Mycolactone activation of Wiskott-Aldrich syndrome proteins underpins Buruli ulcer formation

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Mycolactone is a diffusible lipid secreted by the human pathogen *Mycobacterium ulcerans*, which induces the formation of open skin lesions referred to as Buruli ulcers. Here, we show that mycolactone operates by hijacking the Wiskott-Aldrich syndrome protein (WASP) family of actin-nucleating factors. By disrupting WASP autoinhibition, mycolactone leads to uncontrolled activation of ARP2/3-mediated assembly of actin in the cytoplasm. In epithelial cells, mycolactone-induced stimulation of ARP2/3 concentrated in the perinuclear region, resulting in defective cell adhesion and directional migration. In vivo injection of mycolactone into mouse ears consistently altered the junctional organization and stratification of keratinocytes, leading to epidermal thinning, followed by rupture. This degradation process was efficiently suppressed by coadministration of the N-WASP inhibitor wiskostatin. These results elucidate the molecular basis of mycolactone activity and provide a mechanism for Buruli ulcer pathogenesis. Our findings should allow for the rationale design of competitive inhibitors of mycolactone binding to N-WASP, with anti–Buruli ulcer therapeutic potential.

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Mycolactone activation of Wiskott-Aldrich syndrome proteins underpins Buruli ulcer formation

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physiological signals normally delivered by Rho GTPases to hijack WASP-dependent actin polymerization. We showed that mycolactone-induced activation of N-WASP in epithelial cells and the consequent dynamic rearrangements of the actin cytoskeleton dramatically impaired the integrity of the epidermis, thus providing a molecular mechanism underpinning Buruli ulcer pathogenesis.

**Results**

*Mycolactone binds selectively to the WASP/N-WASP regulators of actin polymerization.* To analyze the effect of mycolactone on the cytoskeleton, we selected the HeLa cell line, as a model of anchorage-dependent cells of the human epithelial system, and the Jurkat cell line, as a model of nonadherent, immune cells. Cells were treated with mycolactone for 5 minutes, then spun onto poly-L-lysine–coated glass slides. Actin structures were visualized by phalloidin in cells exposed to mycolactone (Myco) or solvent as control (Ctrl). Scale bars: 10 μm. Graphs show mean cell proportions (± SEM) on more than 50 cells from at least 2 independent experiments. **P < 0.01, ***P < 0.001, unpaired 2-tailed t test. (B) Amino acid sequences of the GBD of WASP, N-WASP, and PAK, with boxes outlining the BR and CRIB motifs. (C) Constructs used in this study. (D) Binding of biotinylated mycolactone to these constructs in ELISA. Data are mean ± SD A<sub>490</sub> nm of duplicates.

Mycolactone binds to WASP/N-WASP with high affinity and specificity. (A) HeLa cells were plated on Y-shaped, fibronectin-coated micropatterns, then exposed to 20 nM mycolactone for 30 minutes. Jurkat T cells were exposed to 20 nM mycolactone for 5 minutes, then spun onto poly-L-lysine–coated glass slides. Actin structures were visualized by phalloidin in cells exposed to mycolactone (Myco) or solvent as control (Ctrl). Scale bars: 10 μm. Graphs show mean cell proportions (± SEM) on more than 50 cells from at least 2 independent experiments. **P < 0.01, ***P < 0.001, unpaired 2-tailed t test. (B) Amino acid sequences of the GBD of WASP, N-WASP, and PAK, with boxes outlining the BR and CRIB motifs. (C) Constructs used in this study. (D) Binding of biotinylated mycolactone to these constructs in ELISA. Data are mean ± SD A<sub>490</sub> nm of duplicates.
Together, these data support the view that a ternary complex can be formed in which CDC42-GTP and mycolactone bind distinct sites on WASPs. Like CDC42, mycolactone activate WASPs by releasing the VCA from the GBD. However, with a 100-fold higher affinity (26), mycolactone is a much stronger activator of N-WASP–mediated actin assembly.

