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## Fetal and early postnatal lead exposure measured in teeth associates with infant gut microbiota



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### ABSTRACT

**Background:** Lead (Pb) is an environmentally ubiquitous heavy metal associated with a wide range of adverse health effects in children. Both lead exposure and the early life microbiome—which plays a critical role in human development—have been linked to similar health outcomes, but it is unclear if the adverse effects of lead are partially driven by early life gut microbiota dysbiosis. The objective of this study was to examine the association between *in utero* and postnatal lead levels (measured in deciduous baby teeth) and early life bacterial and fungal gut microbiota in the first year of life.

**Methods:** Data from the Wayne County Health, Environment, Allergy and Asthma Longitudinal Study (WHEALS) birth cohort were analyzed. Tooth lead levels during the 2nd and 3rd trimesters and postnatally (< 1 year of age) were quantified using high-resolution microspatial mapping of dentin growth rings. Early life microbiota were measured in stool samples collected at approximately 1 and 6 months of age, using both 16S rRNA (bacterial) and ITS2 (fungal) sequencing. Of the 1,258 maternal-child pairs in WHEALS, 146 had data on both tooth metals and early life microbiome.

**Results:** *In utero* tooth lead levels were significantly associated with gut fungal community composition at 1-month of age, where higher levels of 2nd trimester tooth lead was associated with lower abundances of *Candida* and *Aspergillus* and higher abundances of *Malassezia* and *Saccharomyces*; 3rd trimester lead was also associated with lower abundances of *Candida*. Though lead did not significantly associate with the overall structure of the infant gut bacterial community, it associated with the abundance of some specific bacterial taxa, including the increased abundance of *Collinsella* and *Bilophila* and a decreased abundance of *Bacteroides* taxa.

**Conclusions:** The observed associations between lead exposure and infant gut microbiota could play a role in the impact of lead on childhood development. Given the paucity of research examining these associations in humans—particularly for fungal microbiota—further investigation is needed.

### 1. Introduction

The human gut microbiome can be considered an ancillary organ

capable of directly influencing host health status (Evans et al., 2013). It produces a vast range of bioactive metabolites that affect metabolic function and is involved in both energy harvest and storage, as well as

**Abbreviations:** Pb, Lead; WHEALS, Wayne County Health, Environment, Allergy and Asthma Longitudinal Study; HFHS, Henry Ford Health System; ITS, Internal Transcribed Spacer Region; OTU, Operational Taxonomic Unit; BMI, Body Mass Index; ETS, Environmental Tobacco Smoke; ICC, Intraclass Correlation Coefficient; PERMANOVA, Permutational Multivariate Analysis of Variance; FDR, False Discovery Rate; ASD, Autism Spectrum Disorder

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immune system development (Clemente et al., 2012). Early life gut microbiota in particular are thought to affect health later in life, with dysbiosis associated with an increased risk of allergy and asthma (Johnson and Ownby 2017), obesity (Koleva et al., 2015), and neurodevelopmental outcomes (Sordillo et al., 2019). The human microbiome poses a unique research challenge because, unlike many environmental factors that are analyzed as external exposures, the microbiome exists within us, and thus the human physiology serves as the environment for the microbiota. However, with the advent of high-throughput approaches, we have only recently begun to appreciate the role that microbiota play and have perhaps only scratched the surface. Human microbiomes are dynamic and complex, and studies examining how common epidemiological factors relate to the infant gut microbiota only explain a small percentage of its variability, even when considered jointly (Levin et al., 2016). Therefore, taking into account environmental exposures may help to explain additional variability and improve prediction of infant gut microbiota assembly. In particular, the gut microbiome is influenced by environmental toxicants (Adamovsky et al., 2018). Environmental xenobiotic agents may act directly or indirectly on the microbiome to influence its composition; the gut microbiome may also modulate the influence of a toxic exposure by altering the pharmacokinetics of chemicals (National Academies of Sciences Engineering and Medicine, 2018). In short, there is growing recognition of the need to study the role of environmental toxicants on the microbiome (National Academies of Sciences Engineering and Medicine, 2018).

Lead (Pb) is considered the most important toxic heavy element and is ubiquitous in the environment, with potential sources of exposure including food, water, air, dust, soil, and lead-based paint (Wani et al., 2015). Upon exposure, Pb bioaccumulates throughout the human body and therefore has been implicated in a wide range of adverse health effects, including neurological, renal, immunological, and developmental (Abadin et al., 2019). There is no safe blood Pb level according to the Centers for Disease Control and Prevention (Betts, 2012), particularly for children, in which the toxicity of lead has the largest impact due to high soft tissue absorption (Wani et al., 2015). Both Pb exposure and microbiome composition have been linked to similar health outcomes in humans, including neurological (Abadin et al., 2019; Chu et al., 2019) and immune function (Abadin et al., 2019; Gao et al., 2018), and the microbiome may play a role in mediating the association between lead exposure and health outcomes. Therefore, it is important to understand how lead exposure impacts microbiota development in the infant gut.

In addition to lead exposure in children, it is not yet clear how prenatal exposure to lead may influence the development and trajectory of the human gut microbiome. In a study using a mouse model (Wu et al., 2016), pregnant mice were dosed with lead through drinking water, and their offspring showed altered gut microbiomes and body weight in adulthood, compared to controls. Potential mechanisms of these effects could be through preprogramming of the immune system or by altering offspring microbiome through microbial vertical transmission (either before, during, or after birth). However, it is unknown how prenatal exposures to lead or other environmental toxicants may influence the gut microbiome in humans. This analysis is the first to examine this relationship, focusing on associations between prenatal (2nd and 3rd trimester) and early postnatal (birth to approximately 1 year of age) lead exposure and the human gut microbiome in infancy (bacterial and fungal microbiota at 1 and 6 months of age). Lead levels were measured in naturally shed deciduous teeth, which uniquely allows for retrospective quantification of fetal and early life lead levels with precise windows of exposure.

