Metabolomic Profiling of Burkholderia Cenocepacia in Synthetic Cystic Fibrosis Sputum Media Reveals Nutrient Environment-Specific Production of Virulence Factors

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Abstract

Infections by *Burkholderia cenocepacia* lead to life-threatening disease in immunocompromised individuals, including those living with cystic fibrosis (CF). While genetic variation in various *B. cenocepacia* strains has been reported, it remains unclear how the chemical environment of CF lung influences the production of small molecule virulence factors by these strains. Here we compare metabolomes of three clinical *B. cenocepacia* strains in synthetic CF sputum media (SCFM2) and in a routine laboratory media (LB), in the presence and absence of the antibiotic trimethoprim. Using a mass spectrometry based untargeted metabolomics approach, we identify several compound classes which are differentially produced in SCFM2 compared to LB media, including siderophores, antimicrobials, quorum sensing signals, and various lipids. Furthermore, we describe that specific metabolites are induced by the antibiotic trimethoprim only in SCFM2 when compared to LB. Herein, C13-acyl-homoserine lactone, a quorum sensing signal previously not known to be produced by *B. cenocepacia* as well as pyochelin-type siderophores were exclusively detected during growth in SCFM2 in the presence of trimethoprim. The comparative metabolomics approach described in this study provides insight into environment-dependent production of secondary metabolites by *B. cenocepacia* strains and suggests future work which could identify personalized strain-specific regulatory mechanisms involved in production of secondary metabolites.

Introduction

Cystic fibrosis (CF) patients experience chronic lung infections by pathogenic microorganisms with accompanying airway inflammation, leading to a progressive decline in lung function and respiratory failure^{1,2}. The *Burkholderia cepacia* complex (Bcc) is a group of over 20 related *Burkholderia* species which can cause chronic infections in CF patients^{3,4}. Bcc bacteria display intrinsic and acquired resistance to many common antibiotics⁵. Additionally, Bcc spp. are capable of spreading between patients and can lead to cepacia syndrome, a condition characterized by rapidly progressing necrotizing pneumonia, bacteremia, and a sharp decline in lung function⁵⁻⁹. The majority of Bcc infections are caused by either *Burkholderia cenocepacia* or *Burkholderia multivorans*¹⁰. While Bcc infections in general are associated with negative clinical outcomes, *B. cenocepacia* in particular is one of the most feared CF pathogens^{11,12}. *B. cenocepacia* is associated with reduced survival rates and an increased risk of developing cepacia syndrome compared to other Bcc species¹³⁻¹⁵.

Burkholderia spp. bacteria are known to produce a variety of small organic molecules also known as natural products, allowing these bacteria to adapt and survive in diverse ecological niches^{9,16-18}. These natural products play important role in swarming, biofilm formation, iron acquisition, combating competition, and quorum sensing (QS). As such, many of these natural products serve as potential virulence factors with important roles implicated in infection. Genomic approaches have been utilized extensively to compare the potential of various Bcc spp. to produce virulence factors^{19,20}. However, pathogens display variable phenotypes when cultured in diverse nutritional environments^{21,22}. While

genomic analysis provides insight into the metabolic potential of these bacteria, it does not accurately represent their metabolic state under specific environmental conditions. On the other hand, metabolomics is a complementary technique that provides a global “snapshot” of an organism’s metabolic state under a given condition and therefore is a relevant representation of the organism’s molecular phenotype. Thus, comparative metabolomics is an attractive approach for monitoring metabolic responses to external stimuli.

Previous work has demonstrated that nutrient availability in infection sites is associated with phenotypic changes in CF pathogens, including increased virulence^{23,24}. Along these lines, clinical *B. cenocepacia* isolates cultured in a minimal media supplemented with 12.5% (w/v) CF sputum were shown to exhibit differential expression of genes involved in virulence processes, such as antimicrobial resistance, iron uptake, protection against reactive oxygen/nitrogen species, motility, and secretion²⁵. Therefore, it is evident that nutrient composition is an important consideration for developing a clinically relevant *in vitro* model of CF sputum to be used in comparative metabolomic studies. To this end, an assortment of artificial sputum media formulations have been established for studying CF pathogens *in vitro* that overcome the challenges with the inherent heterogeneity associated with culturing in sputum from patients^{22,23,26-31}. Among these, a chemically defined synthetic CF sputum media (SCFM) was developed, and subsequently refined (SCFM2) by adding DNA, mucin, 1,2-dioleoyl-sn-glycero-3-phosphocholine (DOPC), and *N*-acetyl glucosamine (GlcNAc) to approximate the average physical and chemical environment of CF sputum^{31,32}. The sputum of CF patients is a complex mixture of amino acids, carbohydrates, mucins, lipids, and micronutrients which support pathogen growth *in vivo*^{23,24,33-38}. A transposon sequencing (Tn-seq) approach was used to demonstrate that genes required for fitness *in vitro* are practically identical in CF sputum and in SCFM2 media for the prominent CF pathogen, *Pseudomonas aeruginosa*³¹. Additionally, the transcriptomic profiles of *P. aeruginosa* in CF sputum samples were best represented by an SCFM2-based *in vitro* model compared to any other single laboratory model tested³⁹. While no individual laboratory model completely captured gene expression profiles observed during *P. aeruginosa* CF infections, the combination of SCFM2 and murine lung infection model came closest³⁹.

These findings support the validity of SCFM2 media as a model for studying CF pathogens in a controlled laboratory setting. To better understand small molecule production under disease-relevant conditions, we set out to compare the metabolomes of three *B. cenocepacia* strains when cultured in SCFM2 media versus a standard laboratory growth media. In this study, we demonstrate that *B. cenocepacia* strains exhibit altered metabolomic profiles when cultured in SCFM2 media compared to Luria broth (LB). Specifically, we observed that growth in SCFM2 media upregulates production of pyochelin-type siderophores while downregulating production of ornibactin siderophores. We also report production of unique *N*-acyl-homoserine lactone (AHL) QS signals in SCFM2, with C13 acyl chains containing AHLs detected exclusively in SCFM2 samples in the presence of trimethoprim. Additionally, we note that several families of lipids (including hopanoids, cyclopropane fatty acids, monoacylglycerols, and phosphatidylethanolamines) exhibit increased production in SCFM2 compared to LB. Finally, we

show that sublethal concentrations of trimethoprim lead to media-dependent changes in secondary metabolite production, and that metabolism of trimethoprim itself is significantly upregulated during growth in SCFM2 compared to LB. Overall, this study provides valuable insight into *B. cenocepacia* secondary metabolite production under disease-relevant environmental conditions and highlights the importance of use of appropriate *in vitro* models for metabolomic studies.

Methods

Bacterial strains. *B. cenocepacia* C5424, K56-2, and J2315 clinical isolates were used for culturing during this study^{59,118,119}. These strains belong to the highly transmissible and epidemic ET12 lineage, wherein the J2315 and K56-2 strains are clonally related. As reported, the C5424 and J2315 strains produced observable amounts of the pigment pyomelanin whereas the K56-2 strain did not⁴⁶.