Mycolactone activates cellular WASP/N-WASP to promote actin assembly. To evaluate the physiological relevance of these findings, we then coated latex beads with mycolactone, incubated them with N-WASP, and placed them in Jurkat cell extracts spiked with fluorescently labeled actin. A fluorescent signal readily developed at the surface of mycolactone-coated beads incubated with N-WASP, and placed them in Jurkat cell extracts spiked with fluorescently labeled actin. A fluorescent signal readily developed at the surface of mycolactone-coated beads incubated with N-WASP, but not control beads that either had not been coated with mycolactone or had not been incubated with N-WASP (Figure 3A), which demonstrated that N-WASP activated by mycolactone could induce the formation of branched actin filaments in a cellular environment. We then took advantage of a fluorescent derivative of mycolactone to study its distribution in living cells (Supplemental Figure 4 and ref. 27). As shown in Figure 3B, mycolactone was distributed primarily in the cytosolic compartment of treated Jurkat cells. A limited, but significant, colocalization with active WASP was observed after 1 hour, using an antibody that preferentially binds the open form of the protein (Figure 3B). Notably, spots of ARP2/3 complex were detected in the area of Bodipy-mycolactone/active WASP colocalization, which indicates that mycolactone reaches the sites of WASP-mediated ARP2/3 activation. After 4 hours, immunofluorescence detection of ARP2/3 revealed an increased level of activated complex in the perinuclear area of HeLa cells (Figure 3C). These alterations persisted for more than 16 hours and were efficiently suppressed by coincubation with the N-WASP inhibitor wiskostatin (28). They were not due to increased protein level, as expression of WASP, N-WASP, the p34 subunit of ARP2/3 (p34-ARP2/3), and actin was unchanged in mycolactone-treated cells (Figure 3D). Taken together, the data in Figure 3 indicated that mycolactone promotes perinuclear recruitment and activation of ARP2/3 by N-WASP.

Activation of N-WASP in epithelial cells mediates mycolactone-induced adhesion defects. Since N-WASP is a major regulator of cell adhesion...
(21, 22), we next explored its contribution to mycolactone effects on cell-cell and cell-matrix contacts. Live imaging of HeLa cells exposed to mycolactone revealed that the early induction of spike-like protrusions was followed by lamellipodium retraction (Figure 4A). If exposure to mycolactone was prolonged over 24 hours, this process culminated in cell rounding and detachment (Figure 4, A and B). Induction of cell apoptosis, as measured by caspase activation (early stages) and membrane permeability to propidium iodide (late stages), was primarily observed in the detached fraction of mycolactone-treated cells (Figure 4C), which strongly suggested that mycolactone kills epithelial cells by anokis. Strikingly, 4 hours of exposure to mycolactone was sufficient to induce a near-complete and persistent incapacity to adhere to plastic wells (Figure 4D). This effect was counteracted by wiskostatin or by prior transfection of the cells with WASP CR1 (Figure 4, D and E), demonstrating its association with activated N-WASP.

In HeLa cell monolayers treated with mycolactone for 16 hours, cell retraction was associated with a marked reduction in the number of intercellular contacts (Figure 5A). Cadherin staining showed that mycolactone suppressed both the maintenance of mature cell-cell junctions and the active formation of new interfaces via an adhesion zipper (Figure 5B). In polarized systems, such as the MDCK cell line model, mycolactone induced marked suppression of adherens and tight junctions, as shown by the decrease in the fluorescent ring–like signal of cadherins and zona occludens–1 (ZO-1; Figure 5C). Confocal imaging of basal sections revealed decreased stress fibers, increased perinuclear actin rings, and impaired membrane localization of cadherin in mycolactone-treated cells, resulting in disruption of the honeycomb-like organization and, eventually, rupture of the monolayer (Figure 5, D and E). Mycolactone impairs direction sensing in migrating epithelial cells. The directed migration of epithelial cells requires not only activation...