## 2. Methods

### 2.1. Study information

Analyses were based on data from the Wayne County Health, Environment, Allergy and Asthma Longitudinal Study (WHEALS), a racially-diverse unselected birth cohort of 1,258 maternal-child pairs based in Detroit, Michigan (Wegienka et al., 2015, 2016). WHEALS recruited pregnant women with due dates from September 2003 through December 2007, and who were seeing a Henry Ford Health System (HFHS) obstetrics practitioner at one of five clinics to establish an unselected birth cohort (Cassidy-Bushrow et al., 2012; Havstad et al., 2011; Wegienka et al., 2011). All women were in their second trimester or later, were aged 21–49 years, and were living in a pre-defined geographic area in Wayne and Oakland counties of Michigan that included the city of Detroit as well as the suburban areas immediately surrounding the city. All participants provided written, informed consent. Study protocols were approved by the HFHS Institutional Review Board.

Details of the Tooth Fairy Study, a sub-study of the WHEALS cohort, have previously been reported (Cassidy-Bushrow et al., 2017, 2019). Briefly, advertisements for the WHEALS Tooth Fairy Study were placed in study newsletters, a save-the-tooth refrigerator magnet was sent in a holiday mailer to all participants, and families were asked if they wanted to donate a tooth for the study during planned recruitment phone calls for other WHEALS activities. A total of 512 teeth were received from 203 participants between December 2011 and September 2019. Teeth were selected for metal measurement if (1) the child had at least some outcome data available (birth outcomes and/or a 2-year clinic visit) or early life microbiome data; and (2) the tooth sample met laboratory quality control/quality assurance guidelines. A total of 180 participants had tooth metals quantified.

### 2.2. Stool specimens

Home visits with participants were conducted targeting infant ages 1 and 6 months. Families were asked to retain the most recent soiled diaper prior to the home visit and stool samples were banked at  $-80^{\circ}\text{C}$ . Detailed information on DNA extraction methods are presented elsewhere (Fujimura et al., 2016).

### 2.3. Polymerase chain reaction conditions and library preparation for sequencing

The V4 region of the 16S rRNA gene was amplified, as described in detail elsewhere (Caporaso et al., 2012). Briefly, 16S rRNA amplification was performed in 25- $\mu\text{L}$  reactions using 0.025 U Takara Hot Start ExTaq (Takara Mirus Bio Inc., Madison, WI), 1X Takara buffer with  $\text{MgCl}_2$ , 0.4 pmol/ $\mu\text{L}$  of F515 and R806 primers, 0.56 mg/mL of bovine serum albumin (Roche Applied Science, Indianapolis, IN), 200  $\mu\text{M}$  of dNTPs and 10 ng of genomic DNA. Reactions were performed in triplicate with the following: initial denaturation ( $98^{\circ}\text{C}$ , 2 min), 30 cycles of  $98^{\circ}\text{C}$  (20 sec), annealing at  $50^{\circ}\text{C}$  (30 sec), extension at  $72^{\circ}\text{C}$  (45 sec), and final extension at  $72^{\circ}\text{C}$  (10 min). Amplicons were verified using a 2% Tris/Borate/EDTA agarose e-gel (Life Technologies, Grand Island, NY), cleaned and normalized using SequalPrep Normalization Plates (Applied Biosystems, Foster City, CA), and further quantified using the Qubit 2.0 Fluorometer and the double-stranded DNA HS Assay Kit (Life Technologies). Samples were pooled in equal moles at concentrations of 5 ng, purified using AMPure SPRI beads (Beckman Coulter, Brea, CA), denatured and diluted to 2 nM, and 5 pM was loaded onto the Illumina Nextseq cartridge with 40% (v/v) of denatured 12.5 pM PhiX spike-in control.

The internal transcribed spacer region (ITS) 2 of the rRNA gene was amplified using the primer pair ITS7 (5'-GTGARTCATCGAATCT-TTG-3') and ITS4 (5'-TCCTCCGCTTATT GATATGC-3'). PCR reactions

were performed in triplicate in 25  $\mu$ l reaction with 1X Takara buffer (Takara Mirus Bio), 200 nM of each primer, 200  $\mu$ M dNTPs, 2.75 mM of MgCl<sub>2</sub>, 0.56 mg ml<sup>-1</sup> of BSA (Roche Applied Science), 0.025 U Takara Hot Start ExTaq and 50 ng of gDNA. Reactions were conducted under the following conditions: initial denaturation (94 °C for 5 min) followed by 30 cycles of 94 °C (30 sec), annealing at 54 °C (30 sec), extension at 72 °C (30 sec) and a final extension at 72 °C (7 min). PCR verification and purification were performed as described above for bacterial library preparation. Samples were quantified using KAPA SYBR (KAPA Biosystems, Wilmington, MA) qPCR as recommended by the manufacturers. Samples were then pooled in equal moles at 5 ng, purified, denatured, and diluted similar to the 16S amplicon library described above. 10 pM of the ITS2 amplicon library was loaded onto the Illumina MiSeq cartridge with 25% (v/v) of denatured 10 pM PhiX spike-in control.

#### 2.4. Sequence data processing and quality control

Bacterial paired-end sequences were assembled using FLASH v1.2.7 (Magoč and Salzberg 2011) requiring a minimum base pair overlap of 25 bp and demultiplexed by barcode using QIIME v1.8 (Caporaso et al., 2010b). Quality filtering was performed using USEARCH v8.0.1623 (Edgar, 2010) to remove reads with > 2 expected errors. Quality reads were dereplicated at 100% sequence identity, clustered at 97% sequence identity into operational taxonomic units (OTUs), filtered of chimeric sequences by UCHIME (Edgar et al., 2011), and mapped back to resulting OTUs using UPARSE (Edgar, 2013); sequence reads that failed to cluster with a reference sequence were clustered *de novo*. Taxonomy was assigned to the OTUs using the Greengenes database v13.5 (McDonald et al., 2012). Sequences were aligned using PyNASt (Caporaso et al., 2010a), and FastTree 2.1.3 (Price et al., 2009) was used to build a phylogenetic tree. A rarefaction depth of 60,000 reads per sample was selected by using rarefaction curves, based on the plateau of reads and maximizing sample inclusion. Note that those included versus excluded from the current analysis did not have significantly different total read depths ( $p = 0.42$ ), making it unlikely that a different rarefaction depth would be more appropriate if only done within the analysis subset. Additionally, rather than selecting a single rarefaction table, we multiply rarefied the data in order to ensure a representative community was selected for each sample (Fujimura et al., 2016). Briefly, multiple rarefied OTU tables are calculated, and for each sample, the distance between the sample-specific rarefied vectors calculated. The rarefied vector that is the minimum average distance from itself to all other rarefied vectors is selected, as it is considered the most representative for that sample.

