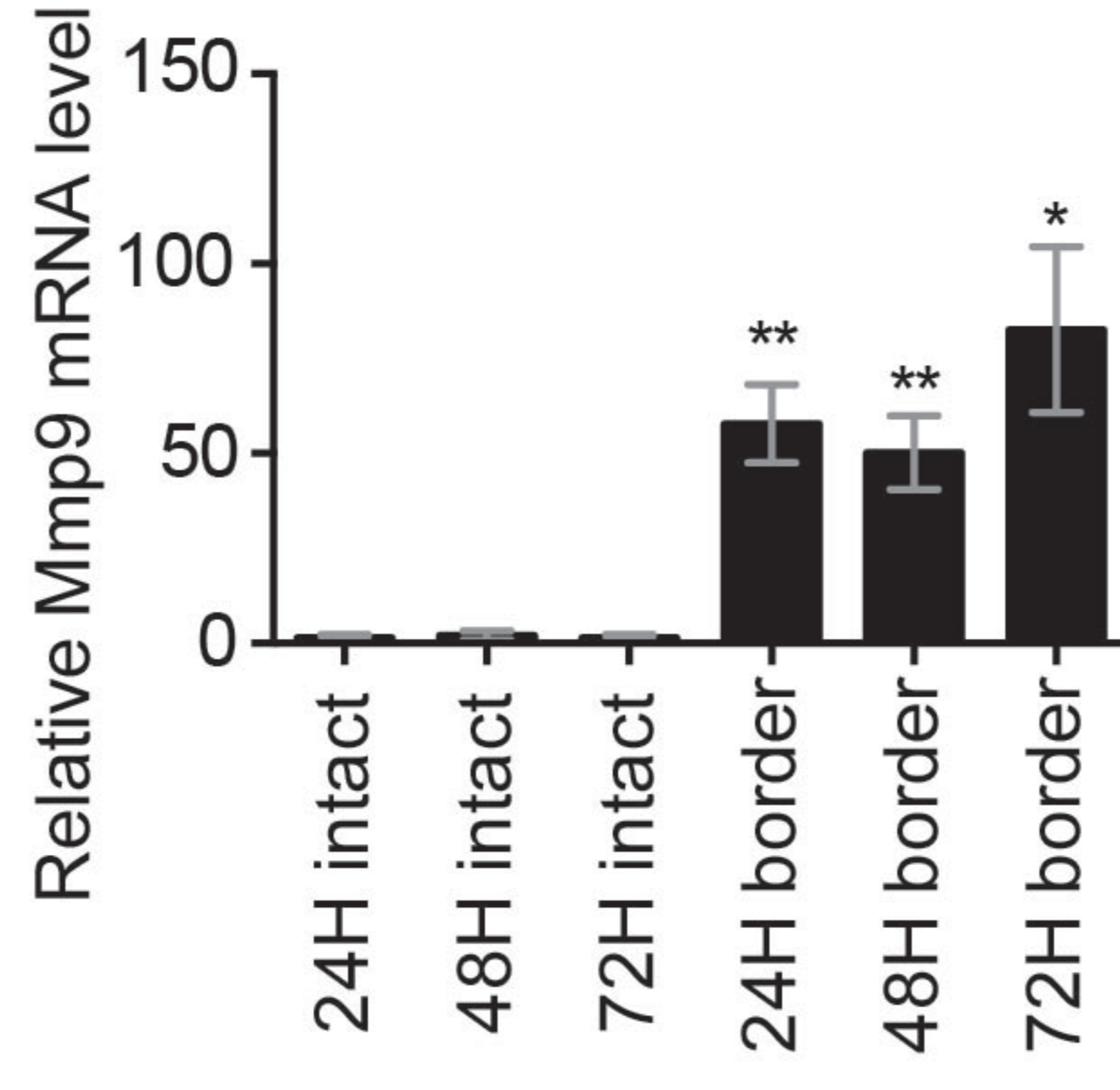
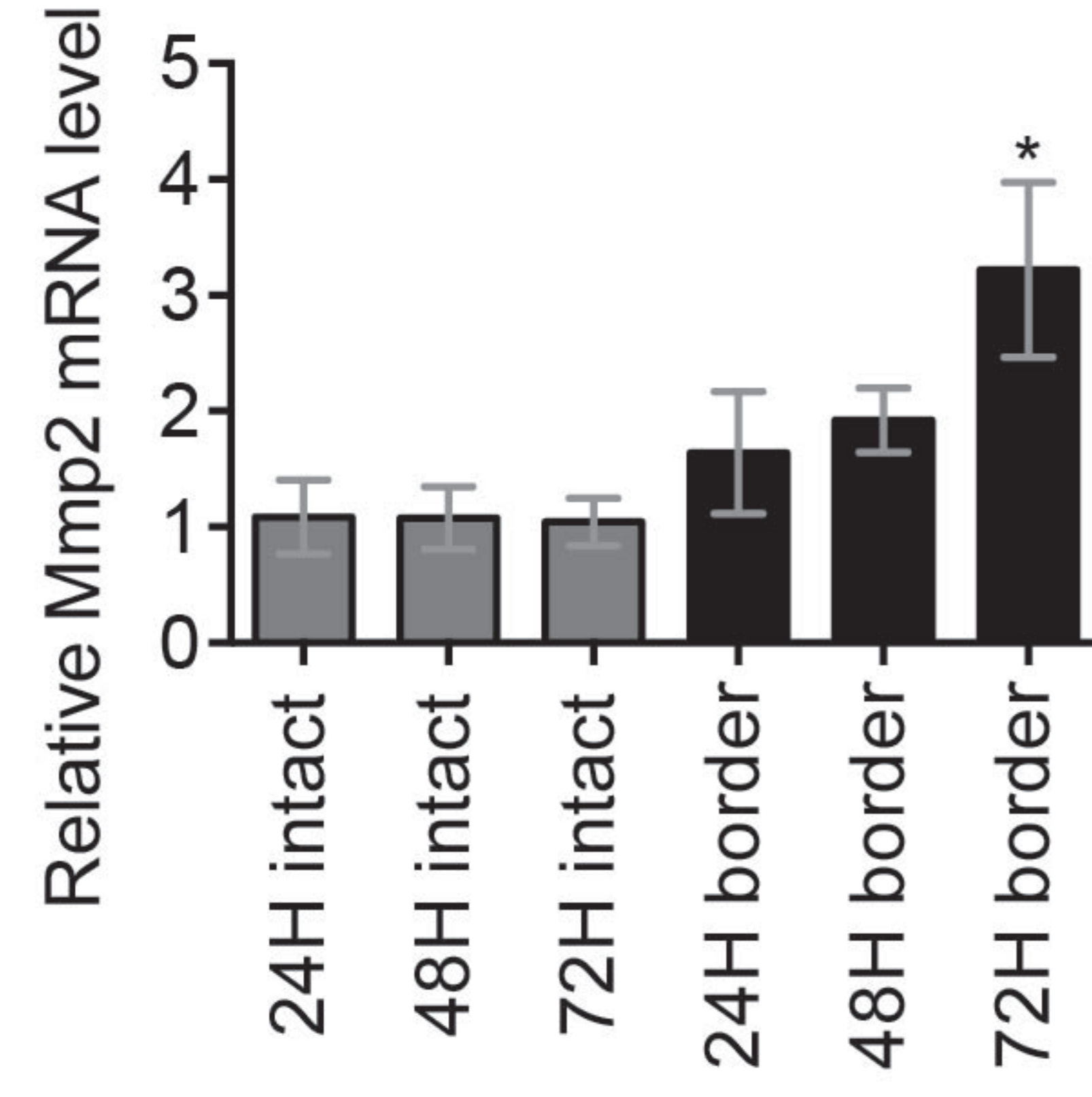
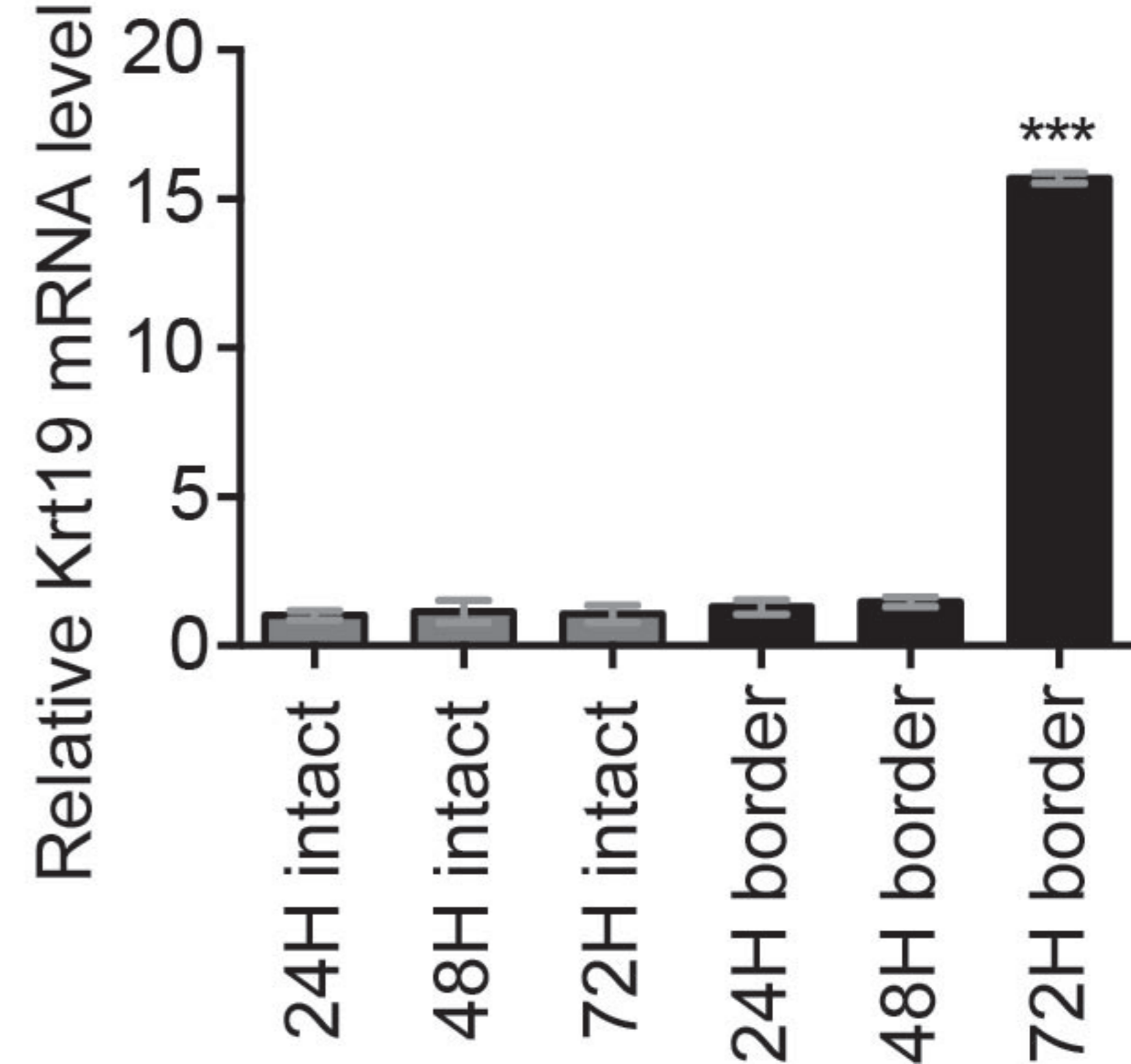
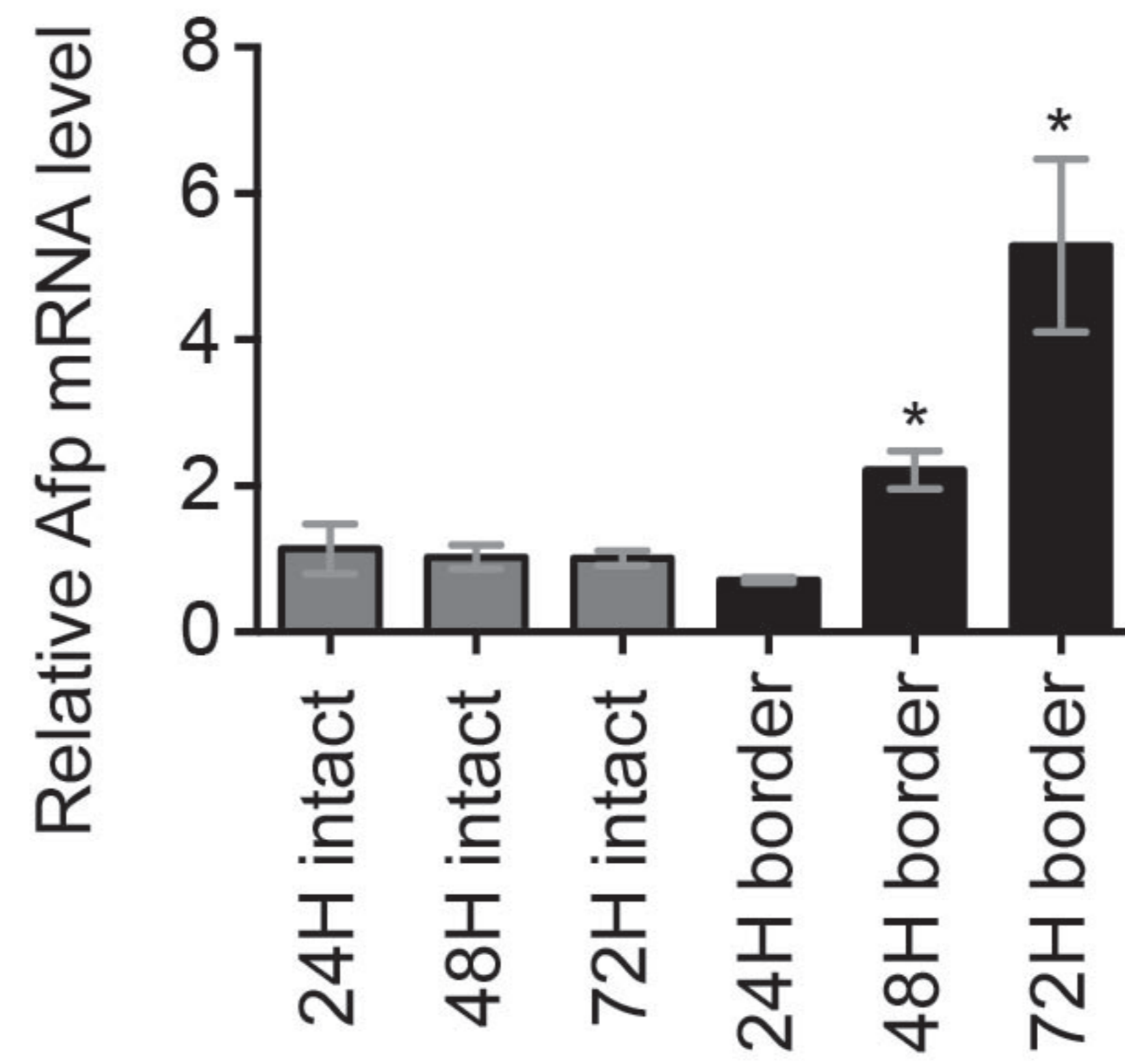
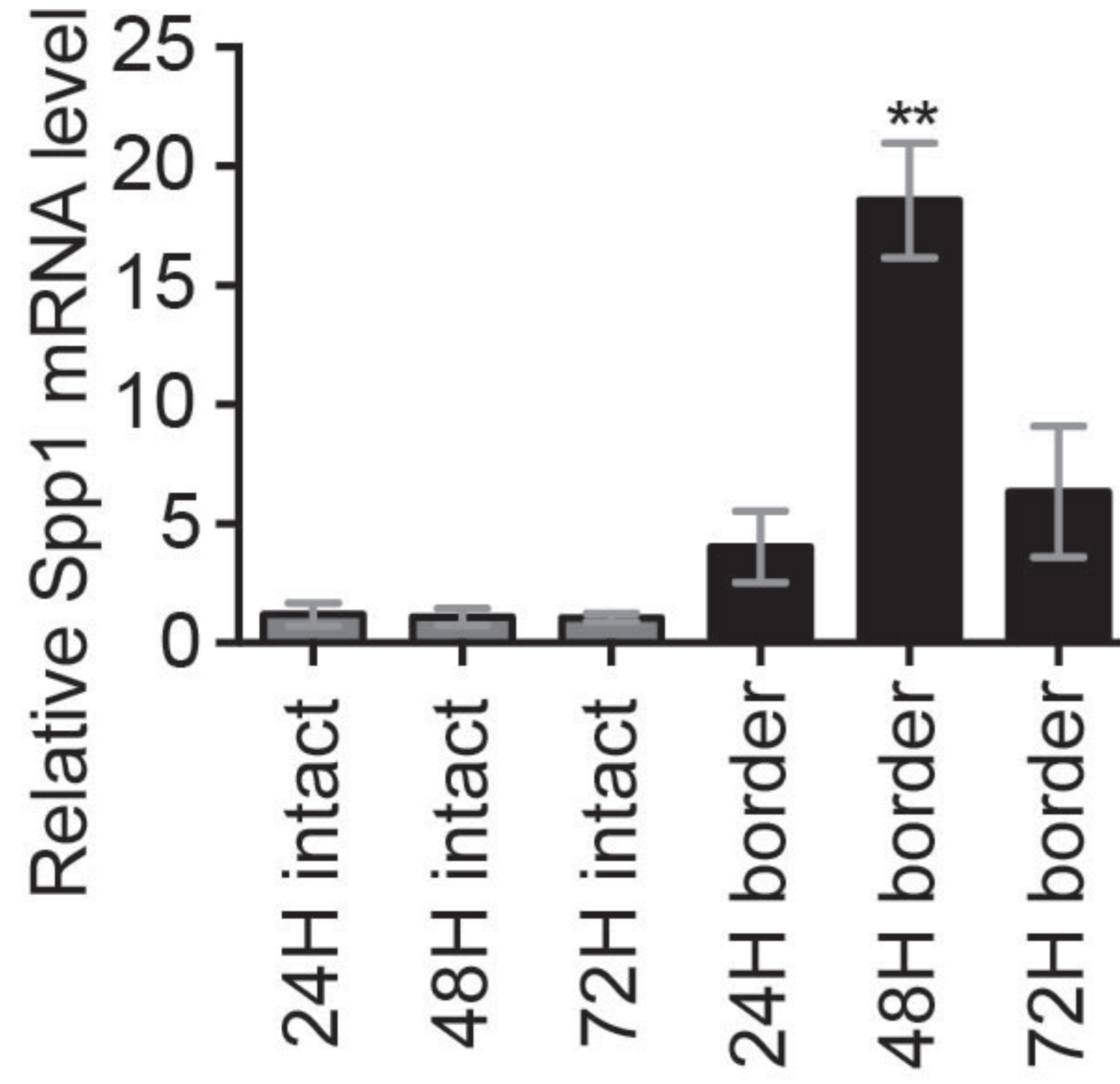


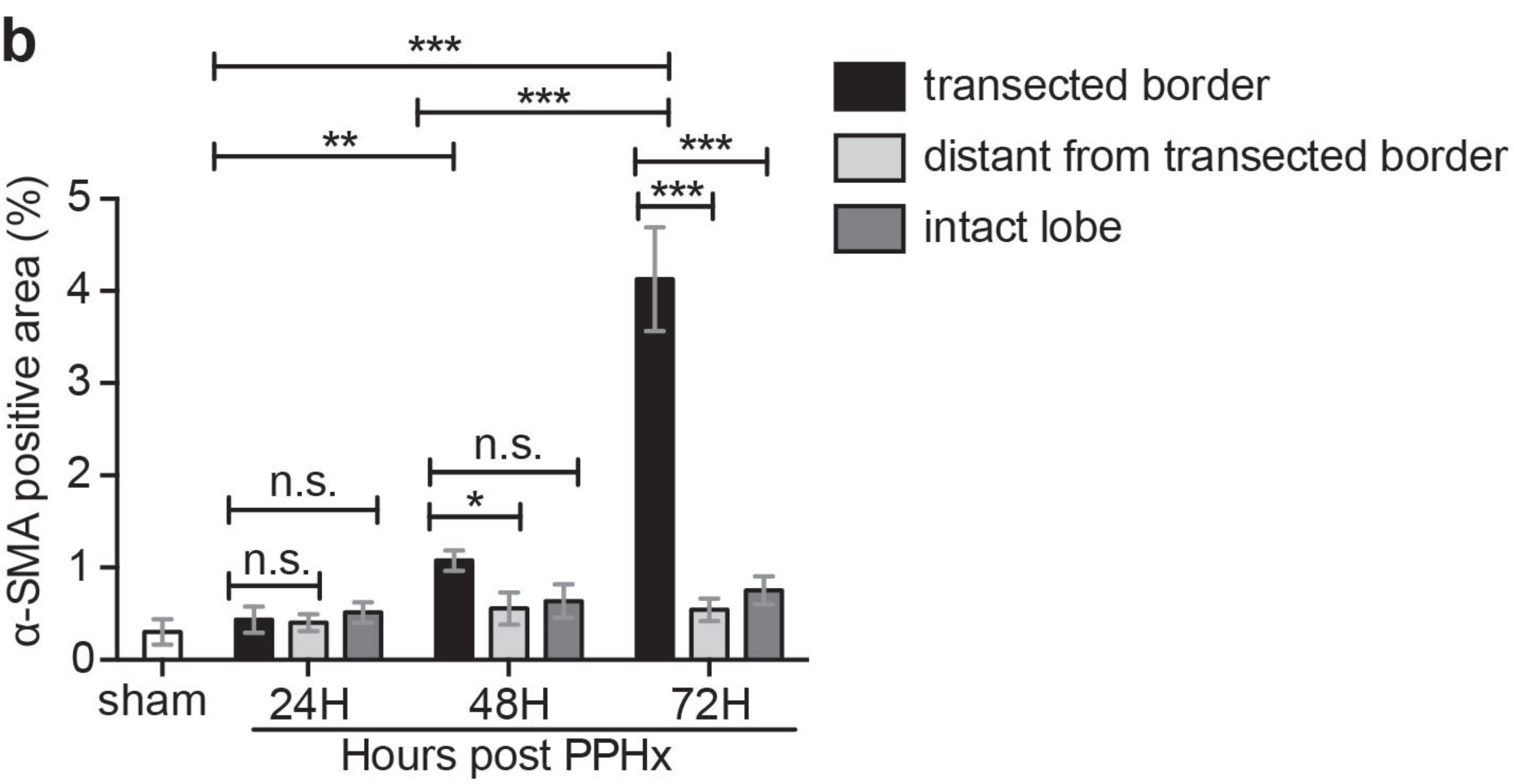
