Bacteriophage adhering to mucus provide a non–host-derived immunity

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Mucosal surfaces are a main entry point for pathogens and the principal sites of defense against infection. Both bacteria and phage are associated with this mucus. Here we show that phage-to-bacteria ratios were increased, relative to the adjacent environment, on all mucosal surfaces sampled, ranging from cnidarians to humans. In vitro studies of tissue culture cells with and without surface mucus demonstrated that this increase in phage abundance is mucus dependent and protects the underlying epithelium from bacterial infection. Enrichment of phage in mucus occurs via binding interactions between mucin glycoproteins and Ig-like protein domains exposed on phage capsids. In particular, phage Ig-like domains bind variable glycan residues that coat the mucin glycoprotein component of mucus. Metagenomic analysis found these Ig-like proteins present in the phages sampled from many environments, particularly from locations adjacent to mucosal surfaces. Based on these observations, we present the bacteriophage adherence to mucus model that provides a ubiquitous, but non–host-derived, immunity applicable to mucosal surfaces. The model suggests that metazoan mucosal surfaces and phage co-evolve to maintain phage adherence. This benefits the metazoan host by limiting mucosal bacteria, and benefits the phage through more frequent interactions with bacterial hosts. The relationships shown here suggest a symbiotic relationship between phage and metazoan hosts that provides a previously unrecognized antimicrobial defense that actively protects mucosal surfaces.

symbiosis | host-pathogen | virus | immunoglobulin | immune system

Mucosal surfaces are the primary zones where animals meet their environment, and thus also the main points of entry for pathogenic microorganisms. The mucus layer is heavily colonized by bacteria, including many symbionts that contribute additional genetic and metabolic potential to the host (1, 2). Bacterial symbionts associated with a variety of other host surfaces also provide goods and services, e.g., nutrients (3–6), bio-luminescence (7, 8), and antibiotics (9, 10). These resident symbionts benefit from increased nutrient availability (5, 11–13), as well as the opportunity for both vertical transmission and increased dissemination (14–16).

Within the mucus, the predominant macromolecules are the large (up to $10^9$ Da) mucin glycoproteins. The amino acid backbone of these proteins incorporates tandem repeats of exposed hydrophobic regions alternating with blocks bearing extensive O-linked glycosylation (17). Hundreds of variable, branched, negatively charged glycan chains extend 0.5–5 nm from the peptide core outward into the surrounding environment (17, 18). Other proteins, DNA, and cellular debris also are present. Continual secretion and shedding of mucins maintain a protective mucus layer from 10–70 μm thick depending on species and body location (19–22).

By offering both structure and nutrients, mucus layers commonly support higher bacterial concentrations than the surrounding environment (11, 23). Of necessity, hosts use a variety of mechanisms to limit microbial colonization (24–27). Secretions produced by the underlying epithelium influence the composition of this microflora (12, 27, 28). When invaded by pathogens, the epithelium may respond by increased production of antimicrobial agents, hypersecretion of mucin, or alteration of mucin glycosylation patterns to subvert microbial attachment (29–31).

Also present in the mucus environment are bacteriophage (phage), the most common and diverse biological entities. As specific bacterial predators, they increase microbial diversity through Red Queen/kill-the-winner dynamics (32, 33). Many phages establish conditional symbiotic relationships with their bacterial hosts through lyogamy. As integrated prophages, they often express genes that increase host fitness or virulence (34–36) and protect their host from lysis by related phages. As free phage, they aid their host strain by killing related competing strains (37–39). Phages participate, along with their bacterial hosts, in tripartite symbioses with metazoans that affect metazoan fitness (40–43). However, no direct symbiotic interactions between phage and metazoans are known.

Recently, Minot et al. (44) showed that phages in the human gut encode a population of hypervariable proteins. For 29 hypervariable regions, evidence indicated that hypervariability was conferred by targeted mutagenesis through a reverse transcription mechanism (44, 45). Approximately half of these encoded proteins possessed the C-type lectin fold previously found in the major tropism determinant protein at the tip of the Bordetella phage BPP-1 tail fibers (46); six others contained Ig-like domains. These Ig-like proteins, similar to antibodies and T-cell receptors, can accommodate large sequence variation (>1013 potential alternatives) (47). Ig-like domains also are displayed in the structural proteins of many phage (48, 49). That most of these displayed Ig-like domains are dispensable for phage growth in the laboratory (45, 49) led to the hypothesis that they aid adsorption to their bacterial prey under environmental conditions (49). The possible role and function of these hypervariable proteins remain to be clarified.

