

# Long-term antibiotic exposure promotes mortality after systemic fungal infection by driving lymphocyte dysfunction and systemic escape of commensal bacteria

Drummond, Rebecca A; Desai, Jigar V; Ricotta, Emily E; Swamydas, Muthulekha; Deming, Clay; Conlan, Sean; Quinones, Mariam; Matei-Rascu, Veronika; Sherif, Lozan; Lecky, David; Lee, Chyi-Chia R; Green, Nathaniel M; Collins, Nicholas; Zelazny, Adrian M; Prevots, D Rebecca; Bending, David; Withers, David; Belkaid, Yasmine; Segre, Julia A; Lionakis, Michail S

DOI:

[10.1016/j.chom.2022.04.013](https://doi.org/10.1016/j.chom.2022.04.013)

License:

Creative Commons: Attribution-NonCommercial-NoDerivs (CC BY-NC-ND)

*Document Version*

Peer reviewed version

*Citation for published version (Harvard):*

Drummond, RA, Desai, JV, Ricotta, EE, Swamydas, M, Deming, C, Conlan, S, Quinones, M, Matei-Rascu, V, Sherif, L, Lecky, D, Lee, C-CR, Green, NM, Collins, N, Zelazny, AM, Prevots, DR, Bending, D, Withers, D, Belkaid, Y, Segre, JA & Lionakis, MS 2022, 'Long-term antibiotic exposure promotes mortality after systemic fungal infection by driving lymphocyte dysfunction and systemic escape of commensal bacteria', *Cell Host & Microbe*, vol. 30, no. 7, pp. 1020-1033.e6. <https://doi.org/10.1016/j.chom.2022.04.013>

[Link to publication on Research at Birmingham portal](#)

## General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

## Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.

Download date: 09. Nov. 2024

# Long-term Antibiotics Promote Mortality After Systemic Fungal Infection by Driving Lymphocyte Dysfunction and Systemic Escape of Commensal Bacteria

Rebecca A. Drummond<sup>1,2†</sup>, Jigar V. Desai<sup>1†</sup>, Emily E. Ricotta<sup>1</sup>, Muthulekha Swamydas<sup>1</sup>, Clay Deming<sup>3</sup>, Sean Conlan<sup>3</sup>, Mariam Quinones<sup>4</sup>, Veronika Matei-Rascu<sup>2</sup>, Lozan Sherif<sup>2</sup>, David Lecky<sup>2</sup>, Chyi-Chia R. Lee<sup>5</sup>, Nathaniel M. Green<sup>1</sup>, Nicholas Collins<sup>6</sup>, Adrian M. Zelazny<sup>7</sup>, D. Rebecca Prevots<sup>1</sup>, David Bending<sup>2</sup>, David Withers<sup>2</sup>, Yasmine Belkaid<sup>6,8</sup>, Julia A. Segre<sup>3</sup>, Michail S. Lionakis<sup>1\*&</sup>

<sup>1</sup>Fungal Pathogenesis Section, Laboratory of Clinical Immunology and Microbiology (LCIM), National Institute of Allergy & Infectious Diseases (NIAID), National Institutes of Health (NIH), Bethesda, MD, 20892, USA

<sup>2</sup>Institute of Immunology & Immunotherapy, Institute of Microbiology & Infection, University of Birmingham, Birmingham B15 2TT, UK

<sup>3</sup>Microbial Genomics Section, Translational and Functional Genomics Branch, National Human Genome Research Institute, NIH, Bethesda, MD, 20892, USA

<sup>4</sup>Bioinformatics and Computational Bioscience Branch, NIAID, NIH, Bethesda, MD, 20892, USA

<sup>5</sup>Laboratory of Pathology, Center for Cancer Research, National Cancer Institute, NIH, Bethesda, MD, 20892, USA

<sup>6</sup>Metaorganism Immunity Section, Laboratory of Host Immunity and Microbiome, NIAID, NIH, Bethesda, MD, 20892, USA

<sup>7</sup>Department of Laboratory Medicine, NIH Clinical Center, NIH, Bethesda, MD, 20892, USA

<sup>8</sup>NIAID Microbiome Program, NIAID, NIH, Bethesda, MD, 20892, USA

†These authors contributed equally

\*Correspondence: [r.drummond@bham.ac.uk](mailto:r.drummond@bham.ac.uk); [lionakism@mail.nih.gov](mailto:lionakism@mail.nih.gov)

&Lead contact

## Summary

Antibiotics are a modifiable iatrogenic risk factor for the most common human nosocomial fungal infection, invasive candidiasis, yet the underlying mechanisms remain elusive. We found that antibiotics enhanced susceptibility to murine invasive candidiasis due to impaired lymphocyte-dependent IL-17A- and GM-CSF-mediated antifungal immunity within the gut, which led to non-inflammatory bacterial escape and systemic bacterial co-infection and could be ameliorated by IL-17A or GM-CSF immunotherapy. Vancomycin alone similarly enhanced susceptibility to invasive fungal infection and systemic bacterial co-infection. Mechanistically, vancomycin reduced gut Th17 cells associated with impaired proliferation and ROR $\gamma$ t expression. Vancomycin's effects on Th17 cells were indirect manifesting only *in vivo* in the presence of dysbiosis. In humans, antibiotics were associated with increased risk of invasive candidiasis and death after invasive candidiasis. Our work highlights the importance of antibiotic stewardship in protecting vulnerable patients from life-threatening infections and provides mechanistic insights into a controllable iatrogenic risk factor for invasive candidiasis.

## Introduction

Antibiotics can influence and modulate the immune system, leading to abrogated antimicrobial immunity and altered homeostasis. For example, antibiotic treatment skews the intestinal immune system towards a more regulatory environment, dampening the development of gut Th17 cells by eliminating the commensal bacteria needed to develop these cell types (Ivanov et al., 2008). Recent studies have also shown that antibiotics can exert direct effects on immune cells and may drive phenotypes that extend beyond the gastrointestinal (GI) tract (Erny et al., 2015). Bactericidal antibiotics can also directly disrupt metabolic processes in myeloid cells which limits their ability to kill bacterial pathogens (Kalghatgi et al., 2013; Yang et al., 2017), while a broad-spectrum antibiotic cocktail prevented protective systemic anti-bacterial immunity *in vivo* by disrupting granulopoiesis in a neonatal sepsis model (Deshmukh et al., 2014). Antibiotics can therefore cause immunological injuries that enhance susceptibility to secondary infections. Identifying which antibiotics carry these potentially harmful side-effects, and the mechanisms underlying these relationships, will be important for informing future antibiotic stewardship strategies and designing immune-based therapies that may circumvent the undesirable effects of antibiotics on the host.

Candidiasis is most often caused by *Candida albicans*, a common human fungal commensal in the GI tract, or, less frequently, the skin (Fan et al., 2015; Findley et al., 2013). Invasive candidiasis originates from these commensal populations following breach of GI or cutaneous barrier integrity (Lionakis, 2014). GI-derived invasive candidiasis is typically seen in patients with hematological malignancies during chemotherapy-induced mucositis and neutropenia (Lionakis, 2014). In these cases, broad-spectrum antibiotic pre-exposure is a recognized risk factor, operating to expand gut commensal *C. albicans* populations which then translocate into the bloodstream following barrier integrity disruption and phagocyte depletion (Fan et al., 2015; Koh et al., 2008; Odds et al., 2006; Zhai et al., 2020). Prophylactic fluconazole has dramatically decreased the prevalence of GI-derived invasive candidiasis in these patients (Lionakis, 2014; Pappas et al., 2018). Skin-derived invasive candidiasis has emerged as the most common form of invasive candidiasis in modern hospitals (Ricotta et al., 2020; Strollo et al., 2016). It is typically seen in acutely ill non-neutropenic patients in the intensive care unit, associated with central venous catheter-related breach of the cutaneous barrier. Notably, broad-spectrum antibiotic pre-exposure has also been documented as a risk factor for this form of invasive candidiasis (Keighley et al., 2019; Pappas et al., 2018). However, the mechanisms by which broad-spectrum antibiotics predispose to infection remain unclear as these patients do not typically feature impaired GI barrier integrity to account for fungal translocation.

In this study, we used animal models to analyze how broad-spectrum antibiotics affect systemic immune responses to *Candida albicans*. We demonstrate that pre-exposure to antibiotics, especially vancomycin, disrupt antifungal immunity in an organ-specific manner and that these effects can be ameliorated by targeted immunotherapy. We also analyzed the relationship between broad-spectrum antibiotic pre-exposure and development of invasive candidiasis in humans using a large health record database.

## Results

### **Broad-spectrum Antibiotics Result in GI Tract-Specific Susceptibility to Invasive Fungal Infection**

To determine if antibiotics affect susceptibility to invasive candidiasis, we investigated antibiotic-induced susceptibility to skin-derived invasive candidiasis in mice. Mice are not naturally colonized with *C. albicans* in their GI tract (Fan et al., 2015; Iliev et al., 2012), and therefore are a suitable model system to evaluate how antibiotics may promote invasive candidiasis and/or perturb antifungal immunity in this setting. Mice treated with a broad-spectrum antibiotic cocktail (ampicillin, metronidazole, neomycin, vancomycin [AMNV]) had significantly higher mortality when infected intravenously with *C. albicans* (Fig. 1A), an infection route that mimics human skin-derived candidiasis (Lionakis et al., 2011). Susceptibility was not associated with reduced circulating neutrophils, the critical mediators of defense against invasive candidiasis (Lionakis and Levitz, 2017) (Fig. S1).

Mortality in the mouse model of skin-derived invasive candidiasis is driven by uncontrolled renal fungal growth that causes kidney failure (Lionakis et al., 2011). Unexpectedly, the heightened mortality of antibiotic-treated animals was not due to increased renal fungal proliferation or kidney damage (Fig. 1B-1C), or impaired fungal control in the liver, spleen or brain (Fig. S2). Instead, we found significantly increased fungal growth across GI tissue compartments in antibiotic pre-exposed mice (Fig. 1D). We asked whether the increased GI fungal burden of antibiotic pre-exposed mice was apparent throughout superficial and deeper gut tissue layers or whether it was only seen in the gut lumen reflective of mouse coprophagic behavior post-infection (Fig. S3). We found that antibiotic pre-exposed mice had significantly higher fungal burden in all gut tissue layers tested (mucus, intestinal epithelium, deep intestinal tissue), consistent with hematogenously-derived fungal tissue invasion (Fig. 1E). Together, this data shows that broad-spectrum antibiotics promoted GI tract-specific susceptibility to invasive candidiasis in mice.

### **Commensal Gut Bacteria Disseminate in Antibiotic Pre-exposed Fungal-Infected Mice**

Since antibiotic-treated mice had increased GI tissue fungal proliferation, we wondered whether antibiotic pre-exposure compromised gut barrier function. Antibiotics were previously linked to low-level systemic translocation of commensal bacteria (Chakraborty et al., 2018; Knoop et al., 2016; Spiller, 2018; Tulstrup et al., 2015). Strikingly, we cultured large amounts of Gram-positive and Gram-negative bacteria from the spleen of *C. albicans*-infected antibiotic pre-exposed mice (Fig. 2A). No bacteria were recovered from the spleen of fungus-uninfected antibiotic pre-exposed mice or *C. albicans* infected mice that did not receive antibiotics, indicating that both antibiotics and systemic fungal challenge are required for systemic bacterial translocation in this model (Fig. 2A). In agreement, we found bacterial

translocation in the blood and peritoneal fluid of *C. albicans*-infected antibiotic pre-exposed mice (Fig. 2B). MALDI-TOF-based identification of these bacteria confirmed that they were GI-derived commensals (Fig. 2C). Histological gut examination indicated that the escape of commensal bacteria in *C. albicans*-infected antibiotic pre-exposed mice was non-inflammatory (Fig. S4), which has been previously suggested to occur via dendritic cell migration (Knoop et al., 2016). We observed bacterial dysbiosis in AMNV pre-exposed animals (Fig. 2D-2E), which has been linked to disrupted systemic antifungal immunity (Li et al., 2020). Several families that were depleted by AMNV pre-exposure (*Bacteroidales*, *Deferribacterales*, *Erysipelotrichales*, *Clostridiales*) included species that are major producers of short-chain fatty acids (SCFAs) such as *Ruminococcus*, *Roseburia* and *Butyricimonas* (Fig. S5A). Because these SCFAs promote GI tissue barrier integrity (Kelly et al., 2015), we examined the potential contribution of the loss of these SCFA-producing bacteria to antibiotic-induced susceptibility to invasive candidiasis. We delivered a cocktail of SCFAs (acetate, butyrate, propionate) to AMNV pre-exposed mice and analyzed the effect on mortality following systemic *C. albicans* challenge. We found that SCFA supplementation did not rescue antibiotic-induced susceptibility to invasive *C. albicans* infection (Fig. S5B). Together, we show that broad-spectrum antibiotic pre-exposure causes gut dysbiosis and drives GI tissue fungal proliferation upon systemic fungal challenge, which promotes subsequent bacterial GI tract translocation and systemic bacterial co-infection in a manner that is independent of SCFAs.