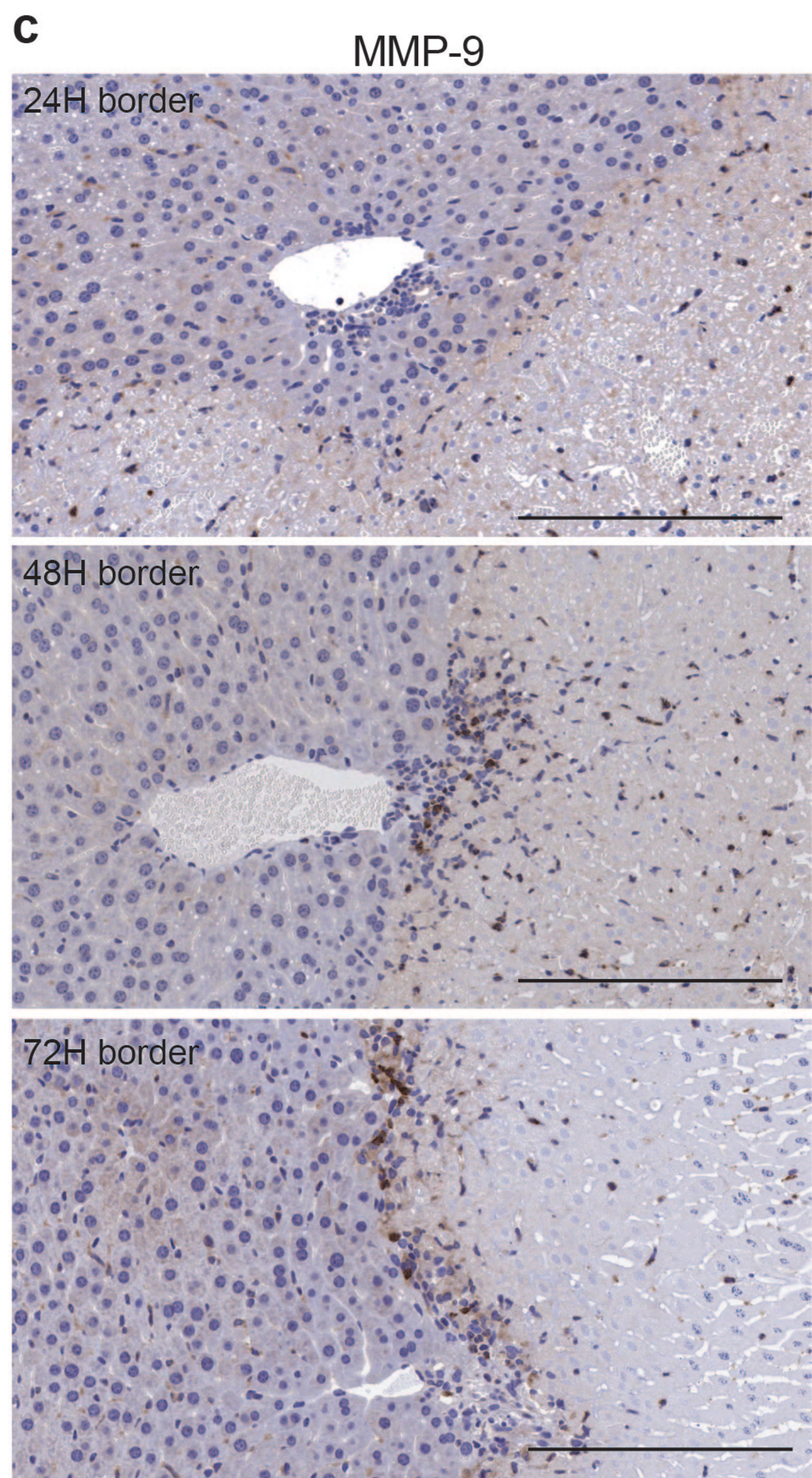
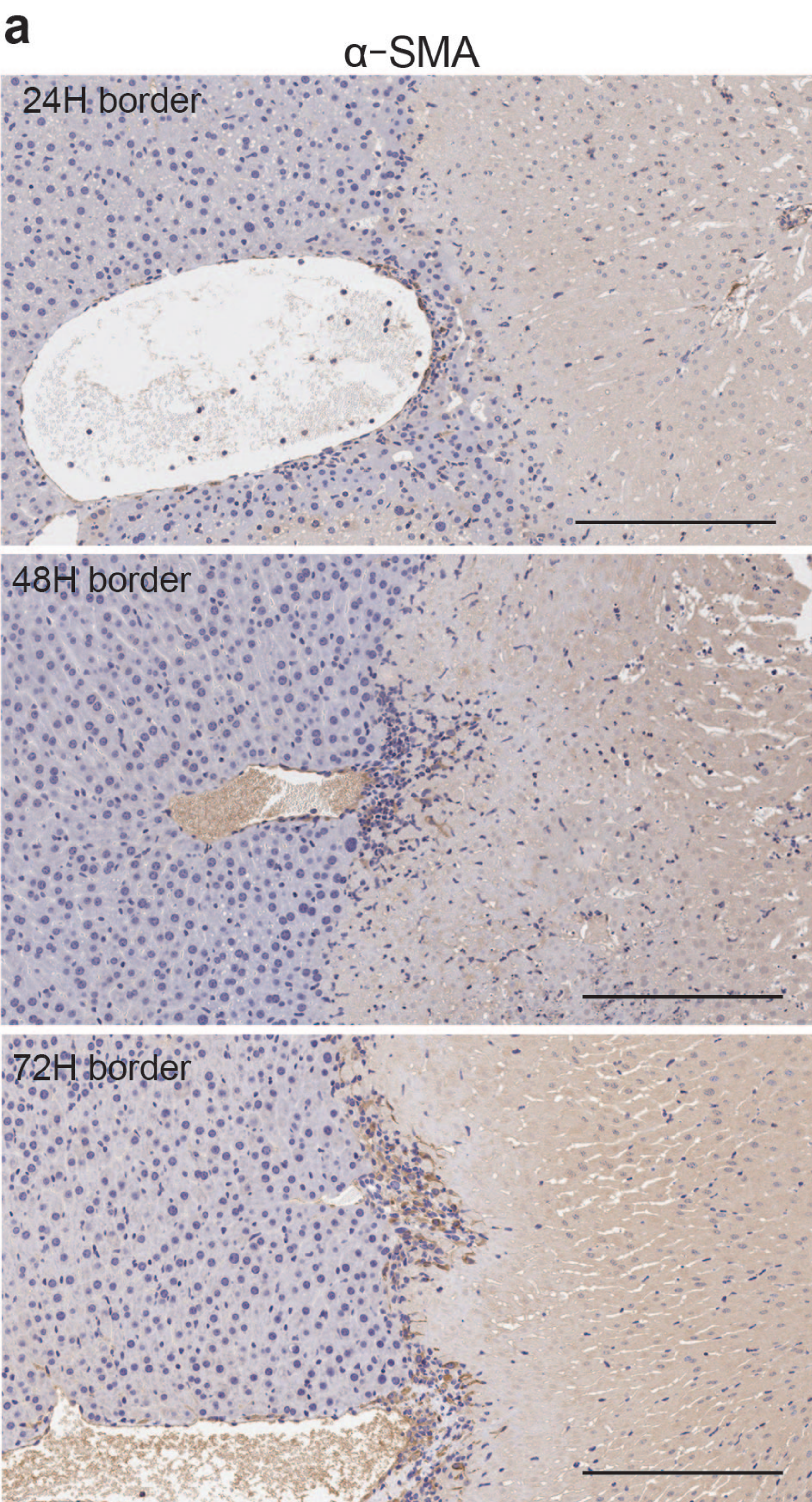
72H PPHx

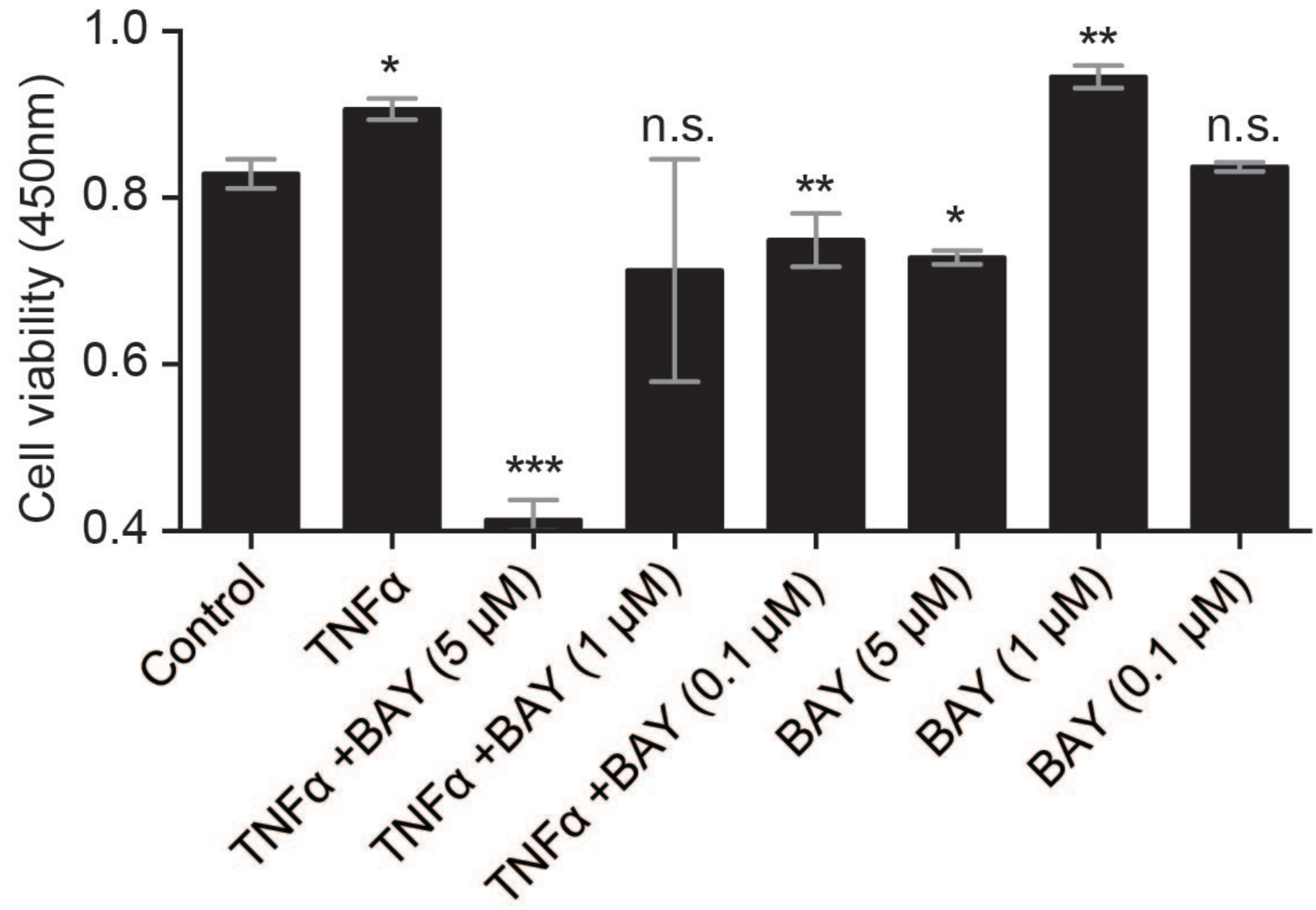




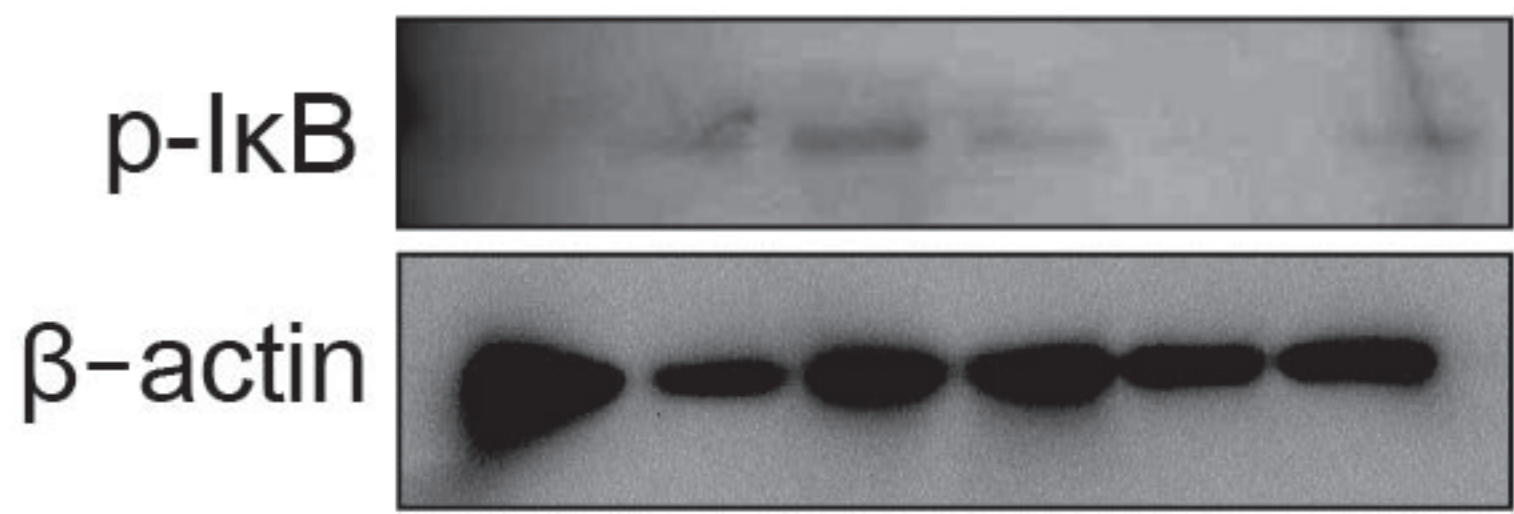


a**b****c**



a**b**

Control
 TNFα (10 ng/ml)
 TNFα (50 ng/ml)
 TNFα (50 ng/ml)+BAY (5 μM)