Here, we show that phage adhere to mucus and that this association reduces microbial colonization and pathology. In vitro studies demonstrated that this adherence was mediated by the interaction between displayed Ig-like domains of phage capsid proteins and glycan residues, such as those in mucin glycoproteins. Homologs of these Ig-like domains are encoded by phages from many environments, particularly those adjacent to mucosal surfaces. We propose the bacteriophage adherence to mucus (BAM) model whereby phages provide a non–host-derived antimicrobial defense on the mucosal surfaces of diverse metazoan hosts.
Results
Phage Adhere to Mucus. Our preliminary investigations of mucosal surfaces suggested that phage concentrations in the mucus layer were elevated compared with the surrounding environment. Here, we used epifluorescence microscopy to count the phage and bacteria in mucus sampled from a diverse range of mucosal surfaces (e.g., sea anemones, fish, human gum), and in each adjacent environment (SI Materials and Methods and Fig. S1). Comparing the calculated phage-to-bacteria ratios (PBRs) showed that PBRs in metazoan-associated mucus layers were on average 4.4-fold higher than those in the respective adjacent environment (Fig. 1A). The PBRs on these mucus surfaces ranged from 21:1 to 87:1 (average, 39:1), compared with 3:1 to 20:1 for the surrounding milieus (average, 9:1; n = 9, t = 4.719; ****P < 0.0002). Earlier investigations of phage abundance in marine environments reported that phage typically outnumber bacteria by an order of magnitude (50–52), but here we demonstrate that this margin was significantly larger in metazoan-associated mucus surface layers.

To determine whether this enrichment was dependent on the presence of mucus rather than some general properties of the cell surface (e.g., charge), phage adherence was tested with tissue culture (TC) cells with and without surface mucus (SI Materials and Methods). In these assays, T4 phage were washed across confluent cell monolayers for 30 min, after which nonadherent phage were removed by repeated washings and the adherent phage quantified by epifluorescence microscopy. Two mucus-producing cell lines were used: T84 (human colon epithelial cells) and A549 (human lung epithelial cells). For these cells, mucin secretion was stimulated by pretreatment with phorbol 12-myristate 13-acetate (53, 54). Comparison of the T84 cells with the non–mucus-producing Huh-7 human hepatocyte cell line showed that T4 phage adhered significantly more to the mucus-producing T84 cells (Fig. 1B; n = 18, t = 8.366; ****P < 0.0001). To demonstrate the mucus dependence of this adherence, the mucus layer was chemically removed from A549 cells (Huh-7; MUC–) and these cells were overlaid with T4 phage for 30 min, washed to remove nonadherent phage, and then incubated with E. coli for 4 h. Cells then were scraped from the plates and the attached bacteria were fluorescently stained and counted by epifluorescence microscopy. Phage pretreatment of mucus-producing TC cell lines (T84, A549) significantly decreased subsequent bacterial attachment (Fig. 2A; T84: n > 30, t = 32.05, ****P < 0.0001; A549: n = 30, t = 36.85, ****P < 0.0001). Phage pretreatment of non–mucus-producing cells (Huh-7; MUC–), an A549 mucus knockdown strain) had a much smaller effect on bacterial attachment.

To determine whether this reduced bacterial attachment depended on bacterial lysis and the production of progeny viruses, we repeated these experiments using an amber mutant T4 phage (T4 am4344f). When infecting wild-type E. coli, this phage produces no infective progeny virions, but infection of the E. coli amber suppressor strain SupD yields infective virions. For these experiments, mucus-producing A549 cells were pretreated with amber mutant T4 am4344 phage and then incubated with either wild-type or the amber suppressor strain E. coli (Fig. 2B). Bacterial attachment was reduced by more than four orders of magnitude when phage could replicate and thereby increase the number of infective virions within the mucus (n = 8, ****P < 0.0001). Comparatively, when no phage replication occurred in the mucus, there was no observable change in bacterial colonization and fewer phages were detected (n = 8, *P = 0.0227). These results show that pretreatment of a mucosal surface with phage reduces adherence of a bacterial pathogen and that this protection is mediated by continued phage replication in the mucus.