### **IL-17A and GM-CSF Protect against Antibiotic-Induced Susceptibility to Invasive Candidiasis**

We next aimed to define the molecular and cellular underpinnings of antibiotic-induced dysfunctional GI tract antifungal immunity and barrier integrity. Antibiotics disrupt cytokine production from GI tract-resident CD4 T cells and innate lymphoid cells (ILCs), particularly IL-17A, IL-22, and GM-CSF (Ivanov et al., 2009; Ivanov et al., 2008; Mortha et al., 2014). IL-17A is a critical antifungal molecule at the mucosal barrier as inborn errors of IL-17 immunity cause mucosal candidiasis (Boisson et al., 2013; Levy et al., 2016; Lionakis and Levitz, 2017; Puel et al., 2011). IL-22 and GM-CSF also protect against *C. albicans* infection (Aggor et al., 2020; Bichele et al., 2018; Kasahara et al., 2016), and promote intestinal barrier integrity (Gennari et al., 1994; Sonnenberg et al., 2012). We thus examined whether the production of these three cytokines by GI tract-resident lymphocytes was affected by antibiotic treatment before and during fungal infection (Fig. S6) and found significant reductions in the frequencies of IL-17A-, IL-22-, and GM-CSF-producing lymphoid cells upon antibiotic pre-exposure (Fig. 3A-3B). To determine the impact of these antibiotic-induced cytokine perturbations on mouse susceptibility to *C. albicans* infection, we infected mice

deficient in each of these cytokines and examined whether antibiotics accelerated their mortality upon fungal infection, as we observed in wild-type (WT) mice. We found that antibiotic pre-exposed *Il22*<sup>-/-</sup> animals had greater mortality post-fungal infection compared to *Il22*<sup>-/-</sup> animals that did not receive antibiotics (Fig. 3C), indicating that IL-22 is not required to mediate the susceptibility phenotype in this model. By contrast, mice lacking either IL-17A or GM-CSF exhibited similar susceptibility to fungal infection regardless of antibiotic pre-exposure (Fig. 3C). This indicates that reduced production of IL-17A and GM-CSF, but not IL-22, in antibiotic pre-exposed animals mediates their enhanced susceptibility to subsequent invasive fungal disease.

As we found that antibiotics reduced the production of GM-CSF and IL-17A in the GI tract, we next explored whether restoring these defects with recombinant cytokine treatment may rescue the susceptibility of antibiotic-treated animals when challenged with *C. albicans*. Recombinant GM-CSF is FDA-approved for accelerating myeloid reconstitution following myeloablative chemotherapy or hematopoietic stem cell transplantation, and has provided clinical benefit in some patients with invasive fungal (including *C. albicans*) infections (Dignani et al., 2005; Gavino et al., 2014; Poynton et al., 1998; Rókuscz et al., 2001; van de Veerdonk et al., 2012). Thus, we administered GM-CSF to antibiotic pre-exposed *C. albicans*-infected mice and found that it partially reversed their susceptibility (Fig. 3D). We observed a similar partial rescue with IL-17A administration (Fig. 3E). Our data therefore points to a potential translational avenue for therapeutic intervention to ameliorate antibiotic-induced infection susceptibility in this setting.

### **Both CD4 T cells and ILCs Promote Protection in the Setting of Antibiotic-Induced Invasive Candidiasis**

As both GI tract-resident CD4 T cells and ILCs of antibiotic pre-exposed mice exhibited cytokine production defects (Fig. 3A-3B), we next examined which of these populations' impairment contributes to susceptibility to systemic fungal infection upon antibiotic pre-exposure. Antibiotic-treated *Rag1*<sup>-/-</sup> mice, which lack CD4 T cells (and other  $\alpha\beta$  and  $\gamma\delta$  lymphoid cells), had greater mortality following fungal infection compared to *Rag1*<sup>-/-</sup> animals that did not receive antibiotics (Fig. 4A), indicating that the antibiotic exposure-induced susceptibility phenotype develops even in the absence of  $\alpha\beta$  (and  $\gamma\delta$ ) lymphoid cells. By contrast, we found that antibiotic-treated *Rag2*<sup>-/-</sup>*Ilr2g*<sup>-/-</sup> animals, which also lack ILCs, had similar susceptibility to fungal infection regardless of antibiotic pre-exposure (Fig. 4B). These findings indicate that defects in both innate and adaptive lymphoid populations of antibiotic pre-exposed animals mediate their enhanced susceptibility to subsequent invasive fungal infection. In line with this, non-antibiotic exposed *Rag2*<sup>-/-</sup>*Ilr2g*<sup>-/-</sup> mice phenocopied WT



antibiotic-treated mice at the levels of significantly greater fungal proliferation across the GI tract but not the kidney (Fig. 4C) and systemic escape of commensal bacteria (Fig. 4D), consistent with the notion that innate and adaptive lymphoid cells are necessary to promote fungal control within the GI tract and containment of systemic bacterial translocation. To examine whether broad-spectrum antibiotic pre-exposure is sufficient to compromise innate and adaptive lymphoid cell-mediated protection against invasive candidiasis in this setting, we adoptively transferred CD4 T cells and ILCs into *Rag2<sup>-/-</sup>Il2g<sup>-/-</sup>* mice and found that they rescued the susceptibility phenotype (Fig. 4A, 4E). Notably, adoptive transfer of CD4 T cells and ILCs harvested from antibiotic pre-exposed donors into *Rag2<sup>-/-</sup>Il2g<sup>-/-</sup>* mice significantly blunted their protective effects during invasive candidiasis (Fig. 4E). Together, these studies indicate that broad-spectrum antibiotic pre-exposure impairs lymphocyte-dependent IL-17A- and GM-CSF-mediated antifungal immunity within the GI tract, which is associated with non-inflammatory bacterial translocation, systemic bacterial co-infection, and increased mortality post-fungal challenge.

### **Vancomycin Pre-exposure Specifically Promotes Susceptibility to Invasive Candidiasis**

Different antibiotics can exert specific effects on immune cell function and/or microbiome-dependent immune modulation (Kalghatgi et al., 2013). We therefore sought to examine whether any specific antibiotic was responsible for the enhanced susceptibility to invasive candidiasis of AMNV pre-exposed mice. We single-treated WT mice with each antibiotic and evaluated survival following intravenous *C. albicans* challenge, as before. Notably, pre-exposure to vancomycin, but not metronidazole or neomycin, promoted markedly increased mortality, whereas ampicillin pre-exposure modestly increased susceptibility (Fig. 5A). Vancomycin has been associated with defective immune responses in mice (Brandl et al., 2008; van Opstal et al., 2016) and humans (Von Drygalski et al., 2007) in other settings, therefore we next focused on vancomycin to understand how it impaired GI tract antifungal immunity.

Like AMNV pre-exposed mice, vancomycin pre-exposed mice had impaired control of invasive fungal infection specifically in GI tissues but not kidney (Fig. 5B) and translocated gut commensal bacteria to the spleen and peritoneal fluid (Fig. 5C). Vancomycin-associated susceptibility was evident upon long-term pre-exposure (4 weeks), whereas a 10-day pre-treatment did not enhance susceptibility to invasive candidiasis (Fig. S7A). Moreover, vancomycin-driven susceptibility was specific to invasive fungal infection, since vancomycin pre-exposed mice did not exhibit accelerated mortality when infected systemically with *Yersinia pseudotuberculosis* (Fig. S7B). Lastly, we examined whether parenteral administration of vancomycin also increased susceptibility to invasive fungal infection. We

treated mice every 2 days with intraperitoneal (IP) vancomycin for 4 weeks, followed by intravenous challenge with *C. albicans*. In contrast to oral vancomycin pre-exposure, IP vancomycin administration did not increase susceptibility to subsequent invasive fungal infection (Fig. S7C). Collectively, our data show that vancomycin pre-exposure selectively enhances susceptibility to invasive candidiasis, which is dependent on long-term oral administration.

### **Vancomycin Pre-exposure Results in Reduced Proliferation and ROR $\gamma$ t Expression of Gut Th17 cells in a Microbiome-Dependent Manner**

We next sought to determine whether vancomycin pre-exposure conferred similar defects in intestinal lymphoid cell cytokine production as AMNV pre-exposure did (Fig. 3), focusing primarily on IL-17A. Indeed, we found that vancomycin pre-exposed mice had a significantly reduced frequency of IL-17A<sup>+</sup> CD4 T cells in the small intestine lamina propria, whereas the frequency of IL-17A<sup>+</sup> ILCs was preserved (Fig. 6A and S8A). To understand the mechanism(s) by which vancomycin pre-exposure might cause the reduction of gut IL-17A<sup>+</sup> CD4 T cells, we examined Th17 cell ROR $\gamma$ t expression, proliferation, and viability *in vivo*. Gut IL-17A<sup>+</sup> CD4 T cells from vancomycin pre-exposed mice exhibited significantly reduced expression of ROR $\gamma$ t relative to untreated controls (Fig. 6B). In addition, we found a significant reduction of proliferating CD4 T cells in the intestine of vancomycin pre-exposed mice (Fig 6C). Ki67<sup>+</sup> CD4 T cells produced more IL-17A than IFN $\gamma$ , and vancomycin pre-exposure specifically reduced proliferating IL-17A<sup>+</sup> CD4 T cells, whereas proliferating IFN $\gamma$ <sup>+</sup> CD4 T cells remained unchanged (Fig. 6D). By contrast, we observed no defect in CD4 T cell survival in the intestine of vancomycin pre-exposed mice (Fig. S8B). Moreover, ROR $\gamma$ t expression, proliferation, and viability were preserved in intestinal ILCs of vancomycin pre-exposed mice (Fig. S8B-D). Taken together, this data shows that vancomycin pre-exposure reduced gut Th17 cells associated with decreased ROR $\gamma$ t expression and proliferation of Th17 cells.

Next, we asked whether vancomycin acted directly on Th17 cells to induce the immune defects we observed *in vivo*. We first isolated naïve CD4 T cells from spleen of WT mice and cultured them *in vitro* under Th17 inducing conditions, in the presence of increasing concentrations of vancomycin. These experiments revealed that vancomycin had no effect on Th17 cell differentiation or proliferation *in vitro*, since we found similar production of IL-17A by vancomycin-treated CD4 T cells, and no difference in the expression of ROR $\gamma$ t or Ki67 (Fig S9A-C). In the same experiments, we measured GM-CSF levels by ELISA upon vancomycin treatment *in vitro* and found no difference in GM-CSF secretion by T cells in the presence of increasing concentrations of vancomycin (Fig S9C). In an

independent line of experiments, we isolated naïve CD4 T cells from the spleen of vancomycin pre-exposed mice and cultured them *in vitro* under Th17 inducing conditions to determine whether the decreased frequency of IL-17A<sup>+</sup> cells we observed *in vivo* was also evident *in vitro*. However, we found no defect in the frequency of Th17 cells *ex vivo*, nor did we detect any impairment in the secretion of GM-CSF by these cells (Fig S9D). Together, these experiments show that vancomycin does not directly impair Th17 cell differentiation or proliferation.

Since our *in vitro* and *ex vivo* experiments indicated that the Th17 cell defects only occurred with *in vivo* exposure to vancomycin, we next examined the microbiome changes occurring in the vancomycin pre-exposed mice which may disrupt microbial-derived signals that influence Th17 cell development. Indeed, it has been shown that specific bacterial species and their products can modulate the number and/or function of gut Th17 cells (Alexander et al., 2022; Atarashi et al., 2015; Ivanov et al., 2009; Martínez-López et al., 2019). We found that vancomycin pre-exposure had no effect on the abundance of total bacteria (Fig 6E), but significantly reduced the diversity of bacterial species (Fig 6F). Specifically, vancomycin pre-exposed animals had significant reductions in the relative abundance of *Bacteroides*, and expansions in the relative abundance of *Anaeroplasma*, *Deferribacteria* and *Enterobacteria* (Fig 6G). Because segmented filamentous bacteria (SFB) promote Th17 differentiation in the mouse intestine (Ivanov et al., 2009), and vancomycin has activity against SFB (Klaasen et al., 1991), we quantified the amount of SFB in the stool of vancomycin and AMNV pre-exposed animals. We found that the abundance of SFB was significantly reduced in both vancomycin and AMNV pre-exposed animals (Fig 6H). Importantly, we found no reduction in SFB was seen in mice treated with IP vancomycin (Fig 6H) that did not demonstrate increased susceptibility to systemic candidiasis (Fig S7C). Collectively, these results indicate that the reduction of Th17 cells in mice pre-exposed to antibiotics orally may be contributed, at least in part, by the loss of SFB from the microbiota.