Fungal sequences were quality trimmed (Q score, < 25) and removed of adaptor sequences using cutadapt (Martin, 2011). Paired-end sequences were assembled, demultiplexed by barcode, clustered into OTUs at 97% identity and filtered of chimeras using similar methods as described for 16S amplicons. Taxonomy was assigned using UNITE v7.0 (Koljalg et al., 2013). Resulting sequence reads were normalized by again multiply rarefying to 1,000 reads per sample as described previously to assure representative reduced data. Sequencing data was uploaded to the NCBI Sequence Read Archive (SRA) BioProject PRJNA648818 under the accession number SUB7839619.

A total of 580 children had at least 1 stool sample in the final rarefied OTU table; of these, 146 (25%) had tooth metal levels quantified, providing the final analysis subset sample size (Supplemental Fig. 1). Of these 146 children, 43 had 1-month bacterial microbiota only, 35 had 6-month bacterial microbiota only, and 68 had bacterial microbiota at both time points. Sample sizes were smaller for fungal microbiota: as is common in early life cohorts, many samples failed to produce an ITS2 amplicon (53%), with 33 children having 1-month fungal microbiota only, 29 having 6-month fungal microbiota only, and 19 having it at both time points. In the analytical dataset, stool specimens from the 1-month visit were collected at a mean  $\pm$  SD of 38  $\pm$  16 days

(minimum = 16, maximum = 107) and stool specimens from the 6-month visit were collected at a mean  $\pm$  SD of 205  $\pm$  27 days (minimum = 172, maximum = 290). Throughout, “1 month” and “6 month” are used as labels of the targeted time period of sample collection.

#### 2.5. Covariate Measurement

Maternal race, household income, marital status, exposure to environmental tobacco smoke (ETS), smoking during pregnancy, year home was built, and exposure to indoor pets prenatally were self-reported. Year residence was built was dichotomized as 1980 or after or before 1980, to indicate risk of lead exposure due to lead-based paint (Dixon et al., 2009). Address during pregnancy was recorded and used to define whether the fetus was mainly exposed to an urban residence (defined as within the confines of the city of Detroit) or a suburban residence. Prenatal and delivery records for WHEALS women and children were abstracted to obtain body mass index (BMI) at the first prenatal care visit, delivery type, gestational age at delivery, and birthweight. Breastfeeding status was determined via questionnaires at the 1-month and 6-month study visits.

#### 2.6. Analysis of lead in tooth samples

We directly measured lead in baby teeth using laser ablation-inductively coupled plasma-mass spectrometry (LA-ICP-MS) and assigned developmental times as detailed elsewhere (Arora et al., 2014, 2012). Teeth were sectioned and the neonatal line (a histological feature formed in enamel and dentine at the time of birth) and incremental markings were used to assign temporal information to sampling points. Second trimester, third trimester, postnatal (birth through 1-year), and childhood (age 1 to tooth shedding) lead levels were distinguished using previously validated methods that rely on incremental markings in teeth, akin to growth rings in trees (Arora et al., 2014, 2012). The childhood time point was excluded from this analysis because exposure time was after microbiota quantification. We used an ArF excimer laser ablation system (ESI, USA) attached to an Agilent Technologies 8800 triple-quad ICP-MS. Data were analysed as metal-to-calcium ratios (e.g. <sup>208</sup>Pb:<sup>43</sup>Ca) to control for variations in mineral content within a tooth and between samples. Samples were analysed in three batches. Tooth attrition, which is the amount of tooth lost due to grinding or wear, was also measured; teeth with excessive attrition that would impede the chemical analysis are excluded. National Institute of Standards and Technology SRM 612 was used for calibration and quality control. The detection limit was 0.05  $\mu$ g/g for lead. Across all time points, approximately 2–7% of lead measurements were below the detection limit; values below the detection limit are excluded from statistical analysis.

A small number of children ( $N = 17$ ) had two teeth analyzed for quality control procedures; in these cases, metals levels were averaged over teeth within each child. We have previously shown moderate to excellent agreement for lead concentrations between teeth from the same child (second trimester intraclass correlation coefficients [ICC] = 0.55; third trimester ICC = 0.74; postnatal ICC = 0.87) (Cassidy-Bushrow et al., 2017).

#### 2.7. Statistical analysis

Tooth lead levels were tested for differences by maternal and child characteristics using the Kruskal-Wallis test for categorical covariates and Spearman correlations for continuous covariates. For these tests, tooth lead levels were shown as metal-to-calcium ratios; for further modelling, lead levels were divided by the standard deviation to represent a one standard deviation increase. Additionally, all tests of the relationship between lead levels and infant gut microbiota were adjusted for tooth type, tooth attrition, tooth batch, exact age at stool

sample collection, and child race, for which we have previously shown associations with tooth lead levels (Cassidy-Bushrow et al., 2017) and infant gut microbiota (Levin et al., 2016) in this cohort. Note that sex-specific effects were not explored. In addition to the relatively small sample size, while there is clear evidence of a sex-specific effect of lead on health (Polanska et al., 2018; Wu et al., 2016), there is a lack of evidence of a sex-specific effect of metals on the gut microbiota (Wu et al., 2016).

Infant gut alpha diversity metrics (bacterial: richness, Pielou's evenness, Faith's phylogenetic diversity, Shannon's diversity; fungal: richness, Pielou's evenness, Shannon's diversity) were tested for differences in lead levels using covariate-adjusted linear regression models. The association between fungal amplification failure and lead levels was modelled using covariate-adjusted logistic regression models. Adjusted permutational multivariate analysis of variance (PERMANOVA) was used to test for infant gut compositional differences by lead levels (Anderson, 2001) using the R package *vegan* (Oksanen et al., 2019). In order to capture different features of community composition, both phylogenetic [bacterial only: unweighted and weighted UniFrac (Lozupone et al., 2011)] and non-phylogenetic [bacterial and fungal: Bray-Curtis, Canberra] distances were used in PERMANOVA tests. Principal coordinates analysis (PCoA) was performed on these distances using the R package *labdsv* (Roberts, 2019); plots of these distances are provided in Supplemental Fig. 2.