To test whether the observed reduced bacterial adherence was accompanied by reduced pathology of the underlying TC cells, mucus-producing A549 and non–mucus-producing MUC– TC cells were exposed to E. coli overnight, either with or without a 30-min pretreatment with T4 phage. Infection was quantified as the percentage of cell death. Adherence of phage effectively protected the mucus-producing cells against the subsequent bacterial challenge (Fig. 2C; n = 12, ***P < 0.0001). Phage pretreatment showed a reduced protection to the non–mucus-producing MUC–...
cells, decreasing cell death only twofold. Evaluating the importance of mucus production for effective protection, we found that phage pretreatment of mucus-producing A549 cells resulted in a 3.6-fold greater reduction in cell death (n = 12, *P = 0.0181) than the same pretreatment of the mucus knockdown MUC- cells.

**Role of Capsid Ig-Like Domains in Phage Adherence.** Minot et al. (44) recently reported that phage communities associated with the human gut encode a diverse array of hypervariable proteins, including some with hypervariable Ig-like domains. Four Ig-like domains are found in highly antigenic outer capsid protein (Hoc), a T4 phage structural protein of which 155 copies are displayed on the surface of the major capsid protein (49). Results suggested that most Ig-like domains function in recognition and adhesion (Hoc), a T4 phage structural protein of which 155 copies are displayed on the surface of the major capsid protein (49). Results indicated a similar increase in adherence to mucin for the Igβ, but not the Igβ, T3 phage (Fig. S6). Thus, adherence of these phage to mucus requires the Ig-like protein domains.

Second, a competition assay using hoc+ and hoc− T4 phage and mucus-producing TC cells was performed to demonstrate the role of mucin in phage adherence. Phage suspended in mucin solutions ranging from 0% to 5% (wt/vol) were washed over confluent layers of mucus-producing A549 TC cells; phage adherence then was assayed as above. Adherence of hoc+ T4 phage, but not of hoc− T4 phage, was reduced by mucin competition in a concentration-dependent manner (Fig. 3B).

Third, interaction of the Hoc protein domains displayed on the capsid surface with mucin glycoproteins was hypothesized to affect the rate of diffusion of T4 virions in mucin. To evaluate this, we used multiple-particle tracking (MPT) to quantify transport rates of phage particles in buffer and in mucin suspension. The ensemble average effective diffusivity (Deff) calculated at a time scale of 1 s for both hoc+ and hoc− T4 phage in buffer was compared against that in 1% (wt/vol) mucin suspensions (SI Materials and Methods). Both hoc+ and hoc− phage diffuse rapidly through buffer (Fig. 3C). Whereas hoc+ phage diffused in 1% mucin at the same rate as in buffer, the mucin decreased the diffusion rate for hoc− phage particles eightfold. Thus, all three of these experimental approaches supported our hypothesis that...

**Fig. 2.** Effect of phage adsorption on subsequent bacterial infection of epithelial cells. (A) Bacterial attachment to mucus-producing (T84 and A549) and non-mucus-producing (Huh-7, MUC) TC cells, with and without phage pretreatment. T4 phage pretreatment significantly decreased subsequent bacterial adherence to mucus-producing TC cell lines (T84: n > 30, t < 32.05, ****P < 0.0001; A549: n > 30, t = 36.85, ****P < 0.0001; unpaired t tests). Less dramatic shifts were seen for non-mucus-producing cells (Huh-7: n > 30, t < 2.72, **P = 0.0098; MUC: n > 30, t > 3.52, ***P = 0.0007; unpaired t tests). (B) Mucus-producing A549 cells were pretreated with T4 am4344 phage (Materials and Methods) and then incubated for 4 h with either wild-type (wt) or amber-suppressor (SupD) E. coli. Phage replication in the SupD E. coli strain significantly reduced bacterial colony-forming units (CFU) in the mucus (n = 8, ****P < 0.0001, Tukey’s two-way ANOVA) and increased phage-forming units (PFU) relative to the no-phage replication wt. E. coli (n = 8, *P = 0.0227). (C) Mortality of mucin-producing (A549) and mucin knockdown (MUC) A549 lung epithelial cells following overnight incubation with E. coli. Phage pretreatment completely protected mucus-producing A549 cells from bacterial challenge (n = 12, ****P < 0.0001, Tukey’s one-way ANOVA); protection of MUC− cells was 3.1-fold less (n = 12, *P = 0.0181). ns, not significant.