### **Broad-spectrum Antibiotic Exposure is Associated with Susceptibility to and Mortality from Invasive Candidiasis in Humans**

To examine the potential human relevance of these findings, we utilized the Cerner *HealthFacts* database, which includes linked microbiological and clinical data for >60 million patients across the US, to determine whether broad-spectrum antibiotic pre-exposure in hospitalized patients may increase the risk of invasive candidiasis, mortality after invasive candidiasis, and/or development of systemic bacterial co-infection during invasive candidiasis. We extracted records for all hospital encounters from 2009 through 2017 where any *Candida* species was isolated from blood or other sterile site and identified and evaluated 8,294 patients with invasive candidiasis. Pre-exposure to broad-spectrum

antibiotics was defined as having received >7 days of a systemic antibiotic in the 31 days prior to the first positive *Candida* culture. The mean duration of pre-exposure to broad-spectrum antibiotics in all evaluable patients was 13 days (median, 11 days; range, 8-79 days). Among the patients with invasive *Candida* infection, the mean duration of pre-exposure to broad-spectrum antibiotics was 14.6 days (median, 12 days; range, 8-52 days). To assess the odds of developing invasive candidiasis, we used logistic regression controlling for patient, encounter, and hospital-level characteristics. Among patients with invasive candidiasis, we calculated the cumulative incidence function of the incidence of bacterial co-infection, considering mortality as a competing risk.

We found that receipt of broad-spectrum antibiotics for >7 days was significantly and independently associated with increased odds of developing invasive candidiasis (OR: 1.13, 95% CI: 1.03 – 1.25), and increased the risk of death following invasive candidiasis (HR: 1.18, 95% CI: 1.02 – 1.36) (Table 1). The receipt of >7 days of broad-spectrum antibiotics exhibited a trend towards an increased risk of bacterial co-infection in individuals with invasive candidiasis not yet experiencing the event (HR: 1.20, 95% CI: 0.90 – 1.60) (Table 1). Although additional studies are required to verify these results, this data suggest that broad-spectrum antibiotics may be a risk factor for developing invasive candidiasis, increased mortality following invasive candidiasis, and increased risk of systemic bacterial co-infection during invasive candidiasis in patients.

## Discussion

Antibiotics are a well-known risk factor for invasive candidiasis in humans, yet the mechanisms for this relationship have remained unclear. Here, we show that long-term antibiotic treatment, particularly with vancomycin, enhances susceptibility to invasive candidiasis and drives non-inflammatory escape of gut bacteria resulting in systemic co-infection and death. Using animals deficient in lymphocytes and cytokines associated with mucosal immunity and barrier integrity, we found that antibiotic treatment significantly alters the function of CD4 T cells and ILCs during invasive fungal infection, and that protection against antibiotic-associated mortality is partially dependent on IL-17A and GM-CSF.

Our data revealed that although antibiotic pre-exposed mice had increased mortality following systemic candidiasis, these animals did not have enhanced fungal infection of tissues that typically correlate with mortality (Spellberg et al., 2005; Spellberg et al., 2003). Instead, we found a GI tract-specific susceptibility to fungal infection following antibiotic pre-exposure. Laboratory-bred mice are not naturally colonized with *C. albicans* (Iliev et al., 2012), and antibiotics are needed to break this colonization resistance which is dependent on HIF1 $\alpha$ , LL-37, and certain microbiota (Fan et al., 2015). Penicillin and streptomycin are most often used to permit fungal colonization of laboratory-bred mice, as other antibiotics - including vancomycin, are less able to allow for persistent *C. albicans* GI colonization (Shankar et al., 2015). While we found evidence that our antibiotic pre-exposed animals did become colonized post-systemic infection, it is important to note that this colonization likely occurred initially from the bloodstream and was maintained by coprophagic behavior. Higher fungal burdens in the stool of intravenously-infected animals has also been observed in mice deficient in DECTIN-1, which are also unable to control fungal invasion of GI tissues (Drummond et al., 2016; Vautier et al., 2012). Mechanisms enabling *C. albicans* translocation across the intestinal barrier are incompletely understood; recent work implicated hypha formation and secretion of the peptide toxin, Candidalysin, as essential for transcellular migration across enterocyte barriers (Allert et al., 2018). Whether similar mechanisms operate to expel *C. albicans* from the deep GI tissue into the GI lumen is unknown.

In addition to uncontrolled GI fungal infection, AMNV and vancomycin pre-exposure was associated with systemic escape of gut bacteria. Although we were able to culture gut bacteria from several distal anatomical sites, we did not find evidence of overt intestinal inflammation or generalized permeability defect of the gut barrier in antibiotic pre-exposed mice. Our study adds to the growing body of evidence that antibiotics may induce the translocation of commensal bacteria across the intestinal barrier independently of epithelial cell disruption and tight-junction damage (Chakraborty et al., 2018; Yu et al., 2014). The mechanisms by which this occurs remain incompletely understood, although one study

indicated that dissemination of bacteria to the mesenteric lymph nodes requires microbial signaling to colonic goblet cells and CX<sub>3</sub>CR1<sup>+</sup> dendritic cells (Knoop et al., 2016). In these studies, antibiotic pre-exposure caused limited bacterial translocation without morbidity. By contrast, our data reveals that antibiotic-induced translocation of bacteria is significantly exacerbated by invasive fungal infection of the GI tract, which markedly elevated bacterial translocation leading to mortality.

Antibiotics are known to cause disruption to mucosal lymphocytes (Ivanov et al., 2009). IL-17A production by CD4 T cells is especially sensitive to antibiotic treatment, as we confirmed here. IL-17A is a key antifungal cytokine at mucosal sites since patients with IL-17 receptor signaling defects are susceptible to mucosal candidiasis and enhanced *C. albicans* GI colonization (Puel et al., 2011; Puel et al., 2010). Indeed, we found that IL-17A was required to control *C. albicans* infection of GI tissues in the setting of antibiotic pre-exposure, and that AMNV and vancomycin treatment significantly reduced the number of intestinal Th17 cells. A recent study by Li *et al* utilized a similar antibiotic treatment in WT animals and also found that antibiotics promoted mortality following systemic candidiasis, which was associated with reduced IL-17A production since recombinant IL-17A rescued antibiotic-treated animals (Li et al., 2020). However, in that study, antibiotics promoted uncontrolled fungal growth in several organs including the kidney and liver, and dissemination of commensal bacteria was not investigated. It is possible that the discrepancy in renal fungal control between that and our study is due to microbiota differences in the mouse colonies. In our study, both AMNV and vancomycin pre-exposed animals had reductions in *Bacteroidetes* whereas Li *et al* found little change in this phylum (Li et al., 2020). Microbiota composition is important for determining immune responses, including to fungi. Indeed, LPS produced by commensal bacteria prevents the development of protective antifungal immunity by limiting neutrophil expansion (Jiang et al., 2014), while GI colonization with *C. albicans* helps drive Th17 expansion and neutrophil reactivity, which in turn promotes protection against systemic fungal infection (Shao et al., 2019). Therefore, the composition of the commensal microbiota is an important determinant of how IL-17A-dependent responses are regulated which in turn may affect the outcome of systemic candidiasis in mice.

In line with that, we found that vancomycin-induced changes to the microbiome caused significant reductions in the proliferation and ROR $\gamma$ t expression of gut Th17 cells. Vancomycin did not directly affect the development of Th17 cells, in line with similar studies that found no direct effect of vancomycin on autoimmune Th17 cells (Almeida et al., 2021). Interestingly, the same study also found no difference in the frequency of infiltrating Th17 cells within the inflamed CNS of vancomycin-treated mice (Almeida et al., 2021), which may indicate that vancomycin exerts tissue-specific effects on Th17 cells that are context-

dependent. The commensal microbiota are important for maintaining the proliferative potential of CD4 T cells in mucosal tissues, as antibiotic treatment has been associated with reduced CD4 T cell division (Cording et al., 2013; Dutzan et al., 2018), in line with our findings. In addition, specific bacterial species or their products are required for the maintenance of functional CD4 T cell phenotypes. In particular, SFB drives the differentiation of Th17 cells in the small intestine (Ivanov et al., 2009), a process shown to be indirect and occur following bacterial binding to the intestinal epithelium (Atarashi et al., 2015). Another driver of gut Th17 cells is microbial-derived ATP. Both SFB and bacterial ATP are reduced by antibiotics which has been linked to reduced IL-17A production (Atarashi et al., 2008; Kusu et al., 2013). Furthermore, orally delivered antibiotics have a greater effect on microbiota-dependent immune modulation compared to parenteral antibiotics (Nakamura et al., 2016; Strzępa et al., 2017). Indeed, SFB abundance was reduced in mice pre-exposed to vancomycin orally but not intraperitoneally, although we cannot rule out the possibility that other parenteral routes and/or dosing schemes may also reduce SFB and thus enhance susceptibility to invasive candidiasis. Collectively, these studies may explain our findings that show vancomycin specifically enhanced susceptibility to *C. albicans* following long-term oral treatment and only indirectly via dysbiosis. Of note, oral administration of vancomycin for *Clostridioides difficile* infections has been associated with the development of invasive candidiasis in patients (Falcone et al., 2016; Russo et al., 2015).

In addition to IL-17A, we found that GM-CSF production was reduced by antibiotics and that the presence of GM-CSF was required for the development of antibiotic-induced susceptibility during invasive candidiasis. GM-CSF has been shown to have a protective benefit against invasive candidiasis in humans (Dignani et al., 2005; Gavino et al., 2014). We show here that GM-CSF can ameliorate mortality in the setting of antibiotic-associated invasive candidiasis in mice. GM-CSF treatment may be effective at boosting the antifungal activity of myeloid cells (Brown et al., 2017), which were previously shown to be susceptible to antibiotic-induced damage, including with vancomycin (Ha et al., 2014). While we did not observe antibiotic-induced quantitative defects in neutrophils, it is possible that there might be functional defects since neutrophil dysfunction is linked with enhanced susceptibility to invasive candidiasis (Lionakis and Levitz, 2017), and antibiotics have been previously shown to affect neutrophil development (Deshmukh et al., 2014). We previously found that GM-CSF therapy does not enhance neutrophil recruitment to *Candida*-infected tissues (Drummond et al., 2018), but whether this immunotherapy boosts the functional capacity of these cells, especially in the context of antibiotic treatment, remains unknown. To understand how GM-CSF exerts this protective benefit, future studies will also need to examine the mechanism by which antibiotics disrupt GM-CSF production in the GI tract. Gut bacteria have been shown to regulate GM-CSF production from ILCs and this can be disrupted by treatment with

certain antibiotics (Mortha et al., 2014). We found that vancomycin did not have a direct effect on GM-CSF production by CD4 T cells *in vitro*. Whether vancomycin (or other antibiotics) affect GM-CSF production via other mechanisms is unclear. Moreover, antibiotic pre-exposed mice may have effects beyond IL-17A/GM-CSF that could promote susceptibility to *C. albicans* infection. Future studies will need to carefully assess the direct and/or indirect mechanisms that specific antibiotics (such as vancomycin and others) perturb cellular antimicrobial immunity.

Lastly, we used a large database of patient health records to determine whether broad-spectrum antibiotic pre-exposure may be associated with the development of (and death from) invasive candidiasis, and whether trans-kingdom infections may occur more frequently in the setting of antibiotic-associated invasive candidiasis. This clinical data corroborated our observations with the mouse model. Patients treated with broad-spectrum antibiotics were more likely to develop and die from invasive candidiasis. We also noted a trend towards developing bacterial co-infections in the setting of antibiotic-associated invasive candidiasis in this dataset. Additional data will be needed to confirm our findings, however bacterial co-infections in patients with systemic *C. albicans* infections has been reported in observational studies, and this was associated with enhanced mortality and prior antibiotic treatment (Isenmann et al., 2002; Sipsas et al., 2009). Therefore, enhanced vigilance for trans-kingdom systemic infections in antibiotic-treated patients is warranted. Analysis of our patient dataset did not provide the power to examine the role of pre-exposure to different classes of antibiotics, whether oral and/or parenteral, in the development of invasive candidiasis, its mortality, and bacterial co-infection. This will be an important direction of future investigation using larger datasets and prospective study design.