Media Formulation and Growth Conditions. SCFM2 media was prepared as previously described by Turner and co-workers³¹. All culturing was performed following previously established methods⁴³. Briefly, *B. cenocepacia* C5424, J2315, and K56-2 frozen glycerol stocks were streaked onto LB agar plates, and then incubated overnight at 37 °C. Subsequently, 5 mL of LB medium was inoculated using colonies from the overnight incubated plates, which was then incubated overnight at 37°C while shaking at 250 rpm. After 16h, sterile 125 mL Erlenmeyer flasks containing 20 mL of either LB-50 mM MOPS (pH 7.0) or SCFM2 media were inoculated with each overnight culture at an initial OD₆₀₀ of 0.05. For each strain and media combination, four flasks were supplemented with 50 µL of a 12 mM trimethoprim stock (suspended in DMSO) to a final concentration of 30 µM, and four flasks were supplemented with 50 µL of DMSO as a vehicle control. These cultures were grown at 37°C while shaking at 250 rpm. Half of the culture (10 mL) from each combination of strain and media type was harvested after 24h (four containing trimethoprim and four with DMSO instead of trimethoprim), and the remaining half (10 mL) was harvested at 48h.

Sample Extraction. Features were extracted using our previously established methods⁴³. Briefly, each culture was extracted using two different methods: liquid-liquid extraction with ethyl acetate (EtOAc) and solid-phase extraction (SPE) using C18 columns (100 mg bed weight). For EtOAc extractions, 20 mL of the solvent was added to whole cultures and mixed every 30 min for 2h before centrifuging at 2,000×g for 3 min. The organic layer (top) was aspirated off with a glass pipette and transferred to a scintillation vial before drying *in vacuo* with a centrifugal evaporator. For solid-phase extractions, cultures were centrifuged at 10,000×g for 10 min and decanted to remove supernatant fluid. SPE columns were initially washed with 10 mL of 100% MeCN, subsequently equilibrated with water (10 mL) and then loaded with the culture supernatant. Analytes were then sequentially eluted from the column using 5 mL each of 20%, 50% and 100% MeCN. These three fractions were pooled together and dried *in vacuo* with a centrifugal evaporator.

Ultra-high-Performance Liquid Chromatography Tandem Mass Spectrometry (UHPLC-MS/MS) Data Acquisition. Dried extracts were dissolved in 600 µL of 80% MeOH (LC-MS grade) containing 1 µM

sulfadimethoxine as an internal standard before analyzing on an Agilent 1290 Infinity II UHPLC system (Agilent Technologies) coupled to an ImpactII ultra-high resolution Qq-TOF mass spectrometer (Bruker Daltonics, GmbH, Bremen, Germany) equipped with ESI source. Chromatographic separation was achieved on a KinetexTM 1.7 μ m C18 reversed phase UHPLC column (50 $\times$ 2.1 mm). Solvent A consisted of water with 0.1% (v/v) formic acid and solvent B consisted of MeCN with 0.1% (v/v) formic acid. The gradient used for chromatographic separation consisted of 5% B and 95% A for 2 min, a linear increase to 95% B over 17 min, a hold for 3 min, a linear decrease to 5% B in 1 min, and hold for 1 min with a constant flow rate of 0.5 mL/min throughout. For MS spectra acquisition, the instrument was set to capture features from m/z 50 - 2000 Da in positive ion mode. External calibration was performed prior to data collection using ESI-L Low Concentration Tuning Mix (Agilent Technologies), with hexakis(1H,1H,2H-perfluoroethoxy)phosphazene employed as internal lock-mass calibrant throughout the run. Ion source parameters were set to 4500 V for capillary voltage, 2 bar for nebulizer gas pressure (N_2), 200°C for ion source temperature, 9 L/min for dry gas flow, and a spectral rate of Hz for MS¹ and 6 Hz for MS². For MS² data acquisition, the eight most intense precursor ions per MS¹ scan were selected for MS² fragmentation. A basic stepping function was used to fragment ions at 50% and 125% of the CID calculated for each m/z , with timing of 50% for each step. Similarly, a basic stepping of collision RF of 550 and 800 Vpp with a timing of 50% for each step and transfer time stepping of 75 and 90 μ s with a timing of 50% for each step was employed. The MS/MS active exclusion parameter was set to 2 and the active exclusion was released after 30 s. The mass of the internal lock-mass calibrant was excluded from MS² acquisition. A mixture of 6 compounds (amitryptiline, sulfamethazine, sulfamethizole, sulfachloropyridazine, sulfadimethoxine, coumarin-314) was run as quality control every 8 samples to ensure consistent instrument and column performance.

MS³ experiments for further structural analysis of the metabolite feature with m/z 339.083 annotated as thiazostatin A/B (**1**) were performed using a Waters Corporation Cortecs UPLC T3 column (2.1 $\times$ 150 mm, 1.6 μ m particle size) coupled to a high-resolution accurate mass Orbitrap ID-X tribrid mass spectrometer. The chromatographic method for sample analysis involved elution with 100% water 0.1% formic acid (mobile phase A) and MeCN and 0.1% formic acid (mobile phase B) using the following gradient program: 0 min 95% A; 0.5 min 95% A; 6 min 0% A (curve 7); 9.4 min 0% A; 9.5 min 95% A; 11 min 95% A. The flow rate was set at 0.4 mL/min. The column temperature was set to 40 °C, and the injection volume was 1 μ L. The Orbitrap ID-X is a tribrid spectrometer that utilizes quadrupole isolation with dual detectors, an orbitrap and an ion trap, with a maximum resolving power of 500,000 full width at half maximum (FWHM) at m/z 200 and mass accuracy of <1 ppm. The heated electrospray ionization (HESI) source was operated at a vaporizer temperature of 275 $\pm$ 5°C, a spray voltage of 3.5 kV, and sheath, auxiliary, and sweep gas flows of 40, 8, and 1 in arbitrary units, respectively. The instrument acquired full MS data in the 100-1000 m/z range in positive ionization mode at 30000 resolution. MS³ data was collected with a MS¹ isolation window of 0.8 m/z , HCD activation of 30% $\pm$ 50%, a MS² isolation window of 1.8 m/z , MS² HCD activation of 40%, and product ion detection in the orbitrap at 30000 resolution.