**Figure 3**

Mycolactone activates cellular WASPs. (A) Streptavidin-latex beads were coated with biotinylated mycolactone or solvent control, then incubated with N-WASP and placed in Jurkat T cell extracts supplemented with ATP, MgCl₂, and Alexa Fluor 488–labeled actin monomers for 2 hours. Representative beads are shown in phase-contrast and fluorescence. Mean fluorescence signal of more than 30 beads from 3 independent experiments are compared. ****P < 0.0001, unpaired 2-tailed t test, Welch corrected. (B) Jurkat cells were treated with a mycolactone fluorescent derivative (bodipy-Myc) for 1 hour, then processed for immunofluorescence with antibody binding the open form of WASP (active-WASP) and antibody recognizing p34-ARP2/3. Each image corresponds to a single confocal plane. 8 of 11 randomly picked cells had a Pearson coefficient greater than 0.5 (calculated on a z stack), indicative of Bodipy-mycolactone and active WASP colocalization. (C) Differential ARP2/3 complex recruitment in HeLa cells treated with vehicle (control), 20 nM mycolactone for 4 hours, or mycolactone plus 1 µM wiskostatin (Wisko). Representative immunofluorescence images and integrated p34-ARP2/3 intensities (n > 50) in the perinuclear region (see Methods) are shown. Perinuclear enrichment was calculated as mean ± SEM intensity and presented relative to control. *P < 0.05, ANOVA with Dunn post-test. (D) Western blot analysis of N-WASP, p34-ARP2/3, and actin expression in HeLa cells exposed to 20 nM mycolactone for 6 or 16 hours, compared with vehicle-treated control cells and untreated cells (Unt). GAPDH served as an internal control. Scale bars: 5 µm (A and B); 25 µm (C).
of ARP2/3-mediated actin assembly, but also its localization at the leading edge (29). The potential effect of mycolactone on HeLa cell displacement was therefore evaluated in a wound-healing assay. As shown in Figure 6A, mycolactone-exposed cells had an impaired capacity to close wounds. This was not due to decreased cell motility, as mycolactone-treated cells moved faster than vehicle-exposed controls (Figure 6B). However, the directionality of their movement was altered: cell trajectories were not linear and perpendicular to the wound (x-axis) like those of control cells, resulting in shorter displacements (Figure 6B). Prior transfection of the cells with WASP CR1 abrogated the mycolactone-induced alterations in speed and directionality and partially restored healing capacity (Figure 6, C and D), which suggests that mycolactone binding to endogenous N-WASP is critically involved in these defects.

**Hyperactivation of epithelial N-WASP underpins epidermal rupture.** The human epidermis is a stratified squamous epithelium composed of proliferating basal keratinocytes attached to a basement membrane and suprabasal keratinocyte layers at various stages of differentiation (from base to top: spinosum, granulosum, lucidum, and corneum). The maintenance of cell-cell contacts and the directionality of cell migration perpendicularly to the basement membrane have been shown to be critical for the maintenance of skin integrity (reviewed in ref. 30). Since mycolactone-induced activation of N-WASP affects both adhesion and direction sensing of epithelial cells, we investigated its effect on epidermal integrity in the mouse ear model, focusing on the events preceding tissue destruction and ulcer formation (Supplemental Figure 5).

As soon as 2 days after injection, we observed progressive thinning of the external stratum granulosum and lucidum (Figure 7, A and B), resulting in a significant decrease in epidermis width after 54 hours (Figure 7C). In agreement with our observation on cultured cells, remodeling of the epidermal layers was associated with a dramatic loss of E-cadherin–adhesive contacts, particularly in the stratum spinosum (Figure 7D). Importantly, coadministration of wiskostatin suppressed mycolactone-induced dissociation and thinning of the epidermis (Figure 7, D and E), demonstrating the importance of N-WASP activation in this process.

![Figure 4](image-url)

**Figure 4**
Effect of mycolactone on epithelial cell adhesion. (A) Phase-contrast images of a HeLa cell treated with 20 nM mycolactone for 0, 3, 8, and 24 hours. Original magnification, ×63. (B) Mean number of adherent HeLa cells after treatment with 20 nM mycolactone or vehicle control for up to 3 days. (C) Proportion of cells undergoing early and late apoptosis, in the adherent (A) and detached (D) fractions of HeLa cells treated with 20 nM mycolactone or solvent as control. Data are mean percentages on triplicates. (D) Adhesion of HeLa cells, as measured by calcein-AM assay, after 4 hours of treatment with vehicle control, 20 nM mycolactone, or mycolactone in the presence of 1 μM wiskostatin. ***P < 0.001, Kruskal Wallis with Dunn post-test. (E) Adhesion of HeLa cells transfected with an expression vector encoding WASP CR1 or with an empty vector (no WASP CR1) 24 hours prior to treatment with vehicle control or 10 nM mycolactone for 16 hours.
Discussion