**Fig. 3.** Effect of Hoc protein on phage–mucin interactions. (A) Adherence of hoc+ and hoc− T4 phage to agar coated with mucin, DNA, or protein reported as an increase relative to plain agar controls (n > 11, t = 3.977, ***P = 0.0007, unpaired t test). (B) Competitive effect of mucin on phage adherence when hoc+ and hoc− T4 phage in 0.5% (wt/vol) mucin solution (1× PBS) were washed over mucus-producing A549 cells (n = 25 per sample). (C) Diffusion of fluorescence-labeled hoc+ (Left) and hoc− (Right) T4 phage in buffer and 1% mucin as determined by MPT. Mucin hindered diffusion of hoc+ T4 phage but not hoc− phage (10 analyses per sample, trajectories of n > 100 particles for each analysis; error bars represent SE).
the Hoc proteins displayed on the T4 phage capsid interact with mucin.

Phage Capsid Ig-Like Domains Interact with Glycans. It is known that ~25% of sequenced tailed dsDNA phages (Caudovirales) encode structural proteins with predicted Ig-like domains (48). A search of publicly available viral metagenomes for homologs of the Ig-like domains of the T4 Hoc protein yielded numerous viral Ig-like domains from a variety of environments (Fig. 4A). These domains were more likely to be found in samples collected directly from mucus (e.g., sputum samples) or from an environment adjacent to a mucosal surface (e.g., intestinal lumen, oral cavity). All homologs displayed high structural homology (Phyre2 confidence score average, 96 ± 5%) with a plant-sugar binding site. A unique example of a meta-
hoc phage on mucosal surfaces is mediated by weak binding interaction between these Ig-like domains and glycans.

Mucins are complex glycoproteins with highly variable glycan groups exposed to the environment. To investigate whether Hoc interacts with glycans and, if so, to determine whether it interacts with a specific glycan or with a diverse array of glycans, we assayed phage adherence to microarrays printed with 610 mammalian glycans. The hoc+ T4 phage adhered to many diverse glycans and showed a preference for the O-linked glycan residues typically found in mucin glycoproteins (Fig. 4B, SI Materials and Methods, and Table S1). The hoc+ T4 phage exhibited significantly lower affinity for all tested glycans. This indicates that Hoc mediates interactions between T4 phage and varied glycan residues.

Discussion

In diverse metazoans, body surfaces that interact with the environment are covered by a protective layer of mucus. Because these mucus layers provide favorable habitats for bacteria, they serve as the point of entry for many pathogens and support large populations of microbial symbionts. Also present are diverse phages that prey on specific bacterial hosts. Moreover, phage concentrations in mucus are elevated relative to the surrounding environment (an average 4.4-fold increase for a diverse sample of invertebrate and vertebrate metazoans; Fig. 4A). The increased concentration of lytic phage on mucosal surfaces provides a previously unrecognized metazoan immune defense affected by phage lysis of incoming bacteria.

Working with a model system using T4 phage and various TC cell lines, we demonstrated that the increased concentration of phage on mucosal surfaces is mediated by weak binding interactions between the variable Ig-like domains on the T4 phage capsid and mucin-displayed glycans. The Ig protein fold is well known for its varied but essential roles in the vertebrate immune response and cell adhesion. Ig-like domains also are present in approximately one quarter of the sequenced genomes of tailed DNA phages, the Caudovirales (48). Notably, these domains were found only in virion structural proteins and typically are displayed on the virion surface. Thus, they were postulated to bind to bacterial surface carbohydrates during infection (48, 49). However, mucin glycoproteins, the predominant macromolecular constituent of mucus, display hundreds of variable glycan chains that are of interest to researchers in the field of phage therapy.

The presence of an Ig-like protein (Hoc) displayed on the capsid of T4 phage significantly slowed the diffusion of the phage in mucin solutions. In vivo, similar phage binding to mucin glycans would increase phage residence time in mucus layers. Because bacterial concentrations typically are enriched in mucin (Fig. S1), we predict that mucus-adherent phage are more likely to encounter bacteria, potentially increasing their replicative success. If so, phage Ig-like domains that bind effectively to the mucus layer would be under positive selection. Likely, Hoc and other phage Ig-like domains interact with other glycans with different compositions, as well (49, 58).