Tests of differential genera and OTU abundance were performed using covariate-adjusted zero-inflated negative binomial regression using the R package *pscl* (Zeileis et al., 2008); in cases where zero-inflated models failed to converge, the standard negative binomial was implemented as a secondary modelling strategy using the R package *MASS* (Venables and Ripley, 2002). Only genera and OTUs found in 10% or more of samples were tested. Statistical significance was assessed after accounting for multiple testing using the Benjamini & Hochberg false discovery rate (FDR) adjustment (Benjamini and Hochberg, 1995), with  $p_{FDR} < 0.05$  considered significant. Analyses were performed using SAS 9.4 and R 3.6.1. Covariate and lead data are provided in Supplemental Data 1.

## 3. Results

### 3.1. Factors associated with lead exposure

In order to describe the data and assess potential confounders for the main association of interest (i.e., between lead levels and early life microbiota), lead levels were first compared across a variety of maternal and child characteristics (Table 1). Location of residence, prenatal indoor pets, birthweight z-score, and child race were all significantly associated with lead levels. Specifically, black children (all  $p < 0.001$ ) and children living in urban residences had significantly higher lead levels at all three time points ( $p = 0.024, 0.005, 0.003$ , respectively). Children whose mothers had a pet during pregnancy had significantly lower lead levels, but only during the 3rd trimester ( $p = 0.019$ ). Higher lead levels at all three time points were also associated with lower birthweight z-scores ( $p = 0.021, 0.019, 0.036$ , respectively). After adjusting for race, however, associations of tooth lead level with location of residence and prenatal pets was diminished and no longer significant (all  $p \geq 0.067$ ). Of note, no associations were observed between lead levels and mode of delivery or breastfeeding status, which we have previously shown have a large impact on infant gut microbiota in these children (Levin et al., 2016).

### 3.2. Association between lead exposure and bacterial gut microbiota

We next evaluated the association between lead levels and infant gut bacterial alpha diversity metrics (richness, evenness, phylogenetic/Shannon's diversity), a high-level summary statistic of microbiota structure. After adjusting for tooth type, attrition, batch, exact age at

stool sample collection, and child race, *in utero* and postnatal lead levels did not significantly associate with bacterial alpha diversity, at both 1 and 6 months of age (Table 2). When microbiota beta diversity (microbiota composition) was considered, lead did not significantly explain the overall variability in bacterial microbiota composition, after adjusting for covariates (Table 3). Though lead exposure did not appear to profoundly influence the structure or composition of the early life bacterial gut microbiome, it may nevertheless relate to the presence and abundance of specific bacterial taxa. Therefore, we also performed genus (Fig. 1) and OTU (Supplemental Table 1) tests of differences by tooth lead levels. Of note, at the genus level, 2nd and 3rd trimester lead levels were positively associated with *Collinsella* abundance at 1-month of age ( $p_{FDR} < 0.001, 0.022$ , respectively), as well as *Bilophila* abundance at 6-months of age ( $p_{FDR} = 0.023, 0.008$ , respectively) (Fig. 1). At the OTU level, the positive association between lead levels and *Collinsella* abundance appeared to be driven by four *Collinsella aerofaciens* OTUs; lead levels also positively associated with two *Bilophila* OTUs, but these did not have classification available beyond genus (Supplemental Table 1). Additionally, *in utero* and postnatal lead levels negatively associated with several *Bacteroides* OTUs, at both 1 and 6 months of age (Supplemental Table 1).

### 3.3. Association between lead exposure and fungal gut microbiota

After covariate adjustment, lead levels did not significantly associate with fungal alpha diversity, at both 1 and 6 months of age (Table 2). In tests of the association between tooth lead levels and fungal amplification failure, no associations were detected after covariate adjustment (all  $p \geq 0.16$ ). However, 2nd trimester and 3rd trimester lead levels significantly associated with fungal community composition at 1-month of age (Table 3; Bray-Curtis  $p$ -value = 0.048, 0.049, respectively), with the association explaining approximately 4% of the variability in community composition.

In tests of differential fungal genera abundance by tooth lead levels (after covariate adjustment and FDR correction), *in utero* and postnatal lead levels significantly associated with a total of five genera (Fig. 1): *Candida*, *Malassezia*, *Penicillium*, *Saccharomyces*, and *Aspergillus*. Specifically, higher tooth lead levels in the 2nd and 3rd trimesters were associated with significantly lower *Candida* abundances at 1-month of age ( $p_{FDR} = 0.006, 0.003$ , respectively); 2nd trimester and postnatal lead levels also negatively correlated with *Aspergillus* abundance at 1-month of age (Fig. 1;  $p_{FDR} = 0.002, < 0.001$ , respectively). Additionally, postnatal lead levels negatively correlated with *Penicillium* abundance at 6-months of age (Fig. 1;  $p_{FDR} = 0.002$ ). In contrast, higher tooth lead levels in the 2nd trimester and postnatally were associated with significantly higher *Malassezia* abundances at both 1-month (Fig. 1;  $p_{FDR} = 0.002, 0.013$ , respectively) and 6-months of age (Fig. 1; both  $p_{FDR} < 0.001$ ). Additionally, higher tooth lead levels in the 2nd trimester was associated with significantly higher *Saccharomyces* abundance at 1-month of age ( $p_{FDR} = 0.006$ ), but significantly lower *Saccharomyces* abundance at 6-months of age (Fig. 1;  $p_{FDR} = 0.006$ ). Tests of specific fungal OTUs (Supplemental Table 1) suggested that the positive association between lead and *Malassezia* is primarily driven by *Malassezia restricta* and *Malassezia globosa* OTUs, whereas the negative association with *Candida* appears to be primarily driven by a *Candida parapsilosis* OTU.

## 4. Discussion

Our study identified an association between tooth lead levels and overall gut fungal community composition at 1-month of age, where lead levels were negatively correlated with the abundance of *Candida parapsilosis*, which has been shown to be sensitive to lead (Bansal et al., 2019), and positively correlated with the abundance of *Malassezia restricta* and *M. globosa* as well as *Saccharomyces* species, which has been shown to be lead-resistant (Jaroslawska and Piotrowska-Seget, 2014).