In summary, our data uncover an unexpected mechanistic relationship between administration of broad-spectrum antibiotics and the development of and risk of death from life-threatening trans-kingdom systemic infections and stimulate new avenues of investigation into how antibiotics may affect antimicrobial immunity. This research adds to the growing body of evidence that antibiotics can cause immunological injuries, points towards a potential translational immune-based avenue for intervention, and underscores the importance of antibiotic stewardship, which may help prevent antimicrobial resistance and protect vulnerable patients from life-threatening invasive infections.



**Acknowledgments**

This work was supported by the Division of Intramural Research of the National Institute of Allergy & Infectious Disease, National Human Genome Research Institute, the National Cancer Institute, and the NIH Clinical Center. The computational resources of the NIH High-Performance Computation Biowulf Cluster (<http://hpc.nih.gov>) were used for this study.

**Author Contributions**

RAD and MSL designed the study. RAD, JVD, DRP, DB, DW, YB, JAS and MSL designed experiments. RAD, JVD, MS, VRM, LS, DL, CD, NMG, NC and AZ carried out the experiments. RAD, JVD, EER, SC, MQ, CCRL, NMG and AZ performed data analysis. RAD and MSL wrote the manuscript. All authors approved the final manuscript.

**Competing Interests**

The authors declare that no conflicts of interest exist.

## Figure Legends

### **Figure 1: Broad-spectrum Antibiotics Promote Mortality Following Systemic *C. albicans* Challenge.**

(A) Survival curve of untreated (n=33) and AMNV pre-exposed (n=31) C57BL/6 mice. Data is pooled from 3 independent experiments and analyzed by Log-rank Mantel-Cox test.

(B) Kidney fungal burdens in untreated mice (n=16) and AMNV pre-exposed mice (n=15 animals). Data is pooled from 2 independent experiments.

(C) BUN and creatinine levels in the serum of untreated (n=4 uninfected, n=13 infected) and AMNV pre-exposed (n=4 uninfected, n=14 infected) mice at 4 weeks post-antibiotic treatment (uninfected) or 7 days post-*C. albicans* infection (infected). Data shown with mean +/- SEM and is pooled from 2 independent experiments.

(D) Fungal burdens of indicated GI tissues (contents removed prior to homogenizing) in untreated mice (stomach n=13; small intestine n=26; caecum/colon n=15) and AMNV pre-exposed mice (stomach n=10; small intestine n=25; caecum/colon n=14) at day 7 post-infection. Data is pooled from 3 (stomach, caecum, colon) or 4 (small intestine) independent experiments and analyzed by Mann Whitney U-test.

(E) Fungal burdens in the mucus, epithelial and deep tissue layers of GI tissues in untreated (n=10) and AMNV pre-exposed (n=11) mice at day 7 post-infection. Data pooled from 2 independent experiments and analyzed by Mann-Whitney U-test. In all panels, each point represents an individual animal. \* $P < 0.05$ , \*\* $P < 0.01$ , \*\*\* $P < 0.005$ , \*\*\*\* $P < 0.0001$ .

### **Figure 2: Broad-spectrum Antibiotic Pre-exposure Promotes Dissemination of Gut Commensal Bacteria Following Fungal Infection.**

(A) Bacterial burdens in the spleen of untreated (n= 8) and AMNV-preexposed (n= 10) mice at day 7 post-fungal infection. No bacteria were detected in the spleens of AMNV pre-exposed *Candida*-uninfected or AMNV-unexposed *Candida*-infected mice. Data is pooled from 2 independent experiments and analyzed by Mann-Whitney U-test.

(B) Percentage of mice with positive bacterial cultures of the blood and peritoneal lavage at day 7 post-fungal infection. Percentages represent combined data from 2 independent experiments with at least 10 mice in each group. nd, not detected.

(C) Bacterial colonies from the spleen or peritoneal lavage fluid of AMNV pre-exposed mice were identified by MALDI-TOF (*Escherichia coli*, *Lactobacillus murinus*, *Enterococcus gallinarum*). Randomly-picked isolates were chosen from at least 6 different animals across 2 independent experiments.

(D) Relative abundance of indicated bacterial families in stools collected from untreated and antibiotic pre-exposed mice at day 0 and day 6 relative to *C. albicans* challenge. Each bar

shows the analysis for an individual mouse and at least 2 mice per group are from an independent experiment.

(E) *Left*: Shannon index of stool microbiome diversity in mice that were untreated (n=5) and AMNV pre-exposed (n=5) mice at day 6 post-infection. *Right*: Quantification of total bacterial load in stool samples collected from animals at day 6 post fungal infection. Each point represents an individual animal (n=3 per group). Data from uninfected animals showed similar results.

### **Figure 3: Broad-Spectrum Antibiotic Pre-exposure Disrupts Lymphocyte-Derived Cytokine Responses within the Gastrointestinal Tract During Invasive Fungal Infection.**

(A and B) (A) Numbers of cytokine-positive CD4<sup>+</sup> T cells and (B) ILCs in untreated (day 0 n=8; day 3 n=7) and AMNV pre-exposed (day 0 n=8; day 3 n=7) mice at indicated time points relative to fungal infection. Example FACS plots are gated on CD4<sup>+</sup> T cells using the gating strategy shown in Fig. S6. All data shown is pooled from 2 independent experiments; each point represents an individual animal. Data is analyzed by unpaired two-tailed t-tests.

(C) Survival curve of *C. albicans*-infected *Il17a*<sup>-/-</sup> (untreated, n=14; AMNV pre-exposed, n=15), *Gmcsf*<sup>-/-</sup> (untreated, n=12; AMNV pre-exposed, n=11) and *Il22*<sup>-/-</sup> (untreated, n=13; AMNV pre-exposed, n=13) mice. Data in each curve is from 3 (*Il17a*<sup>-/-</sup>) or 2 (*Gmcsf*<sup>-/-</sup>, *Il22*<sup>-/-</sup>) independent experiments and analyzed by Log rank Mantel-Cox test.

(D and E) (D) Survival curve of *C. albicans*-infected AMNV pre-exposed WT mice, with or without administration of recombinant GM-CSF, or (E) recombinant IL-17A (untreated n=20 mice, treated n=15 mice). WT mice were pre-exposed to AMNV as in Fig 1A, and 5 µg GM-CSF or 1 µg IL-17A or diluent control was delivered intraperitoneally, using the following dosing strategy: GM-CSF, days -6, -4, -2, 0, +2, +4 and +6 relative to *C. albicans* infection; IL-17A, -8 hours and +24 hours relative to *C. albicans* infection. Data in D and E is pooled from 2 independent experiments and analyzed by Log rank Mantel-Cox test. \**P*<0.05, \*\**P*<0.01, \*\*\**P*<0.005, \*\*\*\**P*<0.0001.

### **Figure 4: Defects in CD4<sup>+</sup> T cells and ILCs Mediate Broad-spectrum Antibiotic-Induced Susceptibility to Invasive Fungal Infection.**

(A) Survival curve of untreated (n=21) and AMNV pre-exposed (n=17) *Rag1*<sup>-/-</sup> animals following systemic *C. albicans* challenge. Data is pooled from 3 independent experiments and analyzed by Log rank Mantel-Cox test.

(B) Survival curve of untreated (n=10) and AMNV pre-exposed (n=9) *Rag2<sup>-/-</sup>Il2rg<sup>-/-</sup>* animals following systemic *C. albicans* challenge. Data is pooled from 2 independent experiments and analyzed by Log rank Mantel-Cox test.

(C) Fungal burdens within indicated tissues at day 7 post-infection in WT (n=12) and *Rag2<sup>-/-</sup>Il2rg<sup>-/-</sup>* (n=15) mice. Each point represents an individual animal. Data is pooled from 3 independent experiments and analyzed by Mann Whitney U-test. \**P*<0.05, \*\*\**P*<0.005.

(D) Bacterial burdens in the spleen of WT (n=12) and *Rag2<sup>-/-</sup>Il2rg<sup>-/-</sup>* (n=15) mice. Each point represents an individual animal. Data is pooled from 3 independent experiments and analyzed by Mann Whitney U-test. \*\*\**P*<0.005, \*\*\*\**P*<0.0001.

(E) Survival curve of *C. albicans*-infected *Rag2<sup>-/-</sup>Il2rg<sup>-/-</sup>* mice that had received adoptive transfer of lymphocytes from untreated WT mouse donors (n=16 recipients) or AMNV pre-exposed WT mouse donors (n=15 recipients). Data is pooled from 2 independent experiments and analyzed by Log rank Mantel-Cox test.

**Figure 5: Vancomycin increases susceptibility to invasive candidiasis.**

(A) Survival curves of C57BL/6 mice treated with the indicated antibiotics for 4 weeks prior to intravenous infection (n=33 per group), compared with untreated control mice (black lines). Data is pooled from 3 independent experiments and analyzed by Log-rank Mantel-Cox test.

(B) Fungal burdens of the indicated organs in untreated mice (kidney n=16; stomach n=13; small intestine n=26; caecum/colon n=15) and vancomycin pre-exposed mice (kidney n=16; stomach n=10; small intestine n=26; caecum n=13; colon n=15). Data is pooled from 3-4 independent experiments and analyzed by Mann Whitney U-test.

(C) Bacterial burdens in the spleen of untreated (n= 8) and vancomycin pre-exposed (n= 10) mice at day 7 post-fungal infection. Percentages represent combined data from 2 independent experiments with at least 10 mice in each group.

(D) Bacterial colonies from the spleen or peritoneal lavage fluid of vancomycin pre-exposed mice were identified by MALDI-TOF. Randomly-picked isolates were chosen from at least 6 different animals across 2 independent experiments. \*\**P*<0.01.

**Figure 6: Vancomycin pre-exposure results in Th17 cell defects in the GI tract associated with depletion of Th17-inducing bacteria from the gut microbiome.**

(A) Frequency of IL-17A<sup>+</sup> ROR $\gamma$ t<sup>+</sup> CD4 T cells in the small intestine lamina propria of untreated and vancomycin pre-exposed mice. Representative FACS plots are gated on live CD4 T cell singlets.

(B) Mean fluorescence intensity (MFI) of ROR $\gamma$ t in CD4<sup>+</sup> T cells (left panel) or IL-17A<sup>+</sup> CD4<sup>+</sup> T cells (middle panel), normalized to the mean of untreated control mice. Right panel shows representative FACS plot of ROR $\gamma$ t expression gated on Th17 cells.

(C) Frequency of Ki67<sup>+</sup> CD4 T cells and (D) the proportion of Ki67<sup>+</sup> CD4 T cells that were positive for IFN $\gamma$  or IL-17A. Shown are summary data (left panels) and representative FACS plots (right panels) gated on total CD4 T cells (C) or Ki67<sup>+</sup> CD4 T cells (D).

Data in panels A-D are pooled from two independent experiments (n=5 mice per group; each point represents an individual animal) and analyzed using unpaired t-tests. \* $P$ <0.05, \*\* $P$ <0.01, \*\*\*\* $P$ <0.0001.

(E) Quantification of total bacterial load in stool samples collected from the indicated groups of animals at day 6 post-fungal infection. Each point represents an individual animal (n=3 per group).

(F) Shannon index of stool microbiome diversity in mice that were untreated (n=5) and vancomycin pre-exposed (n=5) mice at day 6 post-infection.

(G) Relative abundance of indicated bacterial families in stools collected from untreated and vancomycin pre-exposed mice at day 0 and day 6 relative to systemic *C. albicans* challenge (untreated graphs are also shown Fig 2D). Each bar shows the analysis for an individual mouse and at least 2 mice per group are from an independent experiment.

(H) SFB abundance in the stool as detected by qPCR relative to total eubacteria from untreated (n=10), AMNV pre-exposed (n=5), vancomycin pre-exposed (oral route; n=5), or vancomycin pre-exposed (i.p. route; n=5) uninfected mice. Data analyzed by Kruskal-Wallis with Dunn's multiple correction. Each dot represents an individual animal.

**Table 1: Risk analysis of broad-spectrum antibiotic pre-exposure in humans.** Adjusted risk ratios for the association between broad-spectrum antibiotic pre-exposure and developing invasive candidiasis (Odds ratio), broad-spectrum antibiotic pre-exposure and 31-day mortality post-invasive *Candida* infection (hazards ratio), and broad-spectrum antibiotic pre-exposure and developing a systemic bacterial co-infection within 31 days after developing invasive candidiasis (where 31-day mortality was treated as a competing risk, subdistribution hazard ratio). See main text and methods section for total numbers of patients and details of co-variables used. CI, confidence interval.