Data Processing, Feature-based Molecular Networking, and Feature Annotation. The LC-MS/MS data presented in this manuscript is deposited in the online repository MassIVE and publicly available (MSV000087793). The Compass DataAnalysis software was used to convert all the raw spectral files (.d format) to centroided, lock-mass corrected format (.mzXML) for downstream analyses. The converted spectral files were uploaded to MZmine2 (v.2.5.1) for feature detection⁴⁴. Data files were batch processed and filtered using a MS¹ mass detection signal threshold of 5.0×10^2 counts. The following parameters were applied: (i) chromatogram builder (minimum time span: 0.05 min; minimum intensity of the highest data point in the chromatogram: 1.5×10^3 ; m/z tolerance: 10 ppm); (ii) chromatogram deconvolution (local minimum search, m/z range for MS² scan pairing: 0.025 Da; retention time range for MS² scan pairing: 0.2 min); (iii) isotopic peaks grouper (m/z tolerance: 10 ppm; retention time tolerance absolute: 0.1 min; maximum charge: +3; monotonic shape: true; representative isotope: most intense); (iv) join aligner (m/z tolerance: 10 ppm; retention time tolerance: 0.1 min); (v) peak finder (intensity tolerance: 10%; retention time tolerance (absolute): 0.1; m/z tolerance: 10 ppm). A table with ion intensities for each feature was exported (.csv format) for statistical analyses and the “Export for SIRIUS” module was used to generate an .mgf file for batch analysis with SIRUS 4. Additionally, the “Export for GNPS” module was used to convert and export the feature quantification table (.csv format) and the corresponding list of MS² spectra linked to MS¹ features (.mgf format) needed to generate a feature-based molecular network. The feature quantitation table and .mgf file were submitted to the Global Natural Products Molecular Networking (GNPS) platform along with a metadata file (.txt format) containing sample information, and a feature-based molecular network was created⁴⁵. The molecular network and parameters used can be accessed using the following link: <https://gnps.ucsd.edu/ProteoSAFe/status.jsp?task=a80b05b26a1747cfa6caf7f9446edd75>. Briefly, the data was filtered by removing all MS/MS peaks within +/- 17 Da of the precursor m/z . A parent mass tolerance of 0.01 Da and a MS/MS fragment ion tolerance of 0.05 Da were applied to create consensus spectra. The network was created with the edges filtered to have a cosine score above 0.7 and at least 4 matched peaks, and edges connecting two nodes were set to be kept in the network if each of the nodes appeared in each other's respective top 10 most similar nodes. All the mass spectra in the generated network were queried against the spectral libraries available on GNPS. The matched experimental and library spectra were set to have a similarity score above 0.7 and more than 4 matched peaks. This network was exported to Cytoscape for visualization, and nodes appearing in media or solvent blanks were removed for clarity (unless otherwise indicated)¹²⁰. Boxplots were constructed for features of interest using the Plotter Dashboard (v.0.4) available on the GNPS platform. UpSet plots were generated with the UpSetR package in R⁴⁸.

Metabolomic features of interest that remained unknown after library searching were further annotated by first searching through literature for molecules which are known to be produced by *Burkholderia* spp. and developing an in-house database. Our database was curated by inserting structures manipulated using MarvinSketch (v.20.9.0, ChemAxon Ltd.) into a spreadsheet installed with JChem for Excel (v.20.8.0.62, ChemAxon Ltd.). This database was used to identify candidate annotations for features, and these annotations were confirmed by manually inspecting MS² spectra and either comparing against

published MS² spectra (when available) or against the MS² spectra of commercial analytical standards. We then applied the MS2LDA workflow to recognize patterns of common MS² fragments and neutral losses ("Mass2Motifs") corresponding to molecular substructures in our dataset¹²¹. This MS2LDA analysis is available at the following link: <https://gnps.ucsd.edu/ProteoSAFe/status.jsp?task=628622833d0046048dba64e65ced542c>. In addition, we performed batch analysis on the dataset using SIRIUS 4 integrated with CSI:FingerID and CANOPUS^{91,92,122-125}. SIRIUS 4 (v.4.0.1) was employed (using the default settings for a qTOF instrument) to predict molecular formulas for unknown features and develop fragmentation trees for manual annotation of MS² spectra¹²²⁻¹²⁴. CSI:FingerID was then utilized to predict molecular properties of unknown features, which were then queried against molecular properties predicted for compounds in all available molecular databases¹²⁵. This *in silico* tool lead to a ranked list of predicted structures, even when published MS² spectra were not available for these structures. Next, we deployed CANOPUS to classify features into molecular families using ClassyFire, providing biological insight in the absence of structural annotations^{91,92}. MASST searching was used to find metabolomic datasets collected by other researchers containing a match to the metabolic feature annotated as thiazostatin A/B (1)⁶⁷. This MASST job and the parameters used are available at: https://gnps.ucsd.edu/ProteoSAFe/result.jsp?task=c5d7b5cc7459443286fed543e5140ca4&view=view_all_datasets_matched.

To determine differences in trimethoprim metabolism between *B. cenocepacia* strains grown in SCFM2 and LB media, metabolic features containing a trimethoprim substructure (appearing as Mass2Motif 543 in the MS2LDA analysis) were extracted from the feature quantification table (.csv format) generated by MZmine2. These features were then subjected to further statistical analysis on the MetaboAnalyst web server¹²⁶. After uploading the peak intensity table for trimethoprim metabolites, data was log transformed and Pareto scaling was applied. Volcano plots comparing trimethoprim against non-trimethoprim cultures were then created using a fold-change threshold of 2.0 and an FDR adjusted p value threshold of 0.05, as calculated by a t-test.

ICP-MS analysis of LB and SCFM2 media. ICP-MS was performed on aliquots of LB and SCFM2 media on a Perkin Elmer NexION 2000 with a S10 Autosampler at the Emory University Mass Spectrometry Center. Samples were diluted 1:10 and 1:100 with Type I deionized water before aspirating into the plasma using the NexION 2000 Peristaltic pump. A kinetic energy discrimination method with helium flow set to 5 arbitrary units was used to detect the ⁵⁷Fe isotope using a pulse counting detection method. Calibration was done from 0.1 to 1000 ppb. Samples were analyzed in technical triplicate for each media type. To determine the amount of iron in each media sample, the mean and standard deviation of all concentrations measured across both dilutions was calculated and converted from ppb to micromolar units.

Results And Discussion

Description of global changes in metabolome

The nutritional composition of standard laboratory growth media such as LB is inadequate in modeling bacterial physiology during infection, as nutrient availability can vary markedly between infection sites and standard growth media. To overcome this challenge, the previously developed SCFM2 was used to model the physical and chemical environment of human sputum from patients with cystic fibrosis. In this study, we employed an untargeted metabolomic approach to compare the metabolomes of three different strains of *B. cenocepacia* namely, C5424, K56-2, and J2315 when cultured in SCFM2 and LB in triplicate (Fig. 1a). These strains are clonal and are associated with the highly transmissible, epidemic ET12 lineage of *B. cenocepacia*¹⁹. In addition, comparative metabolomic analysis was also carried out in the presence and absence of trimethoprim, which is an antibiotic used clinically for treatment of *Burkholderia* infections⁴⁰. This antibiotic is known to upregulate biosynthetic pathways involved in production of secondary metabolites in *B. thailandensis* and in select strains of *B. cenocepacia* in LB, but its effect has not been investigated in SCFM2⁴¹⁻⁴³. In effort to capture a diverse range of compound classes, extractions were performed on bacterial cultures using both liquid-liquid (with ethyl acetate, EtOAc) and solid-phase extraction (SPE) methods. Extracts were then analyzed using high resolution tandem mass spectrometry coupled with ultra-high-performance liquid chromatography (UHPLC-HRMS/MS). Metabolite features representing analytes detected at a unique *m/z* and retention time were extracted, aligned, and quantified using the open-source MZmine2 software and feature-based molecular networking was performed (Figure 1b)^{44,45}.