**Table 1**  
Maternal and child characteristics by *in utero* and postnatal lead levels (N = 146).

Variable	Level	2nd Trimester Lead <sup>a</sup>			3rd Trimester Lead <sup>a</sup>			Postnatal Lead <sup>a</sup>		
		N	Median (IQR) or $\rho$	p-value <sup>b</sup>	N	Median (IQR) or $\rho$	p-value <sup>b</sup>	N	Median (IQR) or $\rho$	p-value <sup>b</sup>
Maternal education	≤ HS diploma	13	0.024 (0.018)	0.26	13	0.026 (0.013)	0.087	13	0.023 (0.026)	0.077
	Some college	61	0.027 (0.037)		64	0.033 (0.047)		65	0.030 (0.029)	
	≥ Bachelor's Degree	62	0.022 (0.026)		65	0.031 (0.034)		65	0.024 (0.030)	
Household income	< \$20,000	9	0.024 (0.011)	0.98	9	0.033 (0.030)	0.89	9	0.027 (0.021)	0.49
	\$20,000- < \$40,000	24	0.026 (0.032)		25	0.032 (0.033)		26	0.028 (0.031)	
	\$40,000- < \$80,000	40	0.029 (0.024)		44	0.035 (0.040)		45	0.032 (0.036)	
	\$80,000- < \$100,000	25	0.021 (0.030)		25	0.027 (0.041)		24	0.022 (0.024)	
	≥ \$100,000	23	0.021 (0.028)		24	0.032 (0.034)		24	0.025 (0.028)	
Married	Refused to Answer	15	0.020 (0.053)	0.089	15	0.028 (0.045)	0.14	15	0.022 (0.046)	0.063
	No	36	0.028 (0.051)		38	0.034 (0.066)		39	0.034 (0.040)	
Location of residence	Yes	100	0.023 (0.026)	0.024	104	0.030 (0.033)	0.005	104	0.024 (0.026)	0.003
	Suburban	85	0.021 (0.026)		88	0.027 (0.030)		89	0.022 (0.028)	
	Urban	51	0.028 (0.039)		54	0.041 (0.056)		54	0.033 (0.027)	
Maternal age at birth	—	136	0.056	0.52	142	0.094	0.27	143	0.054	0.52
Maternal BMI at first prenatal visit	—	135	-0.014	0.87	141	0.031	0.72	142	0.064	0.45
Mother Atopic	No	62	0.025 (0.037)	0.79	64	0.032 (0.047)	0.69	66	0.024 (0.035)	0.55
	Yes	71	0.024 (0.026)		75	0.030 (0.033)		74	0.029 (0.023)	
Prenatal antibiotic use	No	54	0.023 (0.027)	0.37	57	0.033 (0.040)	0.74	57	0.024 (0.027)	0.16
	Yes	77	0.025 (0.032)		78	0.031 (0.044)		79	0.029 (0.032)	
Prenatal antifungal use	No	108	0.026 (0.029)	0.20	112	0.033 (0.045)	0.18	113	0.029 (0.031)	0.097
	Yes	23	0.024 (0.030)		23	0.026 (0.025)		23	0.021 (0.022)	
Mother smoked during pregnancy	No	127	0.024 (0.028)	0.33	133	0.031 (0.038)	0.41	134	0.027 (0.028)	0.84
	Yes	9	0.035 (0.045)		9	0.042 (0.075)		9	0.024 (0.033)	
ETS during pregnancy	No	107	0.024 (0.026)	0.13	112	0.030 (0.037)	0.18	112	0.027 (0.028)	0.34
	Yes	29	0.035 (0.044)		30	0.039 (0.059)		31	0.027 (0.033)	
Prenatal indoor pets	No	76	0.029 (0.042)	0.058	80	0.038 (0.048)	0.019	82	0.028 (0.030)	0.19
	Yes	60	0.021 (0.020)		62	0.026 (0.024)		61	0.024 (0.027)	
Year House was Built	1980 or after	26	0.023 (0.030)	0.40	28	0.026 (0.036)	0.16	28	0.022 (0.024)	0.099
	Before 1980	104	0.024 (0.028)		108	0.032 (0.038)		109	0.027 (0.031)	
First born child	No	79	0.025 (0.019)	0.90	83	0.032 (0.032)	0.64	85	0.027 (0.027)	0.81
	Yes	57	0.024 (0.033)		59	0.030 (0.051)		58	0.027 (0.031)	
Season of birth	Winter	29	0.022 (0.022)	0.52	30	0.029 (0.035)	0.65	31	0.029 (0.033)	0.88
	Spring	34	0.029 (0.048)		36	0.034 (0.058)		37	0.025 (0.039)	
	Summer	35	0.021 (0.028)		37	0.030 (0.032)		37	0.024 (0.026)	
	Fall	38	0.026 (0.032)		39	0.032 (0.045)		38	0.029 (0.021)	
Mode of delivery	Vaginal	83	0.023 (0.026)	0.31	87	0.029 (0.034)	0.29	88	0.026 (0.026)	0.49
	C-Section	52	0.028 (0.033)		54	0.034 (0.045)		54	0.028 (0.030)	
Gestational age at birth	—	133	0.013	0.88	142	-0.023	0.79	139	0.01	0.91
Birthweight z-score	—	124	-0.208	0.021	129	-0.207	0.019	131	-0.184	0.036
Child race	White	50	0.018 (0.018)	< 0.001	52	0.022 (0.025)	< 0.001	52	0.020 (0.019)	< 0.001
	Black	56	0.031 (0.050)		58	0.044 (0.061)		59	0.032 (0.033)	
	Other/Mixed	30	0.024 (0.022)		32	0.031 (0.041)		32	0.031 (0.032)	
Child sex	Male	66	0.029 (0.026)	0.072	69	0.036 (0.039)	0.065	70	0.029 (0.020)	0.33
	Female	70	0.021 (0.026)		73	0.027 (0.035)		73	0.024 (0.036)	
Breastfeeding status at 1-month	Formula Feeding	25	0.032 (0.030)	0.32	26	0.042 (0.048)	0.58	26	0.028 (0.033)	0.60
	Mixed Feeding	81	0.023 (0.028)		85	0.031 (0.049)		87	0.027 (0.029)	
	Exclusive Breastfeed	30	0.029 (0.025)		31	0.029 (0.023)		30	0.026 (0.031)	
Breastfeeding status at 6-months	Formula Feeding	25	0.032 (0.030)	0.38	26	0.042 (0.048)	0.58	26	0.028 (0.033)	0.93
	Mixed Feeding	95	0.024 (0.027)		99	0.031 (0.042)		100	0.027 (0.026)	
	Exclusive Breastfeed	16	0.024 (0.023)		17	0.026 (0.029)		17	0.025 (0.033)	

<sup>a</sup> Tooth lead levels shown as metal-to-calcium ratios.