<b>Clinical variable</b>	<b>Adjusted risk ratio</b>	<b>Lower CI</b>	<b>Upper CI</b>
Invasive candidiasis	<b>1.13</b>	<b>1.03</b>	<b>1.25</b>
Mortality post-invasive candidiasis	<b>1.18</b>	<b>1.02</b>	<b>1.36</b>
Bacterial co-infection, post-invasive candidiasis	<b>1.20</b>	<b>0.90</b>	<b>1.60</b>

## STAR METHODS

### RESOURCE AVAILABILITY

#### Lead Contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Michail Lionakis (lionakism@niaid.nih.gov).

#### Materials Availability

This study did not generate new unique reagents.

#### Data and Code Availability

16S sequencing data have been deposited at NCBI BioProject under the ID number PRJNA812495 and are publicly available as of the date of publication. Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

### EXPERIMENTAL MODEL AND SUBJECT DETAILS

#### Animals

8-12 week-old female mice were maintained in individually ventilated cages under specific pathogen-free conditions at the National Institutes of Health (Bethesda, MD, USA). Animal studies were performed in accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health, under the auspices of protocol LCIM14E approved by the Animal Care and Use Committee of the National Institute of Allergy and Infectious Diseases, or under project license PBE275C33 at the University of Birmingham approved by the local Animal Welfare and Ethical Review Board. WT refers to C57BL/6 (Taconic) or the corresponding littermate/genetically matched controls of the genetically-modified lines; *Rag1*<sup>-/-</sup>, *Rag2*<sup>-/-</sup>*Il2rg*<sup>-/-</sup>, *Il17a*<sup>-/-</sup> (all Taconic), *Il22*<sup>-/-</sup> (Jackson) and *Gmcsf*<sup>-/-</sup> (Charles River). In some experiments, transgenic animals that report on IL-17A production were used. These included *Il17a*<sup>Smart</sup> (Price et al., 2012) and *Il17a*<sup>Cre</sup> animals crossed with *Rosa26*<sup>flSTOP-tdRFP</sup> mice (Fiancette et al., 2021); these animals were bred at the Biomedical Services Unit at the University of Birmingham. Animals were euthanized by cervical dislocation following administration of ketamine/xylazine cocktail at defined time-points post-infection (see Figure legends) or when humane endpoints (e.g. hypothermia, weight loss) had been reached, whichever occurred earlier.

### **Mouse Antibiotic Treatments and Fungal Infections**

Ampicillin (1 mg/mL), metronidazole (1 mg/mL), neomycin (1 mg/mL) and vancomycin (0.5 mg/mL) were diluted in autoclaved water and delivered to mice *ad libitum* as their drinking water for up to 4 weeks, either alone or in combination. Waters were refreshed twice weekly. In some experiments, mice were pre-exposed to vancomycin delivered intraperitoneally at 100 µg every 2 days for 4 weeks. *Candida albicans* strain SC5314 was used for all infections. Yeast was serially passaged 3 times in YPD (yeast extract, bacto-peptone and dextrose) broth, grown at 30°C with shaking for 18-24 hours at each passage. Yeast cells were washed in PBS, counted, and  $0.95 \times 10^5$  yeast cells injected intravenously via the lateral tail vein. To control for the background level of sucrose found in the antibiotic cocktail, we treated animals with sucrose in their drinking water and challenged with *C. albicans* as above. From 2 independent experiments, we found no difference in survival between untreated (n=20; survival rate 20%) and sucrose-treated (n=18; survival rate 22%) mice.

### ***Yersinia pseudotuberculosis* Mouse Infection Model**

Infection and analysis of mouse survival was carried out as described previously (Collins et al., 2019). Briefly, WT *Yersinia pseudotuberculosis* (32777) was grown on MacConkey plates at room temperature for 48 hours. A single colony was picked and grown as a liquid bacterial culture in 2X YT media (Sigma) shaking (220 rpm) at 25°C overnight. Mice were infected intravenously with 200 colony forming units (CFU). Survival (defined as 30% weight loss) was assessed daily.

### **Mouse Short-Chain Fatty Acid Treatments**

Acetate, propionate and butyrate (all from Sigma) were diluted in autoclaved water and delivered to mice *ad libitum* as their drinking water for at least one week prior to infection (in combination with antibiotics) and continued throughout the infection period. Final concentrations in the drinking water were 40 mM butyrate, 67.5 mM acetate and 25.9 mM propionate (Smith et al., 2013). Waters were refreshed twice weekly.

### **Mouse Adoptive Transfers**

C57BL/6 mice (Taconic) were treated for 4 weeks with AMNV (see above) or control water, and then used as donors for the transfers. Lungs and small intestines were digested as outlined above (lungs were put directly into the same digest buffer as the intestines, after finely mincing), then stained with anti-CD45, anti-CD90.2, anti-CD127 and lineage markers (CD3, CD19, CD11b). Isolated leukocytes from the lung/intestine were pooled and ILCs were FACS-sorted (defined as CD45<sup>+</sup>CD90.2<sup>+</sup>CD127<sup>+</sup>lineage<sup>-</sup>) into sterile sorting buffer (HBSS supplemented with 2 mM EDTA, 10% FCS, 100 U/mL penicillin, 100 µg/mL



streptomycin) using a FACS Aria instrument. CD4<sup>+</sup> T-cells were isolated from the spleens of donor mice using the Miltenyi CD4<sup>+</sup> T-cell negative selection kit, as per the manufacturer's instructions. Purified ILCs and CD4<sup>+</sup> T-cells were combined in sterile PBS at a ratio of 200:1 (T-cells: ILCs) and 100  $\mu$ L (2x10<sup>6</sup> CD4 T-cells, 1x10<sup>4</sup> ILCs) injected intraperitoneally into *Rag2<sup>-/-</sup>Il2rg<sup>-/-</sup>* recipients. The mice were infected with *C. albicans* approximately 18 hours after the adoptive transfer of CD4 T-cells and ILCs.

## Clinical Data

Cerner *HealthFacts*, a large de-identified electronic health record database from a well-distributed cohort of US academic and community hospitals (Cerner Inc., Kansas City, MO), was used to identify hospital encounters from 2009-2017 where any *Candida* species was isolated from any body source (excluding stool and respiratory samples). Encounters with more than one *Candida* spp isolated were excluded. Invasive disease was defined as *Candida* isolated from blood, sterile abdominal, or other non-abdominal sterile site microbiology test. Bacterial co-infection was assessed in these patients by determining whether they had any positive bacterial microbiology test from blood or a sterile source (including abdominal, respiratory, and urinary tract) 1-31 days after the first positive *Candida* culture was drawn. Patients with evidence of bacterial infection in the 31 days prior to the first positive *Candida* culture were excluded. Prior exposure to antibiotics was defined as having received  $\geq 7$  days of a broad-spectrum systemic antibiotic (i.e. clindamycin, vancomycin, piperacillin/tazobactam, ceftolozane/tazobactam, ceftriazone, levofloxacin) in the 31 days prior to the first positive *Candida* culture. In our analysis, there were 116,337 unique inpatient hospitalizations that were positive for *Candida* (any species), of which 6,681 were excluded due to evidence of bacterial infection prior to *Candida* infection diagnosis. Of the remaining 110,044 encounters, 7.5% (n=8,294) were invasive *Candida* infections, and 15% (n=14,737) of patients died or were discharged to hospice. 5.5% (n=6,022) received at least one systemic antibiotic for >7 days.

## METHOD DETAILS

### Determination of Organ Fungal and Bacterial Burdens

At defined time points post-infection (see Figure legends), animals were euthanized and organs were weighed, homogenized in PBS, and serially diluted before plating onto YPD agar supplemented with 100 U/mL penicillin and 100  $\mu$ g/mL streptomycin (for fungal burdens), or onto MacConkey and CNA selective agar (Remel) for enumeration of Gram-negative and Gram-positive bacteria, respectively. Colonies were counted after incubation at

37°C for 24-48 hours. Contents of the stomach, intestines and caecum were removed prior to homogenization and plating.

### **Serological Determination of Renal Function**

Terminal blood samples were taken via cardiac puncture and serum obtained by centrifugation at 15,000 rpm for 10 minutes at 4°C. Serum samples were diluted in sterile PBS and analyzed for concentrations of blood urea nitrogen (BUN) and creatinine as markers of renal function at the NIH Clinical Chemistry Laboratory.

### **Histopathology**

Organs were removed from infected mice at the indicated time points and fixed in 10% formalin for 24 hours before embedding in paraffin wax. Tissue sections were stained with either periodic acid-Schiff (PAS) and/or hematoxylin and eosin (H&E).

### **Determination of Gastrointestinal Fungal Burdens**

In some experiments, GI fungal burdens were determined in the different layers of GI tissues. Aseptically removed GI tissues were opened up laterally (to create a flat sheet) and soaked in 10 mL sterile PBS before shaking vigorously to remove contents. Washed tissues were then added to 5 mL of HBSS supplemented with 5% FBS and 25 mM HEPES, shaken vigorously and supernatant stored in a fresh tube (this was designated as 'mucus layer'). This step was repeated to create 10 mL of 'mucus layer', which was centrifuged at 4000rpm for 4 minutes and pellets resuspended in 1 mL sterile PBS prior to plating on YPD plates for determination of fungal burdens. After mucus removal, GI tissues were added to 5mL of pre-warmed HBSS supplemented with 15 mM HEPES, 5 mM EDTA, 10% FBS and 1 mM dithioerythritol. Tissues were vigorously shaken twice and supernatants collected and resuspended into 1 mL sterile PBS, as before. Supernatants from this step were designated as 'epithelial layer'. Tissues were finally washed twice in sterile PBS (by shaking in 5 mL each time), and then remaining tissue homogenized in 1mL sterile PBS prior to plating (designated 'deep tissue layer')(Kang et al., 2020).

### **Bacterial Identification by MALDI-TOF MS**

Individual colonies were isolated from MacConkey and CNA plates and re-streaked onto Sheep Blood agar plates (Remel) to create single cultures. Microbiological identification of isolates was performed using MALDI-TOF MS (Bruker Daltonics). For protein extraction, bacterial colonies in sheep blood agar plate were resuspended in 1 ml 70 % ethanol, vortexed for 1 min, and centrifuged at 13,000 rpm for 2 min. The supernatant was removed completely, and the sample was vortexed for 10 seconds with 50 µl of 70 % formic acid and

50 µl acetonitrile. After a 2 min centrifugation at 13,000 rpm, 1 µl of supernatant was spotted onto the target plate and overlaid with 2 µl of alpha-cyano-4-hydroxycinnamic acid. MALDI-TOF MS analysis was performed using a Microflex LT spectrometer (Bruker Daltonics) and the Biotyper version 4.0.0.1. Manufacturer-recommended cutoff scores of  $\geq 2.0$  for species-level identification and  $\geq 1.7$  for genus-level identification, and  $>10\%$  difference of top score from other genera and species were applied.

### **Preparation of Genomic DNA from Murine Stool Samples**

Microbiome analysis was performed on total DNA extracted from stool samples, collected from mice housed in sterilized empty cages using sterilized forceps. Stool samples were stored dry at  $-80^{\circ}\text{C}$  until DNA extraction. Total DNA was extracted from stool samples using the PowerLyzer PowerSoil DNA Extraction Kit (MoBio) as per the manufacturer's instructions, with the following modifications: Proteinase K (5 mg/mL) was added to the stool sample in the provided lysis buffer (C1) and incubated at  $65^{\circ}\text{C}$  for 10 minutes. Bead-beating was performed on the samples for 2 minutes at 30 Hz using the TissueLyzer (MoBio). Samples were then centrifuged and processed as outlined in the manufacturer's instructions.

### **Quantitative PCR for Segmented Filamentous Bacteria**

To assess the abundance of Segmented Filamentous Bacteria (SFB) in total genomic DNA from stool samples, qPCR reactions were carried out using 5 ng of total DNA, Perfecta SYBR Green Fast mix (Quanta Biosciences) and primers specific for SFB (SFB736F: 5'-GACGCTGAGGCATGAGAGCAT - 3'; SFB844R: 5'-GACGGCACGGATTGTTATTCA - 3') or total eubacteria (UniF340: 5' - ACTCCTACGGGAGGCAGCAGT - 3'; UniF514: 5' - TATTACCGCGGCTGCTGGC - 3'), as reported previously (Molloy et al., 2013).