Here, we showed that mycolactone mimicked endogenous regulators of WASP/N-WASP to hijack ARP2/3-mediated assembly of actin filaments in host cells. Numerous bacterial effectors, such as IcsA/VirG of *Shigella flexneri* (31) and TccP/EspFu of the enterohemorrhagic *Escherichia coli* (32), target WASP/N-WASP in host cells. However, mycolactone is the first example of a nonprotein factor interfering with these actin-nucleating factors. Unlike IcsA and EspFu, which require translocation into target cells, mycolactone gains access to WASPs by passive diffusion through the plasma membrane (8). Mycolactone bound N-WASP 100-fold more strongly than its major regulator, CDC42 (26), leading to a much greater capacity to stimulate actin assembly in vitro. The ARP2/3 complex has previously been shown to induce a slight shift in the autoinhibitory equilibrium of WASP, facilitating CDC42-induced opening and activation (26). We found that ARP2/3 comparably potentiated mycolactone-induced release of WASP CR1–bound VCA (data not shown). This property, combined with its high affinity for WASPs, should enable mycolactone to efficiently disconnect WASPs from endogenous regulation in vivo.

Mycolactone counteracted the effects of the N-WASP inhibitor wiskostatin in cellular assays. Consistent with the fact that wisko-
tone-induced activation of N-WASP operates preferentially in this region. Accordingly, the defects caused by mycolactone in migrating epithelial cells (e.g., increased speed of cell migration and loss of directionality) were comparable to those induced by delocalization of ARP2 expression from leading protrusions to the perinuclear area (34). The decreased capacity of mycolactone-exposed cells to extend cadherin-adhesive contacts was similar to that previously observed upon sequestration of ARP2/3 by exogenously expressing the CA region of N-WASP (35).

Our data in Figure 4 showed that mycolactone-induced killing of epithelial cells was primarily caused by N-WASP-mediated detachment and subsequent anoikis. This mechanism obviously

Figure 6
Mycolactone impairs directed migration of epithelial cells. (A) Representative image of the wound area after 24 hours of cell migration in the presence of 25 nM mycolactone or methanol (control). Dashed line denotes wound edges at 0 hours. (B) Mean ± SEM cell speed and x displacement (n > 170), and time-dependent representation of cell trajectories. ***P < 0.001, Mann-Whitney test. (C) Mean ± SEM speed and x displacement of more than 200 HeLa cells transfected with an expression vector encoding WASP CR1 24 hours before assessment of cell migration in the presence or absence of 25 nM mycolactone. Controls are cells transfected with empty vector and treated with vehicle. *P < 0.05, ***P < 0.001, Kruskal Wallis with Dunn post-test. (D) Representative images of the wound area after 24 hours of cell migration. HeLa cells were transfected with an expression vector encoding WASP CR1, or an empty vector (no WASP CR1) 24 hours prior to wound formation and cell migration in the presence of 25 nM mycolactone or vehicle control. Dashed lines denote wound edges at 0 hours. Wound-healing assays were repeated twice with similar results. Scale bars: 100 μm (A and D); 50 μm (B).

statin inhibits N-WASP activity by stabilizing the autoinhibited conformation, we found that mycolactone relieves the intramolecular contacts that maintain WASP autoinhibition. Intriguingly, mycolactone binding to WASPs involved the BR, which is located immediately upstream of the CRIB. Phosphatidylinositol 4,5-disphosphate (Pip2) micelles react with arginine and lysine residues clustered in this region to activate WASPs at both the allosteric level and the oligomerization level (33). NMR studies will help to determine whether mycolactone exploits this mechanism and to provide structural insight into the hyperactive conformation of the mycolactone–N-WASP complex.