<sup>b</sup> Kruskal-Wallis or Spearman correlation p-value.

Interestingly, compositional differences were only found for *in utero* but not postnatal lead levels, highlighting the advantage of capturing multiple exposure time points both before and after birth using novel tooth-matrix biomarkers. In contrast, we did not detect an association between *in utero* or early life tooth lead levels and overall infant gut bacterial community composition, though specific bacterial taxa including *Collinsella*, *Bilophila*, and *Bacteroides* were identified after covariate adjustment and FDR correction.

Several animal studies have shown that lead is associated with the bacterial gut microbiome. In female C57BL/6 mice (~7 weeks old), those exposed to lead in their drinking water (10 ppm) for 13 weeks had significantly different alpha and beta diversity trajectories of the gut microbiome compared to control mice (Gao et al., 2017). Mice (~6 week old female Balb/C) treated for 8 weeks with cadmium or lead had statistically significant lower levels of *Lachnospiraceae* and higher

numbers of *Lactobacillaceae* and *Erysipelotrichaceae* in their gut than control animals. Finally, Wu et al., examined the impact of maternal lead exposure in drinking water (exposed pre-conception through lactation) on the gut microbiome of the adult offspring of non-agouti wild-type mice (Wu et al., 2016). While there were no differences in the richness or diversity of the exposed offspring compared to control mice, *Bacteroidetes* were significantly reduced while *Firmicutes* and *Desulfobriviales* were significantly increased in the mice exposed to lead through their mothers (Wu et al., 2016). Consistent with these findings, we also observed a significant reduction of specific *Bacteroides* OTUs as well as a significant increase in *Bilophila* (which is of the order *Desulfobriviales*) with higher levels of lead exposure; of note, *Desulfobrivium desulfuricans* has been shown to be resistant to Pb (Sani et al., 2003).

In addition to animal models, a small number of studies have

**Table 2**  
Association between tooth lead levels and alpha diversity metrics.

Time of Metal Measurement	Outcome	Time of Stool Measurement	Bacterial			Fungal		
			N	$\beta$ (SE) <sup>b</sup>	p-value	N	$\beta$ (SE) <sup>b</sup>	p-value
2nd trimester	Richness	1-Month	104	5.53 (6.98)	0.43	48	0.29 (1.65)	0.86
		6-Months	96	-7.77 (7.31)	0.29	45	1.7 (1.51)	0.27
	Evenness	1-Month	104	0 (0.01)	0.97	37	0.03 (0.05)	0.56
		6-Months	96	-0.02 (0.01)	0.092	40	-0.02 (0.05)	0.73
	Faith's Diversity <sup>a</sup>	1-Month	104	0.16 (0.39)	0.67			
	6-Months	96	-0.19 (0.37)	0.61				
Shannon's Diversity	1-Month	104	0.01 (0.08)	0.89	48	0.06 (0.15)	0.67	
	6-Months	96	-0.11 (0.07)	0.11	45	0 (0.14)	0.99	
3rd trimester	Richness	1-Month	107	2.52 (6.37)	0.69	49	0.69 (1.82)	0.71
		6-Months	101	-13.11 (8.36)	0.12	47	2.54 (1.56)	0.11
	Evenness	1-Month	107	-0.01 (0.01)	0.43	38	0.03 (0.05)	0.52
		6-Months	101	-0.02 (0.01)	0.19	42	0.03 (0.05)	0.58
	Faith's Diversity <sup>a</sup>	1-Month	107	0.03 (0.35)	0.93			
	6-Months	101	-0.52 (0.42)	0.22				
Shannon's Diversity	1-Month	107	-0.05 (0.07)	0.50	49	0.09 (0.16)	0.57	
	6-Months	101	-0.12 (0.08)	0.15	47	0.15 (0.15)	0.34	
Postnatal	Richness	1-Month	109	2.18 (7.16)	0.76	50	-1.85 (2.54)	0.47
		6-Months	100	-2.55 (6.42)	0.69	46	-0.35 (1.05)	0.74
	Evenness	1-Month	109	-0.02 (0.01)	0.21	39	0.07 (0.1)	0.47
		6-Months	100	-0.01 (0.01)	0.40	41	0.06 (0.06)	0.36
	Faith's Diversity <sup>a</sup>	1-Month	109	-0.08 (0.39)	0.84			
	6-Months	100	0.11 (0.32)	0.74				
Shannon's Diversity	1-Month	109	-0.1 (0.08)	0.22	50	-0.07 (0.23)	0.78	
	6-Months	100	-0.05 (0.06)	0.46	46	-0.05 (0.1)	0.59	

<sup>a</sup> Faith's phylogenetic diversity not able to be calculated for fungal data as no phylogenetic tree can be derived.

<sup>b</sup> linear regression coefficient, which represents the change in alpha diversity metric, for a 1-SD increase in tooth lead level, after adjusting for tooth type, attrition, batch, exact age at stool sample collection, and child race.

examined the association between lead levels and bacterial community composition in humans. One such study was a randomized trial aimed to evaluate the effectiveness of a probiotic yogurt in lowering heavy metal levels among at-risk pregnant women and school-aged children in Tanzania (Bisanz et al., 2014). In the children, they found that elevated blood lead was associated with the increased abundance of *Succinivibrionaceae* and *Gammaproteobacteria* (Bisanz et al., 2014). While no OTUs were classified as *Succinivibrionaceae* in the current study, a total of 10 *Gammaproteobacteria* OTUs were significantly associated with lead levels, but the direction of association depended on more refined taxonomic classification (Supplemental Table 1). Specifically, *Acinetobacter* and *Xanthomonadaceae* OTUs negatively associated with lead levels, while *Enterobacteriaceae* OTUs were primarily positively associated with lead levels. A study by Eggers et al. examined urinary lead concentration in relation to gut microbiota composition in US adults (Eggers et al., 2019), finding that lead levels were significantly associated with increased bacterial richness and an increased abundance of *Proteobacteria*. In addition to observing an increased abundance of *Bifidobacteria* with lead exposure (which is of the phylum *Proteobacteria*), we also identified 9 *Proteobacteria* OTUs significantly and positively associated with lead levels (Supplemental Table 1), providing support for this finding.