### **16S Amplicon Sequencing and Analysis**

16S rRNA sequencing libraries were prepared based on a previously described strategy (Findley et al., 2013). The V1-V3 region of the 16S rRNA gene was amplified using primers 27F (5'-AGAGTTTGATCCTGGCTCAG) and 534R (5'-ATTACCGCGGCTGCTGG). Amplicons were generated and processed as previously described (Fadrosh et al., 2014; Jo et al., 2016). Paired-end reads were quality trimmed and joined using FLASH v1.2.11 (<https://github.com/dstree/FLASH2>) and parameters of Min overlap: 10, Max mismatch density: 0.25. Joined pairs were analyzed using the Nephela platform (v.2.15.0, <http://nephela.niaid.nih.gov>), in particular the DADA2 based pipeline with default parameters. This pipeline de-noised the sequences, removed chimeras and assigned amplicon sequence variants (ASVs) according to the DADA2 algorithm (Weber et al., 2018). Taxonomic assignment was performed with the RDP method against the SILVA

v.132 database (Quast et al., 2013). An unrarefied and rarefied biom file with 895 ASVs (40,000 reads/sample) was produced and used for visualization and downstream analyses. Alpha diversity (Shannon) was performed in R with package phyloseq (McMurdie and Holmes, 2013) and phylosmith <https://github.com/schuyler-smith/phylosmith>.

### **Isolation and Stimulation of Lamina Propria Leukocytes**

The entire small intestine was aseptically removed and contents flushed out with ice-cold PBS. Intestines were opened up laterally and rinsed with ice-cold PBS to wash. Washed intestines were cut into small (1-2 mm) squares and added to 10mL pre-warmed RPMI supplemented with 15% FBS and 5 mM EDTA. Intestine pieces were incubated at 37°C with shaking for 20 minutes, vortexed vigorously for 8 seconds and supernatants removed. 10mL of pre-warmed RPMI supplemented with 5 mM EDTA was added to intestine pieces, vortexed again and supernatants removed. Intestine pieces were washed in PBS and then added to 10 mL digest buffer: RPMI supplemented with 10% FBS, 100 U/mL penicillin, 100 µg/mL streptomycin, 50 µM β-mercaptoethanol, 1 mg/mL collagenase D (Worthington), 1 mg/mL Dispase (Gibco) and 0.2 mg/mL DNase (Roche). Intestines were incubated in digest buffer at 37°C with shaking for approximately 30 minutes until tissue had disintegrated. Digested intestines were poured through a 40µm filter, spun down and resuspended in RPMI supplemented with 10% FBS, 100 U/mL penicillin, 100 µg/mL streptomycin and 50 µM β-mercaptoethanol. In some experiments, intestinal leukocytes were stimulated with PMA (50 ng/mL) and ionomycin (500 ng/mL) in the presence of Brefeldin A (3 µg/mL) for 5 hours at 37°C prior to washing in PBS and staining for flow cytometry analysis.

### **Flow Cytometry**

Isolated leukocytes were resuspended in PBS and stained with Live/Dead stain (Invitrogen) on ice as per manufacturer's instructions. Fc receptors were blocked with anti-CD16/32 and staining with fluorochrome-labelled antibodies was performed on ice. Labelled samples were acquired immediately or fixed and permeabilized for intracellular staining using the Foxp3 staining kit (eBioscience). Anti-mouse antibodies used in this study were: CD45 (30-F11), CD11b (M1/70), CD11c (N418), MHC Class II (M5/114.15.2), CD3 (145-2C11), CD19 (eBio1D3), F480 (BM8), CD127 (A7R34), IL-17A (TC11-18H10.1), IL-22 (1H8PWSR), Ki67 (SolA15), RORgt (AFKJS-9) all from eBioscience, Ly6G (1A8), Ly6C (AL-21), all from BD Biosciences, and GM-CSF (MP1-22E9), CD90.2 (30-H12) all from Biolegend. Samples were acquired on a BD LSR Fortessa equipped with BD FACSDiva software. Analysis was performed using FlowJo (TreeStar).

### ***In vitro* Th17 Cell Differentiation**

Naïve CD4 T cells (from spleens of untreated or vancomycin-preexposed mice) were first isolated using the Mojo negative selection kit (Biolegend). These cells were then further purified using FACS (using the BD FACSAria) based on markers CD4<sup>+</sup>CD44<sup>low</sup>CD25<sup>-</sup> which typically yielded >95% purity. In some experiments, IL-17A reporter animals (*Il17a<sup>Smart</sup>*) were used to determine the transcription of IL-17A, in which IL-17A production is detected by staining for the human nerve growth factor receptor (hNGFR) that is under the same transcriptional control as IL-17A (Price et al., 2012). Naïve CD4 T cells were seeded into 96 well round-bottom plates (1x10<sup>5</sup> cells per well) that had been pre-coated with anti-CD3 (2µg/mL, 145-2C11, Biolegend) and anti-CD28 (10µg/mL, 37.51, BD) for 3 hours at 37°C. T cells were cultured in these pre-coated plates in complete IMDM media (Gibco) supplemented with 10% FBS, 1% Pen/Strep, 1% L-glutamine, 0.1% β-mercaptoethanol, 2ng/mL TGFβ, 25ng/mL IL-6, 10ng/mL IL-1β and 20µg/mL anti-IFNγ (all Biolegend) for 72 hours at 37°C with 5% CO<sub>2</sub>. Some cultures were further supplemented with vancomycin hydrochloride (Sigma; see figure legends for final concentrations). Culture supernatants were collected and stored at -80°C prior to ELISA-based detection for IL-17A or GM-CSF (DuoSet, RnD Systems). CD4 T-cells were stained with anti-CD4-AlexaFluor700 (RM4.4, Biolegend) and anti-HNGFR-APC (ME20.4, Biolegend) and analyzed by flow cytometry as outlined above.

## **QUANTIFICATION AND STATISTICAL ANALYSIS**

### **Statistical analysis of mouse and in vitro data**

All data is expressed as mean +/- SEM unless otherwise stated in the figure legends. Statistical analyses were performed using GraphPad Prism 9.0 software. Details of individual tests are included in the figure legends. In general, data was tested for normal distribution by Kolmogorov-Smirnov normality test and analyzed accordingly by unpaired two-tailed *t*-test or Mann Whitney *U*-test. In cases where multiple data sets were analyzed, two-way ANOVA was used with Bonferroni correction. In all cases, *P* values <0.05 were considered significant.

### **Risk Analysis in Patient Cohort Data**

To assess the odds of developing an invasive *Candida* infection, logistic regression was used controlling for patient, encounter, and hospital-level characteristics. These variables were the patient's gender and age, whether they had received any parenteral nutrition, had a prior abdominal surgery, or had a central venous catheter, and the maximum Elixhauser

Comorbidity Index, as well as the hospital bed size, whether it was a teaching facility, and whether it was urban or rural. The hospital bed sized was used to control for potential differential antibiotic prescribing and treatment practices in various acuity hospital settings. Among those patients with invasive *Candida* infections, the cumulative incidence function of the incidence of bacterial co-infection, taking into account mortality as a competing risk, was calculated. A Fine-Gray proportional hazards model was used to regress development of a bacterial coinfection on antibiotic pre-exposure using the same definitions and covariates as above, with the additional inclusion of days of antibiotics post-*Candida* infection. We also assessed the risk of 31-day mortality using a Cox proportional hazards model, with coinfection as a covariate.

## References

Aggor, F.E.Y., Break, T.J., Trevejo-Nunez, G., Whibley, N., Coleman, B.M., Bailey, R.D., Kaplan, D.H., Naglik, J.R., Shan, W., Shetty, A.C., et al. (2020). Oral epithelial IL-22/STAT3 signaling licenses IL-17-mediated immunity to oral mucosal candidiasis. *Sci Immunol.* 5(48). Published online 2020/06/07 DOI: 10.1126/sciimmunol.aba0570.

Alexander, M., Ang, Q.Y., Nayak, R.R., Bustion, A.E., Sandy, M., Zhang, B., Upadhyay, V., Pollard, K.S., Lynch, S.V., and Turnbaugh, P.J. (2022). Human gut bacterial metabolism drives Th17 activation and colitis. *Cell Host & Microbe.* 30(1), 17-30.e19. DOI: 10.1016/j.chom.2021.11.001.

Allert, S., Förster, T.M., Svensson, C.-M., Richardson, J.P., Pawlik, T., Hebecker, B., Rudolphi, S., Juraschitz, M., Schaller, M., Blagojevic, M., et al. (2018). *Candida albicans*-Induced Epithelial Damage Mediates Translocation through Intestinal Barriers. *mBio.* 9(3), e00915-00918. DOI: 10.1128/mBio.00915-18.

Almeida, L., Dhillon-LaBrooy, A., Castro, C.N., Adossa, N., Carriche, G.M., Guderian, M., Lippens, S., Dennerlein, S., Hesse, C., Lambrecht, B.N., et al. (2021). Ribosome-Targeting Antibiotics Impair T Cell Effector Function and Ameliorate Autoimmunity by Blocking Mitochondrial Protein Synthesis. *Immunity.* 54(1), 68-83.e66. DOI: <https://doi.org/10.1016/j.immuni.2020.11.001>.

Atarashi, K., Nishimura, J., Shima, T., Umesaki, Y., Yamamoto, M., Onoue, M., Yagita, H., Ishii, N., Evans, R., Honda, K., et al. (2008). ATP drives lamina propria TH17 cell differentiation. *Nature.* 455(7214), 808-812. DOI: 10.1038/nature07240.

Atarashi, K., Tanoue, T., Ando, M., Kamada, N., Nagano, Y., Narushima, S., Suda, W., Imaoka, A., Setoyama, H., Nagamori, T., et al. (2015). Th17 Cell Induction by Adhesion of Microbes to Intestinal Epithelial Cells. *Cell.* 163(2), 367-380. DOI: 10.1016/j.cell.2015.08.058.

Bichele, R., Kärner, J., Truusalu, K., Smidt, I., Mändar, R., Conti, H.R., Gaffen, S.L., Peterson, P., Laan, M., and Kisand, K. (2018). IL-22 neutralizing autoantibodies impair fungal clearance in murine oropharyngeal candidiasis model. *Euro J Immunol.* 48(3), 464-470. Published online 2017/12/11 DOI: 10.1002/eji.201747209.

Boisson, B., Wang, C., Pedergrana, V., Wu, L., Cypowyj, S., Rybojad, M., Belkadi, A., Picard, C., Abel, L., Fieschi, C., et al. (2013). An ACT1 mutation selectively abolishes interleukin-17 responses in humans with chronic mucocutaneous candidiasis. *Immunity.* 39(4), 676-686. DOI: 10.1016/j.immuni.2013.09.002.

Brandl, K., Plitas, G., Mihu, C.N., Ubeda, C., Jia, T., Fleisher, M., Schnabl, B., DeMatteo, R.P., and Pamer, E.G. (2008). Vancomycin-resistant enterococci exploit antibiotic-induced innate immune deficits. *Nature.* 455(7214), 804-807. DOI: [http://www.nature.com/nature/journal/v455/n7214/supinfo/nature07250\\_S1.html](http://www.nature.com/nature/journal/v455/n7214/supinfo/nature07250_S1.html).

Brown, R.L., Sequeira, R.P., and Clarke, T.B. (2017). The microbiota protects against respiratory infection via GM-CSF signaling. *Nature Communications.* 8(1), 1512. DOI: 10.1038/s41467-017-01803-x.

Chakraborty, R., Lam, V., Kommineni, S., Stromich, J., Hayward, M., Kristich, C.J., and Salzman, N.H. (2018). Ceftriaxone Administration Disrupts Intestinal Homeostasis, Mediating Noninflammatory Proliferation and Dissemination of Commensal Enterococci. *Infect Immun.* 86(12), e00674-00618. DOI: 10.1128/IAI.00674-18.

Collins, N., Han, S.J., Enamorado, M., Link, V.M., Huang, B., Moseman, E.A., Kishton, R.J., Shannon, J.P., Dixit, D., Schwab, S.R., et al. (2019). The Bone Marrow Protects and Optimizes Immunological Memory during Dietary Restriction. *Cell.* 178(5), 1088-1101.e1015. Published online 2019/08/24 DOI: 10.1016/j.cell.2019.07.049.

Cording, S., Fleissner, D., Heimesaat, M.M., Bereswill, S., Loddenkemper, C., Uematsu, S., Akira, S., Hamann, A., and Huehn, J. (2013). Commensal microbiota drive proliferation of conventional and Foxp3(+) regulatory CD4(+) T cells in mesenteric lymph nodes and Peyer's patches. *Eur J Microbiol Immunol (Bp).* 3(1), 1-10. Published online 2013/03/13 DOI: 10.1556/EuJMI.3.2013.1.1.

Deshmukh, H.S., Liu, Y., Menkiti, O.R., Mei, J., Dai, N., O'Leary, C.E., Oliver, P.M., Kolls, J.K., Weiser, J.N., and Worthen, G.S. (2014). The microbiota regulates neutrophil homeostasis and host resistance to *Escherichia coli* K1 sepsis in neonatal mice. *Nat Med.* 20(5), 528-534. DOI: 10.1038/nm.3542.