 TNFα (50 ng/ml)+BAY (1 μM)
 TNFα (50 ng/ml)+BAY (0.1 μM)

**c**

Control
 TNFα (10 ng/ml)
 TNFα (50 ng/ml)
 TNFα (50 ng/ml)+BAY (5 μM)
 TNFα (50 ng/ml)+BAY (1 μM)
 TNFα (50 ng/ml)+BAY (0.1 μM)

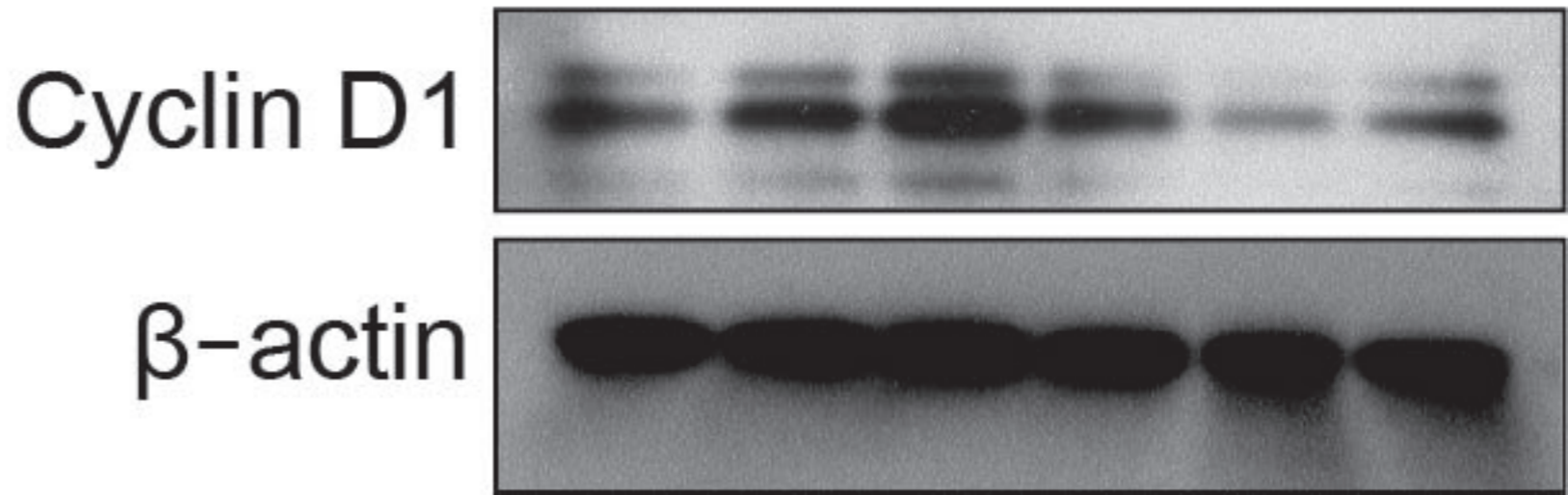


Table 1 Functional annotation clustering using DAVID