In HeLa cells exposed to mycolactone, ARP2/3 was primarily detected in the perinuclear area, which suggests that mycolactone-induced activation of N-WASP operates preferentially in this region. Accordingly, the defects caused by mycolactone in migrating epithelial cells (e.g., increased speed of cell migration and loss of directionality) were comparable to those induced by delocalization of ARP2 expression from leading protrusions to the perinuclear area (34). The decreased capacity of mycolactone-exposed cells to extend cadherin-adhesive contacts was similar to that previously observed upon sequestration of ARP2/3 by exogenously expressing the CA region of N-WASP (35).
gene expression suggest that it may be involved in mycolactone inhibition of cytokine production by T cells (37). Our observation that mycolactone induced lamellipode formation in T cells (Figure 1A) strongly suggests that it may also alter the T cell spreading response to activation. Contrary to N-WASP, WASP is not essential and can be silenced without a major effect on viability. Study of animal models deficient for WASP expression or expressing WASP cannot explain mycolactone cytotoxicity in anchorage-independent cells, such as T lymphocytes (11). Activating mutations in WASP have been shown to increase apoptosis in T cells by promoting genomic instability, providing a possible mechanism for the mild cytopathic activity of mycolactone in this cell population (36). With regard to its immunomodulatory properties, recent studies indicating that WASP is an epigenetic regulator of Th1 gene expression suggest that it may be involved in mycolactone inhibition of cytokine production by T cells (37). Our observation that mycolactone induced lamellipode formation in T cells (Figure 1A) strongly suggests that it may also alter the T cell spreading response to activation. Contrary to N-WASP, WASP is not essential and can be silenced without a major effect on viability. Study of animal models deficient for WASP expression or expressing WASP.
mutants that are unable to bind mycolactone should help to dis- 
ssect the contribution of this key immune regulator to mycolactone 
activity in the hematopoietic cell compartment.

Currently available techniques do not allow for precise quanti- 
fication of mycolactone concentration in infected tissues (38, 39). 
We chose to inject 5 μg mycolactone into mouse ears, as within 2 
weeks, this dose reproduces the characteristic lesions develop- 
ing after months in mice infected with M. ulcerans (40). Consistent 
with our observation that mycolactone suppressed cell-cell con- 
tacts in polarized epithelia (Figure 5), our histopathological analy-
sis of mycolactone-injected skin revealed profound alterations in 
E-cadherin junctions and epidermal architecture, which suggests 
that uncontrolled activation of N-WASP impairs epithelial differ- 
entiation in this tissue. Since mycolactone-induced thinning of the skin was suppressed by wiskostatin, we propose that hyperacti-
vation of N-WASP (and consequent dissociation of epithelial cells) 
is the primary cause of epidermal destruction in Buruli ulcers.

Pathogens often modulate the host actin cytoskeleton to 
enhance their propagation and survival (41). M. ulcerans does not 
survive at the core temperature of the human body and multiplies 
extracellularly in disintegrated skin tissues. This environment, and 
the associated release of nutrients by dead cells, may constitute a 
favorable niche for the growth of M. ulcerans in an infected host. 
By targeting WASP and N-WASP, mycolactone has the potential to 
affect the functional biology of both immune and nonimmune 
cell populations. Further investigations will be required to deter-
mine whether WASP-mediated functions are altered by myocol- 
tone in hematopoietic cells, and to what extent these alterations contribute to the immune defects of M. ulcerans–infected hosts.

Our present results suggest that modulation of mycolactone-
induced N-WASP activation in the skin might counteract the 
ulcerative effects of mycolactone and represent a viable strategy to 
treat Buruli ulcers. This hypothesis was supported by preliminary results showing that local injection of wiskostatin limited the epi-
dermal remodeling preceding ulceration in M. ulcerans–infected mice (our unpublished observations). Resolving the structure of the mycolactone/WASP complex should identify the contact 
amino acids. Drugs mimicking the binding site of mycolactone in 
WASP may represent inhibitors of interest for treating this debili-
tating human disease (42).

**Methods**

**Reagents.** Mycolactone was purified from M. ulcerans 1615 (ATCC 35840) and quantified as described previously (4, 43). Stock solutions were prepared in methanol. Wiskostatin (Sigma-Aldrich) was in DMSO. Both reagents 
were diluted at least x10 in PBS before injection in animals and at least 
x1,000 in culture medium for cellular assays. In all cases, mycolactone- 
and wiskostatin-treated samples were compared with vehicle-treated controls.

**WASP and N-WASP constructs.** The CR1–CR7 constructs corresponding to WASP/N-WASP domains were provided by C. Egle (Institut Pasteur) or 
generated in our laboratory. The CR8–CR11 constructs were derived from 
M biotinylated mycolactone (or vehicle as control) for 15 min-
utes, rinsed, incubated with 400 nM N-WASP for 30 minutes (or buffer as 
control), rinsed, placed into Jurkat cell homogenates spiked with 1.5 
μM biotinylated mycolactone (or vehicle as control) for 15 min-
utes, rinsed, incubated with 400 nM N-WASP for 30 minutes (or buffer as 
control), rinsed, placed into Jurkat cell homogenates spiked with 1.5 μM 
G-actin (10% Alexa Fluor 488) plus 1.2 mM ATP and 2.4 mM MgCl2, and 
and 0.2 mM EGTA. When required, CDC42 was loaded with GTP-gS by a 
30-minute incubation at RT with PBS plus 1% BSA. Image analysis was performed with a custom-designed script developed using Acapela Image analysis software 
(version 2.5; Perkin Elmer). This script was based on nuclei segmentation, 
allowing the definition of a perinuclear ring of fixed thickness (6 pixels). For 
each nucleus (n > 50), the intensity of p34-ARP2/3 staining per unit of ring 
surface was calculated and integrated on 4 consecutive z sections (0.45 μm), 
then exported in a readable file format for statistical analysis.