Dignani, M.C., Rex, J.H., Chan, K.W., Dow, G., deMagalhaes-Silverman, M., Maddox, A., Walsh, T., and Anaissie, E. (2005). Immunomodulation with interferon-gamma and colony-stimulating factors for refractory fungal infections in patients with leukemia. *Cancer.* 104(1), 199-204. Published online 2005/06/02 DOI: 10.1002/cncr.21142.

Drummond, R.A., Dambuza, I.M., Vautier, S., Taylor, J.A., Reid, D.M., Bain, C.C., Underhill, D.M., Masopust, D., Kaplan, D.H., and Brown, G.D. (2016). CD4+ T-cell survival in the GI tract requires dectin-1 during fungal infection. *Mucosal Immunol.* 9, 492-502. DOI: 10.1038/mi.2015.79.

Drummond, R.A., Zahra, F.T., Natarajan, M., Swamydas, M., Hsu, A.P., Wheat, L.J., Gavino, C., Vinh, D.C., Holland, S.H., Mikelis, C.M., et al. (2018). GM-CSF Therapy in Human CARD9 Deficiency. *J Allergy Clin Immunol.* DOI: 10.1016/j.jaci.2018.05.025.

Dutzan, N., Kajikawa, T., Abusleme, L., Greenwell-Wild, T., Zuazo, C.E., Ikeuchi, T., Brenchley, L., Abe, T., Hurabielle, C., Martin, D., et al. (2018). A dysbiotic microbiome triggers T<sub>H</sub>17 cells to mediate oral mucosal immunopathology in mice and humans. *Science translational medicine.* 10(463). DOI: 10.1126/scitranslmed.aat0797.

Erny, D., Hrabe de Angelis, A.L., Jaitin, D., Wieghofer, P., Staszewski, O., David, E., Keren-Shaul, H., Mhlahoi, T., Jakobshagen, K., Buch, T., et al. (2015). Host microbiota constantly control maturation and function of microglia in the CNS. *Nat Neurosci.* 18. DOI: 10.1038/nn.4030

<http://www.nature.com/neuro/journal/vaop/ncurrent/abs/nn.4030.html#supplementary-information>.

Fadrosh, D.W., Ma, B., Gajer, P., Sengamalay, N., Ott, S., Brotman, R.M., and Ravel, J. (2014). An improved dual-indexing approach for multiplexed 16S rRNA gene sequencing on the Illumina MiSeq platform. *Microbiome.* 2(1), 6. Published online 2014/02/25 DOI: 10.1186/2049-2618-2-6.



Falcone, M., Russo, A., Iraci, F., Carfagna, P., Goldoni, P., Vullo, V., and Venditti, M. (2016). Risk Factors and Outcomes for Bloodstream Infections Secondary to *Clostridium difficile* Infection. *Antimicrob Agents Chemother.* 60(1), 252-257. Published online 2015/10/21 DOI: 10.1128/aac.01927-15.

Fan, D., Coughlin, L.A., Neubauer, M.M., Kim, J., Kim, M.S., Zhan, X., Simms-Waldrip, T.R., Xie, Y., Hooper, L.V., and Koh, A.Y. (2015). Activation of HIF-1[alpha] and LL-37 by commensal bacteria inhibits *Candida albicans* colonization. *Nat Med.* 21, 808-814. DOI: 10.1038/nm.3871  
<http://www.nature.com/nm/journal/vaop/ncurrent/abs/nm.3871.html#supplementary-information>.

Fiancette, R., Finlay, C.M., Willis, C., Bevington, S.L., Soley, J., Ng, S.T.H., Baker, S.M., Andrews, S., Hepworth, M.R., and Withers, D.R. (2021). Reciprocal transcription factor networks govern tissue-resident ILC3 subset function and identity. *Nat. Immunol.* 22(10), 1245-1255. DOI: 10.1038/s41590-021-01024-x.

Findley, K., Oh, J., Yang, J., Conlan, S., Deming, C., Meyer, J.A., Schoenfeld, D., Nomicos, E., Park, M., Intramural Sequencing Ctr, N.I.H., et al. (2013). Topographic diversity of fungal and bacterial communities in human skin. *Nature.* 498(7454), 367-370. DOI: 10.1038/nature12171.

Gavino, C., Cotter, A., Lichtenstein, D., Lejtenyi, D., Fortin, C., Legault, C., Alirezaie, N., Majewski, J., Sheppard, D.C., Behr, M.A., et al. (2014). CARD9 Deficiency and Spontaneous Central Nervous System Candidiasis: Complete Clinical Remission With GM-CSF Therapy. *Clin Infect Dis.* 59(1), 81-84. DOI: 10.1093/cid/ciu215.

Gennari, R., Alexander, J.W., Gianotti, L., Eaves-Pyles, T., and Hartmann, S. (1994). Granulocyte macrophage colony-stimulating factor improves survival in two models of gut-derived sepsis by improving gut barrier function and modulating bacterial clearance. *Ann Surg.* 220(1), 68-76. DOI: 10.1097/0000658-199407000-00010.

Ha, Y.E., Kong, K.-H., Cho, M.-H., Kim, D.-H., Song, Y.-S., and Yoon, S.-Y. (2014). Vancomycin blocks autophagy and induces interleukin-1 $\beta$  release in macrophages. *J Antibiotics.* 68, 76. DOI: 10.1038/ja.2014.112.

Iliev, I.D., Funari, V.A., Taylor, K.D., Quoclinh, N., Reyes, C.N., Strom, S.P., Brown, J., Becker, C.A., Fleshner, P.R., Dubinsky, M., et al. (2012). Interactions Between Commensal Fungi and the C-Type Lectin Receptor Dectin-1 Influence Colitis. *Science.* 336(6086), 1314-1317. DOI: 10.1126/science.1221789.

Isenmann, R., Schwarz, M., Rau, B., Trautmann, M., Schober, W., and Beger, H.G. (2002). Characteristics of infection with *Candida* species in patients with necrotizing pancreatitis. *World J Surgery.* 26(3), 372-376. DOI: 10.1007/s00268-001-0146-9.

Ivanov, I.I., Atarashi, K., Manel, N., Brodie, E.L., Shima, T., Karaoz, U., Wei, D., Goldfarb, K.C., Santee, C.A., Lynch, S.V., et al. (2009). Induction of Intestinal Th17 Cells by Segmented Filamentous Bacteria. *Cell.* 139(3), 485-498. DOI: <http://dx.doi.org/10.1016/j.cell.2009.09.033>.

Ivanov, I.I., Frutos, R.d.L., Manel, N., Yoshinaga, K., Rifkin, D.B., Sartor, R.B., Finlay, B.B., and Littman, D.R. (2008). Specific Microbiota Direct the Differentiation of IL-17-Producing T-Helper Cells in the Mucosa of the Small Intestine. *Cell Host Microbe*. 4(4), 337-349.

Jiang, T.T., Chaturvedi, V., Ertelt, J.M., Xin, L., Clark, D.R., Kinder, J.M., and Way, S.S. (2014). Commensal enteric bacteria lipopolysaccharide impairs host defense against disseminated *Candida albicans* fungal infection. *Mucosal Immunol*. DOI: 10.1038/mi.2014.119.

Jo, J.H., Deming, C., Kennedy, E.A., Conlan, S., Polley, E.C., Ng, W.I., Segre, J.A., and Kong, H.H. (2016). Diverse Human Skin Fungal Communities in Children Converge in Adulthood. *J Invest Dermatol*. 136(12), 2356-2363. Published online 2016/08/02 DOI: 10.1016/j.jid.2016.05.130.

Kalghatgi, S., Spina, C.S., Costello, J.C., Liesa, M., Morones-Ramirez, J.R., Slomovic, S., Molina, A., Shirihai, O.S., and Collins, J.J. (2013). Bactericidal Antibiotics Induce Mitochondrial Dysfunction and Oxidative Damage in Mammalian Cells. *Sci Trans Med*. 5(192), 192ra185. DOI: 10.1126/scitranslmed.3006055.

Kang, B., Alvarado, L.J., Kim, T., Lehmann, M.L., Cho, H., He, J., Li, P., Kim, B.-H., Larochelle, A., and Kelsall, B.L. (2020). Commensal microbiota drive the functional diversification of colon macrophages. *Mucosal Immunol*. 13(2), 216-229. DOI: 10.1038/s41385-019-0228-3.

Kasahara, S., Jhingran, A., Dhingra, S., Salem, A., Cramer, R.A., and Hohl, T.M. (2016). GM-CSF signaling regulates neutrophil antifungal activity and the oxidative burst during respiratory fungal challenge. *J Infect Dis*. 213, 1289-1298. DOI: 10.1093/infdis/jiw054.

Keighley, C., Chen, S.C., Marriott, D., Pope, A., Chapman, B., Kennedy, K., Bak, N., Underwood, N., Wilson, H.L., McDonald, K., et al. (2019). *Candidaemia* and a risk predictive model for overall mortality: a prospective multicentre study. *BMC Infect Dis*. 19(1), 445. Published online 2019/05/23 DOI: 10.1186/s12879-019-4065-5.

Kelly, Caleb J., Zheng, L., Campbell, Eric L., Saeedi, B., Scholz, Carsten C., Bayless, Amanda J., Wilson, Kelly E., Glover, Louise E., Kominsky, Douglas J., Magnuson, A., et al. (2015). Crosstalk between Microbiota-Derived Short-Chain Fatty Acids and Intestinal Epithelial HIF Augments Tissue Barrier Function. *Cell Host & Microbe*. 17(5), 662-671. DOI: 10.1016/j.chom.2015.03.005.

Klaasen, H.L.B.M., Koopman, J.P., Vollaard, E.J., Theeuwes, A.G.M., Van Den Brink, M.E., Scholten, P.M., Bakker, M.H., and Beynen, A.C. (1991). Influence of Antimicrobial Drugs on Segmented Filamentous Bacteria in the Ileum of Mice. *Microbial Ecology in Health and Disease*. 4(6), 391-397. DOI: 10.3109/08910609109140155.

Knoop, K.A., McDonald, K.G., Kulkarni, D.H., and Newberry, R.D. (2016). Antibiotics promote inflammation through the translocation of native commensal colonic bacteria. *Gut*. 65, 1100-1109. DOI: 10.1136/gutjnl-2014-309059.

Koh, A.Y., Köhler, J.R., Coggshall, K.T., Van Rooijen, N., and Pier, G.B. (2008). Mucosal Damage and Neutropenia Are Required for *Candida albicans* Dissemination. PLoS Pathog. 4(2), e35. DOI: 10.1371/journal.ppat.0040035.

Kusu, T., Kayama, H., Kinoshita, M., Jeon, S.G., Ueda, Y., Goto, Y., Okumura, R., Saiga, H., Kurakawa, T., Ikeda, K., et al. (2013). Ecto-nucleoside triphosphate diphosphohydrolase 7 controls Th17 cell responses through regulation of luminal ATP in the small intestine. J Immunol. 190(2), 774-783. Published online 2012/12/18 DOI: 10.4049/jimmunol.1103067.

Levy, R., Okada, S., Beziat, V., Moriya, K., Liu, C., Chai, L.Y., Migaud, M., Hauck, F., Al Ali, A., Cyrus, C., et al. (2016). Genetic, immunological, and clinical features of patients with bacterial and fungal infections due to inherited IL-17RA deficiency. Proc Natl Acad Sci U S A. 113(51), E8277-E8285. DOI: 10.1073/pnas.1618300114.

Li, M.M., Li, C.Y., Wu, X.A., Chen, T.T., Ren, L., Xu, B.L., and Cao, J. (2020). Microbiota-driven interleukin-17 production provides immune protection against invasive candidiasis. Critical Care. 24(1), 268. DOI: 10.1186/s13054-020-02977-5.

Lionakis, M.S. (2014). New insights into innate immune control of systemic candidiasis. Med Mycol. 52(6), 555-564. DOI: 10.1093/mmy/myu029.

Lionakis, M.S., and Levitz, S.M. (2017). Host control of fungal infections: lessons from basic studies and human cohorts. Ann Rev Immunol. 36, 157-191. DOI: 10.1146/annurev-immunol-042617-053318.

Lionakis, M.S., Lim, J.K., Lee, C.C.R., and Murphy, P.M. (2011). Organ-specific innate immune responses in a mouse model of invasive candidiasis. J Innate Immun. 3(2), 180-199. DOI: 10.1159/000321157.

Martínez-López, M., Iborra, S., Conde-Garrosa, R., Mastrangelo, A., Danne, C., Mann, E.R., Reid, D.M., Gaboriau-Routhiau, V., Chaparro, M., Lorenzo, M.P., et al. (2019). Microbiota Sensing by Mincle-Syk Axis in Dendritic Cells Regulates Interleukin-17 and -22 Production and Promotes Intestinal Barrier Integrity. Immunity. 50(2), 446-461.e449. DOI: <https://doi.org/10.1016/j.immuni.2018.12.020>.