We first compared the metabolome of each strain irrespective of media type, extraction method, and exposure to antibiotic (Fig. 2a). The largest number of unique metabolite features (20.5%) were detected exclusively in the extracts of the strain K56-2, whereas 8.5% unique features were detected in the extracts of J2315 strain and 1.1% in the extracts of the strain C5424. While 30.2% of features were shared by all three strains, the largest number of features were shared between the strains J2315 and C54524 (31.7%). Among other factors, the similar metabotype of these two strains may be reflective of their ability to produce a brown pigment known as pyomelanin, which is not produced by the strain K56-2^{46,47}. Next, we compared the metabolomes acquired using the SPE and EtOAc extraction methods separately at 48h post-inoculum. This analysis revealed that many metabolite features are exclusively detected in extracts generated using SPE as compared to extracts generated using with EtOAc (Fig. 2b). A total of 32.5% of the metabolite features were detected only with the SPE method, while 9.5% of the total features were unique to EtOAc extraction method (Fig. 2b). Thus, the extraction method employed results in biased metabolomics comparisons when only one type of extraction strategy is employed. Lastly, we generated UpSet plots to visualize the number of features unique to each strain under different growth conditions as well as exposure to sub-lethal dose of antibiotic trimethoprim after subtraction of media background (Fig. 2c)⁴⁸. This analysis revealed that media specific differences were the largest driver of metabolomic diversity within this study, with 878 features shared between all LB samples (●) and 563 features shared between all SCFM2 samples (●). In comparison, 513 features were detected in all samples. The UpSet plot also demonstrates that trimethoprim induces a unique metabolomic response in the K56-2 strain, with 362 metabolomic features exclusively detected in SCFM2 samples (●), 59 exclusive to LB samples (●), and 44 detected in both media types (●). Interestingly, 456 metabolomic features were uniquely

detected in all LB samples except for K56-2 with trimethoprim (●) and 37 in all SCFM2 samples except for K56-2 with trimethoprim (●). While our prior studies have shown that K56-2 exhibits a unique metabolomic response to trimethoprim, our analysis demonstrates that this response is even more apparent in an environment representative of CF sputum⁴³. Another pattern emerging from the UpSet plot is that many features are uniquely detected as shared between the pigmented J2315 and C5424 strains, with 203 features uniquely detected in LB (●), 90 in SCFM2 (●), and 93 across both media types (●). An additional 59 features were uniquely detected in C5424 and J2315 strains cultured in LB and SCFM2 in the presence of trimethoprim (●), hinting at a distinct response to trimethoprim that is associated with pyomelanin production. Such responses are hallmark of the personalized phenotypes of *Burkholderia* strains observed during infection⁴⁹⁻⁵². Future investigations into phenotypic differences as well as differences in gene expression via transcriptomics will provide insights into the biochemical underpinnings driving these observations. Specific metabolites underlying these personalized chemotypes are discussed below.

Differences in siderophore production between LB and SCFM2 cultures

Siderophores are compounds secreted by bacteria to acquire iron from the surrounding environment, and were among the metabolites which were differentially produced by *B. cenocepacia* strains in LB and SCFM2 in this study. In healthy mammalian hosts, the pool of free iron is limited due to poor solubility of iron in its ferric state under physiological conditions, and because the majority of iron is either located in intracellular compartments or bound by host proteins such as hemoglobin, transferrin, lactoferrin, and ferritin^{53,54}. In contrast, levels of both free and ferritin-bound iron are higher in CF sputum compared to sputum from healthy hosts^{55,56}. Since this cofactor is essential for many important biological processes, iron-acquisition is required for survival in the host environment and can influence microbe-microbe interactions^{54,57}. Siderophores which are known to be produced by *Burkholderia* include pyochelin, ornibactins, salicylic acid, and cepabactin^{43,54,58}. Among these, *B. cenocepacia* have been shown to produce ornibactin and pyochelin⁵⁴. In our study, various structural analogs of pyochelin were detected exclusively in SCFM2 in the presence of trimethoprim (Fig. 3). Pyochelin itself was not detected in this study, despite prior reports of low-level production by *B. cenocepacia*. These prior reports did not use mass spectrometry or NMR to report pyochelin production, but relied on fluorescence-based thin layer chromatography⁵⁹. Using MSⁿ analysis, we report production of a methylated derivative of pyochelin rather than pyochelin by *B. cenocepacia* strains used in this study as described below.

The feature with m/z 339.083 had a database annotation as a pyochelin methyl ester in GNPS (Supplementary Fig. S1a)⁶⁰. However, pyochelin methyl ester has only been reported as a synthetic product generated to facilitate NMR characterization, and there is no available biosynthetic evidence that would support methylation on the carboxylic acid to produce an ester⁶¹. This feature was detected exclusively in cultures grown in SCFM2 supplemented with trimethoprim in both the K56-2 and C5424 strains, albeit low levels were observed in C5424 (Fig. 3a). Comparisons with the GNPS library MS² spectra of pyochelin revealed that the fragment peaks with m/z 120.045, 180.048, and 190.032 are

shared between the two molecules while others were shifted by 14.015 Da ($-\text{CH}_2$), supporting the annotation as a methylated analog of pyochelin (Supplementary Fig. S1b). A structure search of pyochelin in SciFinder revealed thiazostatin as a likely candidate. Thiazostatin A/B (**1**) are previously reported stereoisomeric natural products that are related to pyochelin by an additional C4"-methylation of the thiazolidine ring (Fig. 3a)^{62,63}. This thiazolidine C4"-methylation has also been observed in structurally homologous metabolites including isopyochelin, watasemycin, and yersiniabactin⁶⁴⁻⁶⁶. By MS² analysis alone, the location of the methylation cannot be unambiguously determined, although the mass shifts of the fragment ion with m/z 146.027 in the MS² spectrum of pyochelin to m/z 160.043 in our unknown metabolite's MS² spectrum suggests it is found on the terminal thiazolidine ring. To confirm the position of methyl group, we conducted MS³ analysis on the fragment ion with m/z 186.058. This analysis revealed that fragmentation of 186.058 yields a MS³ ion with m/z 158.063, which indicates methylation on the thiazolidine ring rather than at the carboxylic acid (Supplementary Fig. S2). Thus, this feature is reported as thiazostatin A/B. Thiazostatin A/B have been found to display antioxidant activity, and further screening is necessary to determine if this compound possess additional bioactivities⁶². Using MASST through the GNPS platform, we searched the MS² spectrum of this compound against all public spectral datasets (Supplementary Fig. S3)⁶⁷. This MASST search found dataset matches in extracts of *Pseudomonas* spp. grown *in vitro*, as well as in datasets analyzing the metabolomes of humans with various inflammatory diseases (including CF, diabetes, irritable bowel syndrome, rheumatoid arthritis, and HIV). Detection of this molecule in humans with inflammatory disease raises the possibility that this molecule may be important in influencing host microbiome structure, although further studies will be needed to explore whether this observation is truly associated with any biological significance.