A comparison of our finding that lead associates with fungal gut microbiota to that of others is limited due to the paucity of published epidemiological studies. In fact, to our knowledge, no other studies have examined the association between lead levels and fungal community composition in humans. More work has been done in environmental studies, however. For example, in an observational study of how lead levels impact native communities of fungi in soil surrounding an abandoned lead smelting factory, lead levels were associated with decreased fungal richness and distinct community composition (Faggioli et al., 2019); bacterial community composition was not examined. A similar study examining both the bacteria and fungi in the soil of a shooting range heavily polluted by lead found that lead did not alter bacterial microbiota, but was associated with clear changes in the fungal community structure relative to controls (Hui et al., 2012). They

also noted significantly higher fungal diversity in the polluted samples, in contrast to the previous study. The authors hypothesized that bacteria may be more capable of avoiding or adapting to lead exposure by acquiring new traits, whereas replacement may be the typical strategy for fungi (Hui et al., 2012). In a study of experimental rats, yeast levels were significantly decreased following lead exposure; however, yeast levels were determined using a culture-based approach rather than using high-throughput methods (Reddy et al., 2018).

The observed associations between lead exposure and infant gut microbiota could play a role in the impact of lead on childhood development. For example, lead exposure has been implicated as a risk factor for neurocognitive disorders, including Autism Spectrum Disorder (ASD) (Kim et al., 2016; Lakshmi Priya and Geetha, 2011). More recently, studies are beginning to investigate how gut microbiota can affect neurodevelopment disorders through the "gut-brain axis", in which signals from intestinal microbiota traffic to the central nervous system through neural, endocrine, and immune pathways (Sampson and Mazmanian, 2015). Indeed, a study comparing the gut microbiota of 40 individuals with ASD to that of 40 neurotypical controls found that individuals with ASD had a reduced abundance of *Bacteroidetes* and an increased abundance of *Collinsella* (Strati et al., 2017). Given that we found that lead exposure associates with reduced *Bacteroidetes* abundance and increased *Collinsella* abundance, it is plausible that these microbiota lie in the causal pathway between lead exposure and ASD. A mechanistic study using a mouse model of ASD provides further support for this hypothesis, as it demonstrated that oral treatment of *Bacteroides fragilis* corrected autism-related behavioral impairments and gut dysbiosis (Hsiao et al., 2013). The gut microbiome may play a similar role in other disorders affected by lead exposure.

Our study is not without limitations. Given the observational nature of this study, results are associative and not necessarily causative. However, two of the three tooth lead measurements were quantified prior to birth and therefore preceded the measurement of infant microbiota composition. Though birth is a central driver of infant gut microbiota colonization (Milani et al., 2017), there is evidence suggesting that highly limited, but viable bacteria are present in the fetal



**Table 3**  
Association between tooth lead levels and beta diversity metrics.

Time of Metal Measurement	Outcome	Time of Stool Measurement	PERMANOVA <sup>a</sup>			
			N	R <sup>2</sup>	p-value <sup>c</sup>	
<b>Bacterial</b>						
2nd trimester	Unweighted UniFrac	1-Month	104	0.007	0.87	
		6-Months	96	0.011	0.32	
	Weighted UniFrac	1-Month	104	0.012	0.28	
		6-Months	96	0.006	0.70	
	Canberra	1-Month	104	0.009	0.70	
		6-Months	96	0.011	0.46	
	Bray-Curtis	1-Month	104	0.010	0.40	
		6-Months	96	0.013	0.21	
	3rd trimester	Unweighted UniFrac	1-Month	107	0.008	0.75
			6-Months	101	0.011	0.25
Weighted UniFrac		1-Month	107	0.008	0.47	
		6-Months	101	0.006	0.66	
Canberra		1-Month	107	0.009	0.66	
		6-Months	101	0.010	0.33	
Bray-Curtis		1-Month	107	0.006	0.80	
		6-Months	101	0.009	0.55	
Postnatal		Unweighted UniFrac	1-Month	109	0.008	0.64
			6-Months	100	0.008	0.84
	Weighted UniFrac	1-Month	109	0.008	0.49	
		6-Months	100	0.006	0.75	
	Canberra	1-Month	109	0.008	0.84	
		6-Months	100	0.010	0.59	
	Bray-Curtis	1-Month	109	0.007	0.75	
		6-Months	100	0.009	0.58	
	<b>Fungal<sup>b</sup></b>					
	2nd trimester	Canberra	1-Month	48	0.018	0.59
6-Months			45	0.016	0.94	
Bray-Curtis		1-Month	48	0.040	<b>0.048</b>	
		6-Months	45	0.014	0.65	
3rd trimester	Canberra	1-Month	49	0.019	0.51	
		6-Months	47	0.014	0.96	
	Bray-Curtis	1-Month	49	0.039	<b>0.049</b>	
		6-Months	47	0.006	0.94	
Postnatal	Canberra	1-Month	50	0.016	0.78	
		6-Months	46	0.021	0.52	
	Bray-Curtis	1-Month	50	0.021	0.39	
		6-Months	46	0.028	0.22	

<sup>a</sup> adjusted for tooth type, attrition, batch, exact age at stool sample collection, and child race.

<sup>b</sup> UniFrac metrics not able to be calculated for fungal data as no phylogenetic tree can be derived.

<sup>c</sup> For bold p-values indicating  $p < 0.05$ .

intestine at mid-gestation (Rackaityte et al., 2020). Thus, fetal lead exposure may alter early life microbiota either directly or through the fetal microbiota; however, reverse causation cannot be ruled out, as it is unknown if fetal microbiota can impact the uptake of lead by the fetus. Indeed, in addition to previous studies demonstrating that lead can directly alter microbiota composition (Gao et al., 2017), it is also known that microbes can act as bioremediators, capable of directly binding to, detoxifying, and preventing the absorption of lead (Monachese et al., 2012; Tian et al., 2012). Additionally, lead has been detected in breast milk samples (Vollset et al., 2019) and therefore may also impact microbial communities; however, this could not be assessed as breast milk lead levels were not measured in the WHEALS cohort (though, the postnatal tooth measurement may capture this exposure).