McMurdie, P.J., and Holmes, S. (2013). phyloseq: An R Package for Reproducible Interactive Analysis and Graphics of Microbiome Census Data. PLoS One. 8(4), e61217. DOI: 10.1371/journal.pone.0061217.

Molloy, M.J., Grainger, J.R., Bouladoux, N., Hand, T.W., Koo, L.Y., Naik, S., Quinones, M., Dzutsev, A.K., Gao, J.L., Trinchieri, G., et al. (2013). Intraluminal containment of commensal outgrowth in the gut during infection-induced dysbiosis. Cell Host Microbe. 14(3), 318-328. Published online 2013/09/17 DOI: 10.1016/j.chom.2013.08.003.

Mortha, A., Chudnovskiy, A., Hashimoto, D., Bogunovic, M., Spencer, S.P., Belkaid, Y., and Merad, M. (2014). Microbiota-Dependent Crosstalk Between Macrophages and ILC3 Promotes Intestinal Homeostasis. Science. 343(6178), 1249288. DOI: 10.1126/science.1249288.

Nakamura, Y.K., Metea, C., Karstens, L., Asquith, M., Gruner, H., Moscibrocki, C., Lee, I., Brislawn, C.J., Jansson, J.K., Rosenbaum, J.T., et al. (2016). Gut Microbial Alterations Associated With Protection From Autoimmune Uveitis. *Investigative Ophthalmology & Visual Science*. 57(8), 3747-3758. DOI: 10.1167/iovs.16-19733.

Odds, F.C., Davidson, A.D., Jacobsen, M.D., Tavanti, A., Whyte, J.A., Kibbler, C.C., Ellis, D.H., Maiden, M.C.J., Shaw, D.J., and Gow, N.A.R. (2006). *Candida albicans* strain maintenance, replacement, and microvariation demonstrated by multilocus sequence typing. *J Clin Microbiol*. 44(10), 3647-3658. DOI: 10.1128/jcm.00934-06.

Pappas, P.G., Lionakis, M.S., Arendrup, M.C., Ostrosky-Zeichner, L., and Kullberg, B.J. (2018). Invasive candidiasis. *Nat Rev Dis Primers*. 4, 18026. DOI: 10.1038/nrdp.2018.26.

Poynton, C.H., Barnes, R.A., and Rees, J. (1998). Interferon gamma and granulocyte-macrophage colony-stimulating factor for the treatment of hepatosplenic candidosis in patients with acute leukemia. *Clin Infect Dis*. 26(1), 239-240. Published online 1998/02/10 DOI: 10.1086/517077.

Price, A.E., Reinhardt, R.L., Liang, H.-E., and Locksley, R.M. (2012). Marking and quantifying IL-17A-producing cells in vivo. *PLoS One*. 7(6), e39750-e39750. Published online 2012/06/29 DOI: 10.1371/journal.pone.0039750.

Puel, A., Cypowyj, S., Bustamante, J., Wright, J.F., Liu, L., Lim, H.K., Migaud, M., Israel, L., Chrabieh, M., Audry, M., et al. (2011). Chronic Mucocutaneous Candidiasis in Humans with Inborn Errors of Interleukin-17 Immunity. *Science*. 332, 65-68. DOI: 10.1126/science.1200439.

**Puel, A., Daffinger, R., Natividad, A., Chrabieh, M., Barcenas-Morales, G., Picard, C., Cobat, A., Ouachace-Chardin, M., Toulon, A., Bustamante, J., et al. (2010).** Autoantibodies against IL-17A, IL-17F, and IL-22 in patients with chronic mucocutaneous candidiasis and autoimmune polyendocrine syndrome type I. *J Exp Med*. 207(2), 291-297. DOI: 10.1084/jem.20091983.

Quast, C., Pruesse, E., Yilmaz, P., Gerken, J., Schweer, T., Yarza, P., Peplies, J., and Glöckner, F.O. (2013). The SILVA ribosomal RNA gene database project: improved data processing and web-based tools. *Nucleic acids research*. 41(Database issue), D590-D596. Published online 2012/11/28 DOI: 10.1093/nar/gks1219.

Ricotta, E.E., Lai, Y.L., Babiker, A., Strich, J.R., Kadri, S.S., Lionakis, M.S., Prevots, D.R., and Adjemian, J. (2020). Invasive candidiasis species distribution and trends, United States, 2009-2017. *J Infect Dis*. jiaa502. Published online 2020/08/17 DOI: 10.1093/infdis/jiaa502.

Rókusz, L., Liptay, L., and Kádár, K. (2001). Successful treatment of chronic disseminated candidiasis with fluconazole and a granulocyte-macrophage colony-stimulating factor combination. *Scand J Infect Dis*. 33(10), 784-786. Published online 2001/12/01 DOI: 10.1080/003655401317074671.

Russo, A., Falcone, M., Fantoni, M., Murri, R., Masucci, L., Carfagna, P., Ghezzi, M.C., Posteraro, B., Sanguinetti, M., and Venditti, M. (2015). Risk factors and clinical outcomes of candidaemia in patients treated for *Clostridium difficile* infection. *Clin Microbiol Infect.* 21(5), 493.e491-494. Published online 2015/02/24 DOI: 10.1016/j.cmi.2014.12.024.

Shankar, J., Solis, N.V., Mounaud, S., Szpakowski, S., Liu, H., Losada, L., Niernan, W.C., and Filler, S.G. (2015). Using Bayesian modelling to investigate factors governing antibiotic-induced *Candida albicans* colonization of the GI tract. *Sci. Rep.* 5. DOI: 10.1038/srep08131  
<http://www.nature.com/srep/2015/150203/srep08131/abs/srep08131.html#supplementary-information>.

Shao, T.-Y., Ang, W.X.G., Jiang, T.T., Huang, F.S., Andersen, H., Kinder, J.M., Pham, G., Burg, A.R., Ruff, B., Gonzalez, T., et al. (2019). Commensal *Candida albicans* Positively Calibrates Systemic Th17 Immunological Responses. *Cell host & microbe.* 25(3), 404-417.e406. DOI: 10.1016/j.chom.2019.02.004.

Sipsas, N.V., Lewis, R.E., Tarrand, J., Hachem, R., Rolston, K.V., Raad, I.I., and Kontoyiannis, D.P. (2009). Candidemia in patients with hematologic malignancies in the era of new antifungal agents (2001-2007). *Cancer.* 115(20), 4745-4752. DOI: 10.1002/cncr.24507.

Smith, P.M., Howitt, M.R., Panikov, N., Michaud, M., Gallini, C.A., Bohlooly-Y, M., Glickman, J.N., and Garrett, W.S. (2013). The microbial metabolites, short chain fatty acids, regulate colonic Treg cell homeostasis. *Science (New York, N.Y.).* 341(6145), 10.1126/science.1241165. DOI: 10.1126/science.1241165.

Sonnenberg, G.F., Monticelli, L.A., Alenghat, T., Fung, T.C., Hutnick, N.A., Kunisawa, J., Shibata, N., Grunberg, S., Sinha, R., Zahm, A.M., et al. (2012). Innate Lymphoid Cells Promote Anatomical Containment of Lymphoid-Resident Commensal Bacteria. *Science.* 336(6086), 1321-1325. DOI: 10.1126/science.1222551.

Spellberg, B., Ibrahim, A.S., Jr, J.E.E., and Filler, S.G. (2005). Mice with Disseminated Candidiasis Die of Progressive Sepsis. *J. Infect. Dis.* 192(2), 336-343.

Spellberg, B., Johnston, D., Phan, Q.T., Edwards, J.E., French, S.W., Ibrahim, A.S., and Filler, S.G. (2003). Parenchymal organ, and not splenic, immunity correlates with host survival during disseminated candidiasis. *Infect Immun.* 71(10), 5756-5764. DOI: 10.1128/iai.71.10.5756-5764.2003.

Spiller, R.C. (2018). Hidden Dangers of Antibiotic Use: Increased Gut Permeability Mediated by Increased Pancreatic Proteases Reaching the Colon. *Cell Mol Gastroenterol Hepatol.* 6(3), 347-348.e341. DOI: 10.1016/j.jcmgh.2018.06.005.

Strollo, S., Lionakis, M.S., Adjemian, J., Steiner, C.A., and Prevots, D.R. (2016). Epidemiology of Hospitalizations Associated with Invasive Candidiasis, United States, 2002-2012(1). *Emerg Infect Dis.* 23(1), 7-13. Published online 2016/12/17 DOI: 10.3201/eid2301.161198.

Strzępa, A., Majewska-Szczepanik, M., Lobo, F.M., Wen, L., and Szczepanik, M. (2017). Broad spectrum antibiotic enrofloxacin modulates contact sensitivity through gut microbiota in a murine model. *Journal of Allergy and Clinical Immunology*. 140(1), 121-133.e123. DOI: 10.1016/j.jaci.2016.11.052.

Tulstrup, M.V.-L., Christensen, E.G., Carvalho, V., Linnings, C., Ahn, S., Højberg, O., Licht, T.R., and Bahl, M.I. (2015). Antibiotic Treatment Affects Intestinal Permeability and Gut Microbial Composition in Wistar Rats Dependent on Antibiotic Class. *PLoS One*. 10(12), e0144854-e0144854. DOI: 10.1371/journal.pone.0144854.

van de Veerdonk, F.L., Kullberg, B.J., and Netea, M.G. (2012). Adjunctive immunotherapy with recombinant cytokines for the treatment of disseminated candidiasis. *Clin Microbiol Infect*. 18(2), 112-119. DOI: <https://doi.org/10.1111/j.1469-0691.2011.03676.x>.

van Opstal, E., Kolling, G.L., Moore, J.H., 2nd, Coquery, C.M., Wade, N.S., Loo, W.M., Bolick, D.T., Shin, J.H., Erickson, L.D., and Warren, C.A. (2016). Vancomycin Treatment Alters Humoral Immunity and Intestinal Microbiota in an Aged Mouse Model of *Clostridium difficile* Infection. *J Infect Dis*. 214(1), 130-139. Published online 2016/02/24 DOI: 10.1093/infdis/jiw071.

Vautier, S., Drummond, R.A., Redelinguys, P., Murray, G.I., MacCallum, D.M., and Brown, G.D. (2012). Dectin-1 is not required for controlling *Candida albicans* colonization of the gastrointestinal tract. *Infect Immun*. 80(12), 4216-4222. DOI: 10.1128/iai.00559-12.

Von Drygalski, A., Curtis, B.R., Bougie, D.W., McFarland, J.G., Ahl, S., Limbu, I., Baker, K.R., and Aster, R.H. (2007). Vancomycin-Induced Immune Thrombocytopenia. *N. Eng. J. Med*. 356(9), 904-910. DOI: 10.1056/NEJMoa065066.

Weber, N., Liou, D., Dommer, J., MacMenamin, P., Quiñones, M., Misner, I., Oler, A.J., Wan, J., Kim, L., Coakley McCarthy, M., et al. (2018). Nephele: a cloud platform for simplified, standardized and reproducible microbiome data analysis. *Bioinformatics*. 34(8), 1411-1413. Published online 2017/10/14 DOI: 10.1093/bioinformatics/btx617.

Yang, J.H., Bhargava, P., McCloskey, D., Mao, N., Palsson, B.O., and Collins, J.J. (2017). Antibiotic-Induced Changes to the Host Metabolic Environment Inhibit Drug Efficacy and Alter Immune Function. *Cell Host Microbe*. 22(6), 757-765.e753. DOI: <https://doi.org/10.1016/j.chom.2017.10.020>.

Yu, L.C.-H., Shih, Y.-A., Wu, L.-L., Lin, Y.-D., Kuo, W.-T., Peng, W.-H., Lu, K.-S., Wei, S.-C., Turner, J.R., and Ni, Y.-H. (2014). Enteric dysbiosis promotes antibiotic-resistant bacterial infection: systemic dissemination of resistant and commensal bacteria through epithelial transcytosis. *Am J Physiol Gastrointest Liver Physiol*. 307(8), G824-G835. Published online 2014/07/24 DOI: 10.1152/ajpgi.00070.2014.

Zhai, B., Ola, M., Rolling, T., Tosini, N.L., Joshowitz, S., Littmann, E.R., Amoretti, L.A., Fontana, E., Wright, R.J., Miranda, E., et al. (2020). High-resolution mycobiota analysis reveals dynamic intestinal translocation preceding invasive candidiasis. *Nat Med*. 26(1), 59-64. Published online 2020/01/08 DOI: 10.1038/s41591-019-0709-7.