**Actin polymerization assays.** Actin was purified from rabbit skeletal muscle (48), and the monomeric G form was purified by gel filtration and 
and labeled with pyrenyl iodoacetamide or Alexa Fluor 488 succinimidyl ester (Invitrogen). The ARP2/3 complex was purified from bovine brain (31).

The time course of actin polymerization was monitored spectrofluoro-
metrically in a Safas Xenius instrument using the increase in pyrenyl actin 
fluorescence as a probe, with excitation and emission wavelengths of 366 
and 407 nm, respectively. Polymerization was assayed at 2.5 μM G-actin 
(5% pyrenyl labeled), with 28 nm ARP2/3 complex and 32 nm N-WASP in 
the presence of mycolactone or solvent as control. The buffer contained 
5 mM Tris-Cl (pH 8), 0.2 mM ATP, 0.1 mM CaCl2, 0.1 M KCl, 1 mM MgCl2, 
and 0.2 mM EGTA. When required, CDC42 was loaded with GTP-gS by a 
30-minute incubation at RT with 1 mM GTP-gS in 5 mM EDTA. For actin 
polymerization on beads, streptavidin-latex 2-μm beads were incubated 
with 12.5 μM biotinylated mycolactone (or vehicle as control) for 15 min-
utes, rinsed, incubated with 400 nM N-WASP for 30 minutes (or buffer as 
control), rinsed, placed into Jurkat cell homogenates spiked with 1.5 μM 
G-actin (10% Alexa Fluor 488) plus 1.2 mM ATP and 2.4 mM MgCl2, and 
and analyzed for fluorescence after 2 hours at RT.

**Cell adhesion assays.** We used the Vybrant calcine-AM–based assay of cell 
adhesion (Molecular Probes). Briefly, HeLa cells grown in microplate wells 
were treated with mycolactone or vehicle, then detached and loaded with 
the cytoplasmic marker calcine-AM. Labeled cells were then left to adhere
to microplate wells for 2 hours, after which the nonadherent fraction was removed. The number of adherent cells was derived from the intensity of calcein fluorescence in each well.

Wound-healing assays. Wound-healing assays were performed with culture inserts and μ-slide 8 wells from Ibidi. Briefly, HeLa cells (70 μl, 5×10^5 cells/ml) were seeded in culture inserts and incubated for 24 hours at 37°C. After removal of culture inserts, cells were washed 3 times in x1 PBS and resuspended in 300 μl fresh DMEM/F-12 supplemented with 10% FCS and 2 mM l-glutamine. Cell migration was recorded for 24 hours at 37°C using a Nikon Eclipse Ti, capturing images at 10-minute intervals. Cell tracking was performed with Imaris (version 6.0; Bitplane).

Histology and immunocytochemistry. C57BL/6j mice were purchased from Jackson Laboratories and bred under specific pathogen–free conditions. Animals were injected intradermally in the center of the ear with the indicated reagents in a 50 μl PBS volume. At the indicated time points, mice were killed, and the ears were surgically removed and divided into 2 halves. One was fixed, dehydrated with absolute ethanol, plastic resin embedded (Historesin Leica Microsystems), and processed for H&E staining. The other was snap frozen in isopentane cooled with liquid nitrogen before 5-μm cryosectioning and immunostaining with rat anti-mouse E-cadherin ECCD-2 (Invitrogen) followed by secondary antibodies (Jackson ImmunoResearch). Nuclear DNA was visualized by Topro3 (Invitrogen).

Statistics. For each statistical comparison, the test used, n, and respective P value is provided in the corresponding figure legend.

Study approval. All experiments were performed with the guidelines of the National French Veterinary Department.

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