Previous metagenomic studies documented the ubiquity and diversity of bacteria and phage within mucus-associated environments (e.g., human gut, human respiratory tract, corals) (52, 59–64). Known also were some of the essential but adaptable services provided by symbiotic bacteria in these environments (65). However, only recently have efforts been made to investigate the dynamic influences of phage within host-associated ecosystems (37, 44, 66). In this work, we used an in vitro model system to demonstrate a mechanism of phage adherence to the mucus layers that shield metazoan cells from the environment. Furthermore, adherent phage protected the underlying epithelial cells from bacterial infection. Based on these observations and previous research, we proposed the BAM model of immunity, in which the adherence of phage to mucosal surfaces yields a non-host-derived, antimicrobial defense. According to this model (summarized in Fig. 5), the mucus layer, already considered part of the innate immune system and known to provide physical and biochemical antimicrobial defenses (18, 27, 67), also accumulates phage.

The model system we used involved a single lytic phage and host bacterium; the situation in vivo undoubtedly is more complex. Within the mucosal layer reside diverse bacterial lineages and predictably an even greater diversity of phage strains, both enmeshed within complex phage–bacterial infection networks and engaged in a dynamic arms race (68, 69). These and other factors lower the probability that any given phage–bacterium encounter will result in a successful infection. The time dimension adds further complexity. The mucus layer is dynamic. Mucins are secreted continually by the underlying epithelium while mucus is sloughed continually from the outer surface. As a result, there is an ongoing turnover of both the bacterial and phage populations in the mucus layer. Driven by kill-the-winner dynamics, the population of phage types that can infect the dominant bacterial types present will cycle along with the populations of their hosts. Through such mechanisms, we envision that adherent lytic phages provide a dynamic and adaptable defense for their metazoan hosts—a unique example of a metazoan–phage symbiosis.

We posit that BAM immunity reduces bacterial pathogenesis and provides a previously unrecognized, mucosal immunity. This has far-reaching implications for numerous fields, such as human immunity, gastroenterology, coral disease, and phage therapy. Meanwhile, key questions remain. For instance, what role do temperate phages play in the dynamics of BAM immunity? When integrated into the bacterial chromosome as prophages,
they protect their bacterial hosts from infection by related phages: as free phages, they infect and kill sensitive related bacterial strains that compete with their bacterial hosts (37–39). Both mechanisms may benefit their metazoan host by contributing to the maintenance of a selected commensal mucosal microbiota. These possibilities remain to be investigated. Likewise, in vivo investigations are needed to characterize the bacterial and phage diversity present and the consequent effects on BAM immunity. As of now, the relationships shown here open an arena for immunological study, introduce a phage–metazoan symbiosis, and recognize the key role of the world’s most abundant biological entities in the metazoan immune system.

Materials and Methods

Bacterial Strains, Phage Stocks, TC Cell Lines, and Growth Conditions. *E. coli* 1024 strain was used for all *E. coli* experiments and was grown in LB (10 g tryptone, 5 g yeast extract, 10 g NaCl, in 1 L dH2O) at 37 °C overnight. *E. coli* amber-suppressor strain SupD strain CR63 was used as a host for amber mutant phage and grown as above. Bacteriophage T4 was used at ~107 pfu/mL. Hoc T4 phage were kindly supplied by Prof. Venigalla Rao (50), The Catholic University of America, Washington, D.C. T3 am10 Ig− amber mutant phage were kindly supplied by Prof. Ian J. Molineux (70), University of Texas, Austin, TX. T4 replication-negative 43− (amE4332; DNA polymerase) 44− (amN82; subunit of polymerase clamp holder) amber mutant phage were kindly supplied by Prof. Kenneth Kreuzer (71), Duke University School of Medicine, Durham, NC. The human tumorigenic colon epithelial cell line, T84, were obtained from the American Type Culture Collection (ATCC) and cultured in DMEM/F12-K media with 5% FBS and 100 μg/mL penicillin-streptomycin (PS). The human tumorigenic lung epithelial cell line A549 was kindly supplied by Prof. Kelly Doran, San Diego State University, San Diego, CA and cultured in F12-K media with 10% FBS, 100 μg/mL PS. All TC cell lines initially were grown in 50 mL Primaria Tissue Culture Flasks (Becton Dickinson) at 37 °C and 5% CO2.