Another metabolite with m/z 307.021 was detected exclusively in SCFM2 media extracts in both the *B. cenocepacia* strains K56-2 and J2315. In our molecular network, this feature matched with 2'-(2-hydroxyphenyl)-4'-thiazolyl-2,4-thiazoliny-4-carboxylic acid (HPTzTn-COOH, **2**) in the GNPS spectral library, which was further verified by manually comparing experimental MS² spectra with a previously published MS² spectra (Fig. 3a, Supplementary Fig. S4a)⁶⁸. Like pyochelin, HPTzTn-COOH is a siderophore that is dependent on salicylic acid and cysteine as biosynthetic precursors and is capable of chelating Fe^{3+} in addition to other metal ions including Al^{3+} , Ni^{2+} , and Ca^{2+} ^{68,69}. HPTzTn-COOH was detected exclusively in SCFM2 media, and primarily in the K56-2 strain, although also at low levels in J2315 (Fig 3a). This metabolite exhibited increased production in the presence of trimethoprim for the K56-2 strain and was not detected in the absence of trimethoprim for the J2315 strain.

Next, a feature with m/z 222.022 was detected exclusively in K56-2 cultures grown in SCFM2 media in the presence of trimethoprim (Fig. 3b, Supplementary Fig. S4b). This feature was annotated as aeruginoic acid (**3**), which is a shunt product in the pyochelin biosynthesis pathway observed in *Pseudomonas* and *Burkholderia* spp.^{70,71}. Aeruginoic acid is the oxidized form of aeruginaldehyde (also known as the integrated quorum sensing signal, or IQS), which has been proposed as a "fourth QS

molecule” in *P. aeruginosa*, although this claim is disputed^{72,73}. The reactive aldehyde moiety of aeruginaldehyde also leads it to become incorporated into more complex natural products (such as malleonitrone or mindapyrrole B)^{74,75}. Interestingly, pyochelin has been shown to spontaneously undergo cleavage and subsequent transformation into aeruginaldehyde when incubated in buffer solution at 30°C⁷⁴. Another shunt product of pyochelin biosynthesis with *m/z* 210.058 was detected under the same conditions as aeruginic acid, and annotated as aerugine (**4**) (Fig. 3b, Supplementary Fig. S4c). Aerugine has been previously isolated from *Pseudomonas fluorescens* and exhibits selective antifungal activity⁷⁶. Recent work by Kaplan *et. al* has indicated that aeruginaldehyde, aerugine, and aeruginic acid all have iron-binding activity with aeruginic acid binding to Fe³⁺ with a 1:1 ratio, aeruginaldehyde binding with a 2:1 ratio, and aerugine binding with a 3:1 ratio⁷¹. Unlike pyochelin, aeruginic acid has a specific affinity for iron compared to other biologically relevant metals⁷¹. The largest abundance of thiazostatin is observed in the strain K56-2, which is likely why these intermediates are also detected in the strain K56-2.

In contrast to metabolites from the pyochelin pathway described above, the ornibactin class of siderophores was detected in the culture extracts of LB medium and not detected in culture extracts of SCFM2 (Supplementary Fig. S5). The SCFM2 media is prepared by adding 3.60 µM iron in the form of Fe₃SO₄, which was observed to be the average iron concentration present in expectorated sputum collected from CF patients²³. Recently, it has been reported that commercial sources of mucin can be contaminating sources of iron that lead to altered siderophore production in *P. aeruginosa* cultured in SCFM2²². To determine whether the concentration of iron might account for the differential production of siderophores observed between SCFM2 and LB media, inductively coupled plasma mass spectrometry (ICP-MS) was performed on media aliquots. ICP-MS analysis revealed that SCFM2 contained a mean iron concentration of 5.25 µM (standard deviation of 0.52 µM) while LB contained a similar mean concentration of 4.73 µM (standard deviation of 0.62 µM) (Supplementary Fig. S6). Therefore, factors other than iron availability likely play a role in the expression of the pyochelin and ornibactin biosynthetic gene clusters. QS systems have been previously implicated in regulating production of both pyochelin and ornibactin, with the CepR transcriptional regulator repressing production of ornibactin and the CepR2 regulator activating production of pyochelin in *B. cenocepacia* H111⁷⁷. In our experiment, production of siderophores was observed to be dependent on strain and nutritional environment. The production of metabolites from the pyochelin pathway was further induced by the antibiotic trimethoprim, while ornibactins were not. Thus, a specific chemical cue in SCFM2 in presence of trimethoprim might play a role in induction of pyochelin production under these conditions and warrants detailed investigation in the future with knock strains that lack QS circuitry. This observation is noteworthy as pyochelin production by *B. cepacia* was previously suggested to be correlated with morbidity and mortality in patients with CF⁷⁸. Mechanistic investigations into selective induction of siderophore biosynthesis pathways are critical to understanding their relevance to infections, and as such require further inquiry.

N-acyl-homoserine lactones production in LB compared to SCFM2

QS mediates bacterial response to changing environmental conditions through cell-density dependent global changes in gene expression^{79,80}. *Burkholderia* can utilize QS to coordinate various metabolic processes and modulate cellular phenotypes such as swarming, aggregation, spatial structuring, and biofilm formation^{81,82}. QS also plays an important role in infection by regulating genes involved in virulence, and so establishing how the external environment influences production of QS signals is important to understanding their role in pathogenesis⁸³⁻⁸⁵. The CeiIR and CciIR QS systems have been described in *B. cenocepacia* strains which primarily produce and sense *N*-octanoyl-homoserine lactone (C8-AHL, **5**) and *N*-hexanoyl-homoserine lactone (C6-AHL) respectively. In addition, an orphan CeiR2, which is inactivated by C8-AHL, is also present in these strains^{77,86-88}. The gene for the orphan CeiR2 lacks adjacent gene required for synthesis for *N*-acyl-homoserine lactone. In a previous untargeted metabolomic experiment, we observed production of a wide diversity of AHLs by *Burkholderia* spp. grown in LB media, as well as their corresponding acyl-homoserine products formed by hydrolysis of the lactone ring (referred to hereafter as “hydrolyzed AHLs”)⁴³. In this study, we queried whether these signals are differentially detected in SCFM2 compared to LB media. The C8-AHL (*m/z* 228.160, **5**) was detected in both LB and SCFM2 media, and its hydrolyzed form (**6**) was detected in LB media alone (Fig. 4). Additionally, we detected hydrolyzed C13-AHL (**7**), hydrolyzed C13-AHL:1db (**8**), the sodium adduct of hydrolyzed C13-AHL (**9**), and the sodium adduct of hydrolyzed 3-OH C13-AHL (**10**) exclusively in SCFM2 when trimethoprim was present (Fig. 4). Thus, the production of these C13-AHLs were induced by trimethoprim only in SCFM2. Detection of C8-AHL, hydrolyzed C8-AHL and hydrolyzed C13-AHL was confirmed using commercial AHL standards. These AHL standards were treated with sodium hydroxide to promote hydrolysis of the lactone ring to verify the detection of hydrolyzed AHLs (Supplementary Fig. S7). Naturally produced AHLs with an odd number of carbons in the acyl sidechain are relatively rare, and their functions are not well-characterized⁸⁹. In a previous untargeted metabolomics study of 10 different *Burkholderia* strains grown in LB media, we detected hydrolyzed 3-oxo-C13-AHL:1db exclusively in extracts of *B. thailandensis* E264⁴³. To our knowledge, the present study represents the first description of C13 AHLs being produced by *B. cenocepacia*. Further studies will be needed to explore the function and biochemical basis for production of C13-AHLs by *B. cenocepacia* in SCFM2 media, and identify the mechanism by which trimethoprim upregulates the production of this AHL.