Cluster	Gene ontology term	Count	P
Up-regulated at 24-72 h (181 probe sets)	response to wounding	18	2.9E-9
	extracellular region	40	3.7E-9
	extracellular region part	26	1.0E-8
	chemokine activity	8	2.7E-8
Up-regulated at 24 h (626 probe sets)	inflammatory response	15	2.1E-4
	defence response	21	8.9E-4
	response to wounding	17	2.1E-3
	acute inflammatory response	6	2.5E-2
Up-regulated at 48 h (307 probe sets)	MHC class II protein complex	4	6.6E-5
	Antigen processing and presentation of peptide antigen via MHC classII	4	2.4E-4
Up-regulated at 72 h (477 probe sets)	extracellular matrix	20	1.3E-6
	proteinaceous extracellular matrix	19	3.0E-6
	extracellular region part	31	1.8E-5
Up-regulated at 24 h and 72 h (319 probe sets)	extracellular matrix	20	5.3E-8
	extracellular region part	32	7.1E-8
	proteinaceous extracellular matrix	19	1.5E-7
Up-regulated at 48-72 h (357 probe sets)	Cell cycle	72	7.5E-46
	M phase	50	2.0E-39
	cell cycle phase	51	2.0E-37
Down-regulated at any one point of time (564 probe sets)	extracellular region part	28	1.2E-3
	extracellular matrix	14	5.3E-3
	proteinaceous extracellular matrix	13	9.8E-3
Down-regulated at 24 h to 72 h (226 probe sets)	eye development	5	9.6E-3
	sensory organ development	6	1.2E-2

Functional enrichment of data presented in each cluster. P = P-value.