Phage Adherence to Mucus-Associated Macromolecules. LB agar plates were coated with 1 mL of 1% (wt/vol) of one of the following in 1× PBS: type III porcine stomach mucin, DNA from salmon testes, or BSA (all three from Sigma–Aldrich) and then allowed to dry. Stacks of hoc+ and hoc− T4 phage (107 pfu/mL−1) were serially diluted to 1 × 10−1 and 1 × 10−5 per milliliter in LB, and a 5-mL aliquot of each dilution was washed across the plates for 30 min at 37 °C on an orbital shaker. After the phage suspensions were decanted from the plates, the plates were shaken twice to remove excess liquid and dried. Each plate then was layered with 1 mL of overnight *E. coli* culture (106 mL−1) in 3 mL of molten top agar and incubated overnight at 37 °C. The number of adherent phage was calculated from the number of plaque-forming units observed multiplied by the initial phage dilution. To determine whether mucus macromolecules directly affected phage infectivity, hoc+ and hoc− T4 phage (105 pfu/mL−1) were serially diluted as described above into 1 mL LB solutions containing 1% (wt/vol) mucin, DNA, or BSA. After incubation for 30 min at 37 °C, the phage suspensions were combined with *E. coli* top agar as described above and layered over uncoated agar plates (Fig. S5).

Phage Treatment of TC Cells. TC cells were washed twice with 5 mL of serum-free media to remove residual antibiotics, layered with 2 mL of serum-free media containing T4 phage (105−106 mL−1), and incubated at 37 °C and 5% CO2 for 30 min. Cells then were washed five times with 5 mL of serum-free media to remove nonadherent phage.

Phage Adherence to TC Cells. TC cells were treated with phage (106 mL−1; see above), then scraped from plates using Corning Cell Scrappers (Sigma–Aldrich). Adherent phage were counted by epifluorescence microscopy as described above.

Bacterial Adherence to TC Cells With/Without Phage Pretreatment. TC cells with or without pretreatment with T4 phage (106 mL−1) were layered with 2 mL serum-free media containing *E. coli* (105 mL−1), incubated at 37 °C and 5% CO2 for 4 h, and then washed five times with 5 mL serum-free media to remove nonadherent phage and bacteria. Cells were scraped from plates, and adherent phage and bacteria were counted by either epifluorescence microscopy, as described above, or colony-forming and plaque-forming units. Then, 100 μL of a relevant dilution was spread onto an agar plate and incubated overnight at 37 °C, and the number of adherent bacteria was calculated from the colony-forming units observed multiplied by the initial dilution. Plaque-forming units were counted by a top agar assay as described above.

TC Cell Death from Bacterial Infection. Mucus-producing A549 and MUC5AC A549 TC cells were grown to confluence. T4 phage were cleaned using Amicon 50-kDa centrifugal filters (Millipore) and saline magnesium buffer (SM) to remove bacterial lysis products. Cells, with or without T4 phage pretreatment (106 mL−1), were incubated with *E. coli* (106 mL−1) overnight. Afterward, TC cells were recovered from the plates by trypsin/EDTA solution (Invitrogen). Cells were pelleted by centrifugation and resuspended in 1× PBS. Dead cells were identified by staining with 1 mg/mL of propidium iodide (Invitrogen). Samples then were analyzed on a FACSCanto II flow cytometer (BD Biosciences) with excitation at 488 nm and emission detected through a 670 long pass filter. The forward scatter threshold was set at 5,000, and a total of 10,000 events were collected for each sample.

Mucin Competition Assay. Mucus-producing A549 TC cells were grown to confluence. Hoc+ and hoc− T4 phage (105 mL−1) were diluted into mucin solutions ranging between 0% and 5% (wt/vol) in 1× PBS then washed over TC cells for 30 min at 37 °C and 5% CO2. Cells were washed five times with 5 mL serum-free media to remove nonadherent phage, scraped from plates, and adherent phage were quantified as described above.

Graphing and Statistics. Graphing and statistical analyses were performed using GraphPad Prism 6 (GraphPad Software). All error bars represent 5–95% confidence intervals. The midline represents the median and the mean for box plots and bar plots, respectively.
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