Fragin and pyrazine secondary metabolites

Fragin (**11**) is a diazeniumdiolate metallophore with antifungal activity that is produced by *Burkholderia* and *Pseudomonas* spp.⁹⁰. In a previous untargeted metabolomics study, we reported significantly increased production of fragin and its structural analogs in *B. cenocepacia* K56-2 cultures grown in LB media when supplemented with the antibiotic trimethoprim⁴³. Fragin production was only observed in the K56-2 strain, despite the fact that the *ham* gene cluster responsible for fragin biosynthesis is identical in the closely related J2315 and C5424 strains. This result highlighted that even genetically similar *Burkholderia* strains can exhibit markedly different responses to external stimuli, such as trimethoprim exposure. In the current study, fragin was detected in both media conditions, but the majority of other nodes in the cluster (11/18) were detected exclusively in either LB or SCFM2 media (Fig. 5a). These

analogs differ in the length of the acyl group added by the HamF enzyme, likely a result of differential availability of compounds containing variable acyl chain lengths which act as substrates for HamF. The MS² spectra of these analogs showed a characteristic fragment corresponding to the loss of NO group (29.998 Da), that was also observed in the MS¹ spectrum as an in-source fragment⁴³. Fragin analogs with differential production in the two media types include nodes with *m/z* 302.243 (and the corresponding in-source fragment with *m/z* 272.246), 316.223 (in-source fragment with *m/z* 286.225), and 318.239 (in-source fragment with *m/z* 288.240) (Fig. 5a).

We discovered another cluster in our molecular network containing several molecules which, like fragin, are exclusively detected in K56-2 samples and show increased production in the presence of trimethoprim (Fig. 5b). These metabolites were reported by our group in a previous study as unknown metabolites⁴³. Here we employed the recently developed CANOPUS tool to classify these unknown compounds into ClassyFire chemical classes, leading to their annotation as pyrazines^{91,92}. This information led us to conduct a literature search of pyrazines produced by *Burkholderia*, enabling annotation of these metabolites as pyrazine *N*-oxides (PNOs), which was verified using standards provided by Li and colleagues (Supplementary Figs. S8, S9)⁹³. The expression of *pvfB* and *pvfC* genes in animal pathogen *Pseudomonas entomophila* L48 and the plant pathogen *Pseudomonas syringae* pv. *syringae* UMAF0158 were previously shown to lead to production of a family of PNOs, namely PNO B (**12**, *m/z* 181.135), PNO A (**13**, *m/z* 197.126), and dPNO (**14**, *m/z* 199.145)⁹³. The *pvfB* and *pvfC* genes are homologous to the *Burkholderia* genes *hamC* and *hamD* present in the biosynthetic gene cluster of fragin. Disruption of *pvfA-D* cluster in animal pathogen *Pseudomonas entomophila* L48 and the plant pathogen *Pseudomonas syringae* pv. *syringae* UMAF0158 was shown to significantly reduce the virulence of these strains⁹³. Subsequent studies found that the *pvf* gene cluster is involved in synthesis of a signaling molecule which regulates production of small molecule virulence factors such as monalysin in *P. entomophila* and mangotoxin in *P. syringae*⁹⁴⁻⁹⁶. Both dPNO and PNO B appeared in the feature-based molecular network, while PNO A did not due to low abundance. However, a node corresponding to PNO A was observed when data was analyzed with a classical molecular network (Fig. 5b). Two additional nodes in this network were putatively annotated as 2-isopropyl-3-methoxypyrazine (*m/z* 153.102, **15**) and 2,5-diisopropylpyrazine (*m/z* 165.138, **16**) based upon available literature of bacterially produced pyrazines (Fig. 5b, Supplementary Fig. S9)⁹⁷. The MS² spectra of these related molecules did not have fragment ions in common, and so molecular networking methods alone failed to highlight the structural relatedness of these three compounds. Nevertheless, CANOPUS enabled us to independently predict that each of these compounds were pyrazines, ultimately leading to their annotation.

These observations reveal that the *ham* gene cluster that is responsible for fragin biosynthesis also leads to production of PNOs in *B. cenocepacia*. PNO production was also induced by the addition of trimethoprim in both the SCFM2 and LB culture media. Thus, unlike the siderophores described above which showed media-dependent induction by trimethoprim, fragin and PNOs are similarly induced by antibiotic trimethoprim in both LB and SCFM2. The ability of trimethoprim to induce production of fragin and pyrazines including PNOs in both SCFM2 and LB media highlights that antibiotics can serve as

signaling molecules capable of significantly modulating expression of the genes encoding virulence factors across multiple nutritional environments.

Differential detection of lipids in LB and SCFM2

The overall lipid composition of bacterial cells has been reported to be influenced by several local environmental factors, including pH, nutrient availability, oxygen levels, temperature, and buildup of metabolic waste products⁹⁸. Several clusters with hits to lipids in the GNPS spectral database were detected at higher levels in bacterial extracts grown in SCFM2, including hopanoids, phytomonic acid, monosaturated monoacylglycerols (MGs), and phosphatidylethanolamines (PEs), described below (Fig. 6).