The small sample size used in this analysis—particularly for fungal microbiota—limited the ability to extensively adjust for covariates and may result in spurious findings. Though we adjusted for race—which is highly associated with both lead levels and microbiota composition and

explained away the association between other hypothesized confounders and lead levels—residual confounding is still possible. In fact, the only covariate examined that remained associated with lead exposure after adjusting for race was birthweight z-score; however, we posit that this is a potential mediator rather than a confounder, as birth outcomes may be directly affected by lead exposure (Taylor et al., 2015) and lie within the causal pathway. Lastly, metagenomic sequencing was not performed on the infant stool samples, so the functional capacity of organisms was not quantified.

## 5. Conclusions

Using a U.S. birth cohort of racially diverse children, we identified an association between *in utero* tooth lead levels and gut fungal community composition at 1-month of age, where higher levels of 2nd trimester tooth lead was associated with lower abundances of *Candida* and *Aspergillus* and higher abundances of *Malassezia* and *Saccharomyces*. Third trimester lead was also associated with lower abundances of *Candida*. Though lead did not significantly impact the overall structure of the infant gut bacterial community, it associated with the abundance of some specific bacterial taxa, including an increase in *Collinsella* and *Bilophila*, as well as a decrease in *Bacteroides* taxa. Additional studies are needed to confirm these findings, ideally in both epidemiological studies of humans as well as animal models.

## CRedit authorship contribution statement

**Alexandra R. Sitarik:** Conceptualization, Formal analysis, Visualization, Writing - original draft, Writing - review & editing. **Manish Arora:** Resources, Writing - review & editing, Supervision. **Christine Austin:** Resources, Writing - review & editing. **Lawrence F. Bielak:** Writing - review & editing. **Shoshannah Eggers:** Resources, Writing - review & editing. **Christine C. Johnson:** Writing - review & editing, Supervision, Funding acquisition. **Susan V. Lynch:** Resources, Writing - review & editing, Supervision. **Sung Kyun Park:** Writing - review & editing. **Kuan-Han Hank Wu:** Formal analysis, Writing - review & editing. **Germaine J.M. Yong:** Resources, Writing - review & editing. **Andrea E. Cassidy-Bushrow:** Conceptualization, Writing - original draft, Writing - review & editing, Funding acquisition.

## Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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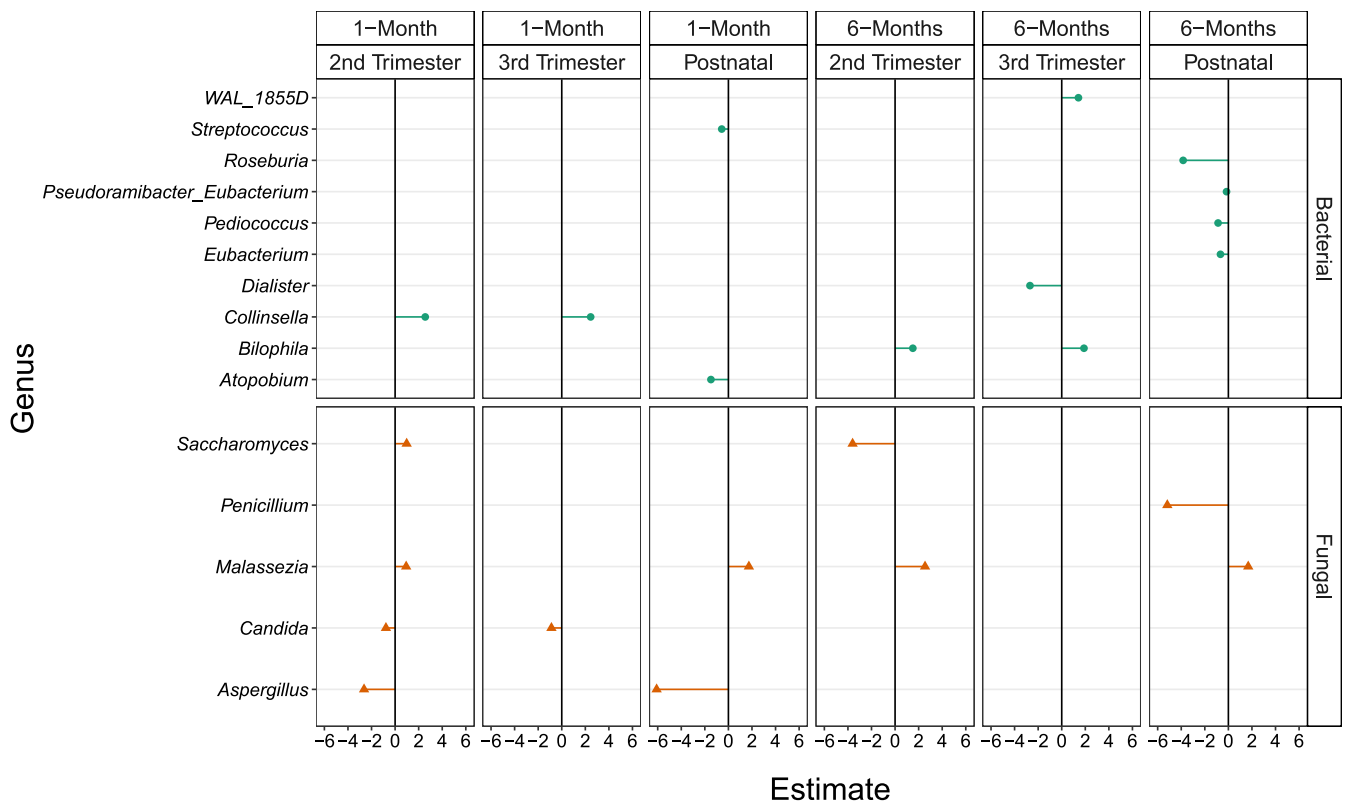
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## Appendix A. Supplementary material

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.envint.2020.106062>.



**Fig. 1.** Bacterial and fungal genera significantly associated with *in utero* and postnatal lead levels ( $P_{FDR} < 0.05$ ). Estimates shown are coefficients from zero-inflated negative binomial models or standard negative binomial models, after adjusting for tooth type, attrition, batch, exact age at stool sample collection, and child race. These can be interpreted as the change in log genera abundance, for a 1-standard deviation increase in lead levels.

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