The annotation of hopanoid cluster was performed by first searching for candidate features with m/z calculated for known bacterial hopanoids. Identified candidate features were then verified by comparing experimental MS² spectra to spectra previously published in literature^{99,100}. Once annotations were supported through spectral matching, these features were used to further propagate annotations to connected nodes within the molecular network based on mass differences. Hopanoids are pentacyclic triterpenoids frequently found in bacterial membranes^{101,102}. These polycyclic lipids are structurally analogous to eukaryotic sterols and are thought to have similar functions in regulating fluidity, permeability, and stabilization of bacterial membranes. Hopanoid biosynthesis has been previously reported in *B. cenocepacia*, where they are involved in resistance to low pH and antibiotics while also being important for swimming and swarming motility^{101,102}. A metabolite feature with m/z 708.540 is annotated as bacteriohopanetetrol (BHT) cyclitol ether (**17**) and was detected only in extracts of SCFM2 (Fig. 6a, Supplementary Fig. S10a)^{99,102}. Production of BHT cyclitol ether has been previously reported in *B. cenocepacia*, and our predicted annotation was supported by comparing experimental MS² spectra with previously published spectra available in literature^{99,102}. In addition, we also annotated the node with m/z 706.526 as unsaturated BHT cyclitol ether (**18**) (Fig. 6a). Both Δ^6 and Δ^{11} monosaturated BHT analogs have been previously characterized in bacteria¹⁰⁰. Although our mass spectrometry analysis alone is insufficient to pinpoint the location of this unsaturation, we have putatively annotated this feature as bacteriohop-6-enetetrol cyclitol ether (**18**) since unsaturation at this location has been reported in *B. cepacia* strains^{102,103}. This feature was detected in all three *B. cenocepacia* strains when grown in SCFM2 media with the highest intensity observed in the K56-2 strain, but when cultured in LB it was only detected in J2315 cultures at low levels. Annotation of another feature with m/z 724.535 is consistent with a gain of a hydroxyl group from BHT cyclitol ether (**17**). This feature was exclusively detected in K56-2 strains grown in SCFM2 media and the associated MS² spectra indicates this feature is likely bacteriohopanepentol (BHP) cyclitol ether (**19**) (Supplementary Fig. S10b). BHP derivatives have been fully characterized only in *Acetobacter* spp., *Azotobacter vinelandii*, and *Nostoc* spp., although our observation is supported by prior evidence suggesting BHP derivatives are produced by *B. cepacia* strains as well¹⁰³⁻¹⁰⁶. The feature with m/z 722.520 was detected in all three *B. cenocepacia* strains (largely in K56-2 samples) grown in SCFM2 media, which is presumably bacteriohop-6-enepentol cyclitol ether (**20**)

although the location of unsaturation cannot be unambiguously determined as mentioned above. Next, a feature with m/z 750.551 was detected exclusively in SCFM2 containing cultures with a mass difference of 42.011 (C_2H_2O) from BHT cyclitol ether representative of acetylation. The final feature in the hopanoid cluster with m/z of 748.543 is annotated as unsaturated analog of the acetylated BHT cyclitol ether (m/z 750.551), with the unsaturation most likely occurring at the C6 position.

Another metabolite in the lipid family is annotated as phytomonic acid (**21**) based on the spectral match to the MS^2 spectrum in the GNPS database. This annotation was confirmed using a commercial analytical standard (Fig. 6b, Supplementary Fig. S11a). Phytomonic acid was detected in both LB and SCFM2 media, although consistently higher levels were detected in SCFM2 media for all three strains (Supplementary Fig. S11b). Production of cyclopropane acids have been previously reported in *B. multivorans*¹⁰⁷. Cyclopropane fatty acids such as phytomonic acid are suggested to regulate membrane fluidity and stability and have been shown to increase extracellular survival in acidic or under conditions of high osmolarity^{108,109}. These lipids also are major components of the membranes of intracellular pathogens such as *Brucella abortus* and *Mycobacterium tuberculosis*^{108,109}. It is interesting to note that similar to *B. abortus* and *M. tuberculosis*, *Burkholderia* spp. are also capable of intracellularly infecting macrophage cells. The specific role of cyclopropane fatty acids in *B. cenocepacia* virulence requires further investigation.

Multiple monounsaturated monoacylglycerol (MG) lipids including nonadecenoyl-glycerol (19:1 MG, **22**), octadecenoyl-glycerol (18:1 MG, **23**), heptadecenoyl-glycerol (17:1 MG, **24**), hexadecanoyl-glycerol (16:1 MG, **25**), and the sodium adduct of heptadecenoyl-glycerol (17:1 MG, **26**) were detected only during growth in SCFM2. Annotations for this class of lipids was performed with a commercial standard of octadecenoyl(d7)-glycerol (18:1-d(7) MG) (Fig. 6c, Supplementary Fig. S12). For both 19:1 MG (**22**) and 17:1 MG (**24**), two unique metabolic features were detected with identical masses and MS^2 patterns but slightly different retention times, possibly corresponding to distinct *cis/trans* isomers. *Burkholderia* spp. are known to produce the lipase LipA, which yields monoacylglycerols as a product during degradation of di- and tri-acylglycerides^{110,111}. In *B. cenocepacia*, production of the LipA lipase is induced as part of the CepIR quorum sensing system^{86,112}. Due to the ability of monoacylglycerols to destabilize bacterial cell membranes, several have been reported to demonstrate antimicrobial activities which vary based on chain length and degree of unsaturation^{113,114}.

Finally, several phosphatidylethanolamines (PEs) were detected in this study, all consistently detected under similar conditions. In Gram-negative bacteria, the inner leaflet of the outer membrane is made up of phospholipids, of which PEs are the major component¹¹⁵. These PEs varied by the length of their acyl-chains, and were annotated as 2-OH-PE(16:1) (**27**), 2-OH-PE(16:0) (**28**), 2-OH-PE(17:1) (**29**), 2-OH-PE(18:1) (**30**), 2-OH-PE(18:0) (**31**), and 2-OH-PE(19:1) (**32**) based on GNPS library matching and manual annotations based on characteristic MS^2 fragmentation patterns described in literature (Fig. 6d)¹¹⁶. Production of 2-OH-PEs has been observed in *B. cepacia*, and is reported to increase during stress observed under growth in high temperature¹¹⁷.

Several compounds that were differentially detected between growth in SCFM2 and LB were found to be related to trimethoprim as evidenced by MS² spectral similarity. As previously described, MS2LDA was used to discover metabolites that contained trimethoprim as substructure. Thus, being higher in molecular weight than trimethoprim, these metabolites represent biochemical transformations of trimethoprim itself by *B. cenocepacia* bacteria⁴³. In this study, trimethoprim substructure was annotated using MS2LDA as Mass2Motif 543. We compared the presence of these metabolites in the extracts of bacteria cultured in LB and SCFM2 (Fig. 7). Metabolism of trimethoprim was carried out by only the pigmented strains J2315 and C5424 and not by the non-pigmented K56-2 strain, as reported previously⁴³. We generated a volcano plot of molecules containing the trimethoprim motif to visualize compounds which were differentially detected across the two media conditions (Fig. 7a). This analysis revealed that majority of the trimethoprim metabolites showed significantly higher production in SCFM2 as compared to LB. Structural characterization of these metabolites will provide insight into the pathways used by pigmented Bcc strains to metabolize xenobiotic compounds like the antibiotic trimethoprim and will facilitate future studies exploring how biotransformation will impact antibacterial activity.

Conclusion

It is well established that environmental conditions are major drivers of secondary metabolite production in bacteria. Therefore, selecting an appropriate culture media for *in vitro* bacterial growth is crucial for designing metabolomic studies relevant to the biological system of interest. In this study, we characterized the metabolomic profiles of three clinical *B. cenocepacia* isolates in SCFM2 and LB media. Sublethal concentrations of trimethoprim has previously been shown to induce secondary metabolite production in *Burkholderia* spp. Thus, culturing was also performed with and without this antibiotic to understand how trimethoprim-induced metabolomic responses vary under CF relevant environmental conditions^{42,43}. We demonstrate considerable metabolic variability between SCFM2 and LB media. In particular, we report that growth in SCFM2 media upregulates production of pyochelin-type siderophores in the presence of trimethoprim while downregulating production of ornibactin siderophores compared to LB media. We also note that production of AHL quorum sensing signals is dysregulated between the two media, with C13-AHLs exclusively detected in SCFM2 in the presence of antibiotic trimethoprim. Thus, trimethoprim induces both the production of pyochelin-type siderophores and a specific AHL signal in SCFM2 media and not in LB. Moreover, we observed that metabolism of trimethoprim itself is significantly upregulated in SCFM2 media compared to LB. Finally, we show that several lipid families (including hopanoids, cyclopropane fatty acids, monoacylglycerols, and phosphatidylethanolamines) exhibit increased production in SCFM2 compared to LB. While further work is needed to fully understand the biochemical mechanisms underlying these findings, this study provides insight into variable production of secondary metabolite production by *Burkholderia cenocepacia* spp. and is important for delineating personalized metabolotypes of different strains of the same species in an *in vitro* model of CF.

Declarations

Data Availability

The data obtained in this study has been deposited with the Mass Spectrometry Interactive Virtual Environment (MassIVE) with the identifier MSV000087793 and is accessible at <ftp://massive.ucsd.edu/MSV000087793/>.

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Author Contributions

Conceptualization—N.G.; methodology—N.G. and A.C.M.; formal analysis—A.C.M., O.J., and N.G.; data curation—D.N.F and A.C.M.; writing—A.C.M. and N.G., review—O.J., supervision—N.G. All authors have read and agreed to the submitted version of the manuscript.

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Figures

Figure 1

Summary of a) experimental design and b) data analysis workflow used in this study.

Figure 2

Overview of global metabolomic profiles across experimental conditions. a) Euler Diagram showing distribution of metabolomic features across three *B. cenocepacia* strains used in this experiment, regardless of culturing conditions, time of harvest, or extraction methods used. b) Euler diagram displaying features detected at 48 hours with ethyl acetate liquid-liquid (EtOAc) or solid-phase extraction methods (SPE), regardless of other experimental conditions. c) UpSet plot displaying number of features detected in each combination of bacterial strain, growth media, and trimethoprim exposure condition. For the sake of clarity, only the top forty combinations of conditions in which the most unique features were detected are shown.

Figure 3

Analysis of pyochelin biosynthetic metabolite production across three *B. cenocepacia* strains. Pie charts within nodes correspond to relative abundance in each strain obtained via integration of area under the chromatographic peak (green: K56-2, purple: C5424, brown: J2315) and media condition (darker shade for SCFM2 and lighter shade for LB). The border color, red vs. blue, corresponds to the presence vs. absence of trimethoprim, respectively. The production of these compounds was only detected in SCFM2 media, and in the presence of the antibiotic trimethoprim. a) Annotated structural analogs of pyochelin, including thiazostatin A/B (1) and HPTzTn-COOH (2). b) Annotated shunt products of pyochelin biosynthesis, including aeruginic acid (3) and aerugine (4).

Figure 4

Distribution of N-acyl-homoserine lactones (AHL) and their hydrolyzed forms detected across three *B. cenocepacia* strains in both LB and SCFM2 media. Pie charts on nodes correspond to relative abundance in strain and media condition, outlines correspond to relative abundance in presence/absence of trimethoprim. The structures for C8-AHL (5), hydrolyzed C8-AHL (6), hydrolyzed C13-AHL (7), and hydrolyzed C13:1db-AHL (8) as well as sodium adducts of hydrolyzed C13-AHL (9) and hydrolyzed 3-OH-C13-AHL are shown (10).

Figure 5

Distribution of fragin and pyrazine metabolites in SCFM2 and LB media. Pie charts on nodes correspond to relative abundance in strain and media condition, outlines correspond to relative abundance in presence/absence of trimethoprim. a) Annotated fragin (11) cluster, which includes structural analogs and their in-source fragment ions which are characteristic of the diazeniumdiolate moiety. b) Annotated nodes corresponding to pyrazines. The structures of pyrazine N-oxides PNO B (12), PNO A (13), dPNO (14), 2-isopropyl-3-methoxypyrazine (15) and 2,5-diisopropylpyrazine (16) are shown.

Figure 6

Distribution of various lipids detected across three *B. cenocepacia* strains in both LB and SCFM2 media. Pie charts within nodes correspond to relative abundance in each strain (green: K56-2, purple: C5424, brown: J2315) and media condition (darker shade for SCFM2 and lighter shade for LB). The border color (red vs. blue) correspond to relative abundance in presence/absence of trimethoprim. a) Annotated hopanoid cluster, including bacteriohopanetetrol (BHT) cyclitol ether (17), bacteriohop-6-enetetrol cyclitol ether (18), bacteriohopanepentol (BHP) cyclitol ether (19), and bacteriohop-6-enepentol cyclitol ether (20). Additionally, an acetylated analog of BHT cyclitol ether (m/z 750.551) and an acetylated analog of bacteriohop-6-enetetrol cyclitol ether (m/z 748.543) were detected. b) Node corresponding to phytomonic acid (21). c) Annotated cluster and nodes representing monoacylglycerols (MGs). Detected MGs vary by number of carbon units on fatty acid chain, and include MG(19:1) (22), MG(18:1) (23), MG(17:1) (24), MG(16:1) (25), and the [M+Na] adduct of MG(17:1) (26). d) Clusters corresponding to phosphatidylethanolamines (PEs). The PEs varied by acyl chain length, and were annotated as 2-OH-PE(16:1) (27), 2-OH-PE(16:0) (28), 2-OH-PE(17:1) (29), 2-OH-PE(18:1) (30), 2-OH-PE(18:0) (31) and 2-OH-PE(19:1) (32).

Figure 7

Distribution of metabolites containing trimethoprim motif which were produced by pigmented *B. cenocepacia* strains (C5424 and J2315) in SCFM2 and LB media. a) Volcano plot showing trimethoprim metabolites which are differentially detected when cultured in either LB (right) or SCFM2 (left). Metabolites exhibiting an absolute fold change greater than 2 and a p value less than 0.05 (indicated by pink dots) were considered differentially detected. b) Venn diagram showing the distribution of trimethoprim metabolites in C5424 and J2315, cultured in SCFM2 and LB media.

Supplementary Files

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- [SupplementalInformationFinal.pdf](#)