Full title

Pan-genome study underling the extent of genomic variation of invasive *Streptococcus pneumoniae* in Malawi

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Abstract

Streptococcus pneumoniae is a common cause of acute bacterial infections in Malawi. Understanding the underlying molecular mechanisms of the pathogen's invasive behavior is crucial for designing new therapeutic strategies. We conducted a pan-genome analysis to identify potential virulence genes in *S. pneumoniae* by comparing the gene pool of the isolates from carriers' nasopharyngeal secretions to isolates from the blood and cerebrospinal fluid of patients. We identified 56 serotypes in the cohort. Most serotypes showed a similar prevalence in carriage and disease groups. However, serotypes 1, 5, and 12F were disproportionately more commonly detected in specimens from patients compared to the carriage group, as expected, given the known invasiveness of these serotypes. These serotypes were genetically distinct from other isolates of other serotypes. The lack of genes from a genomic island known as RD8a was the most pronounced difference between serotypes 1, 5, and 12F compared to significantly prevalent serotypes in the nasopharynx. Genes in RD8a are involved in binding to epithelial cells and performing aerobic respiration to synthesize ATP through oxidative phosphorylation. The absence of RD8a from serotypes 1, 5, and 12F may result in a short duration in the nasopharynx, theoretically because of decreased capacity to bind to epithelial cells and access free oxygen molecules for aerobic respiration (needed to maintain the carriage state). Available ATP may be lower in isolates without RD8a, like serotypes 1, 5, and 12F, causing them to alternatively use phosphotransferase systems to transport carbohydrates since these transporters use phosphoenolpyruvate as the energy source instead of ATP. In conclusion, serotypes 1, 5, and 12F, the most prevalent invasive pneumococcal strains in Malawi, showed a considerable genetic distinction from other strains that may explain their invasiveness.

Author summary

Despite introducing the pneumococcal conjugate vaccine in 2011, *Streptococcus pneumoniae* remains a major cause of bacterial infection in Malawi. Whilst some pneumococcal strains harmlessly colonize the nasopharynx, others find their way into normally sterile sites, such as lungs, blood, and nervous system, resulting in serious illness. We found specific pneumococcal strains the most invasive in Malawi, they had a short colonization period and were genetically distinct from other strains. Their remarkable genetic divergence included the absence of several genes involved in aerobic respiration and the presence of genes that contribute to ATP-independent carbohydrate transport. The presence or absence of these genes may be one of the underlying causes of their high invasiveness with a short colonization period. This hypothesis could make these genes potential candidate targets for future therapeutic strategies. We hypothesized that the specific gene gain and/or loss in invasive vs. other serotypes are possibly associated with invasive pneumococcal disease development.

78 # Introduction

79 *Streptococcus pneumoniae,* also known as *pneumococcus*, is a Gram-positive, facultatively anaerobic
80 bacteria and is one of the leading causes of mortality worldwide. Despite reductions in the incidence of
81 pneumococcal disease in countries that introduced pneumococcal conjugate vaccines (PCV), the
82 pneumococcal mortality rate is still high. Pneumococci are estimated to be responsible for 317,300
83 deaths in children aged 1 to 59 months worldwide in 2015 (1). In the post-PCV13 era, a high disease
84 burden has still been reported in low-income countries in Africa, such as Malawi (2).

85 Although pneumococcal nasopharyngeal colonization is asymptomatic, it is a prerequisite for
86 transmission and disease development (3)(4). Symptoms appear when isolates from the nasopharynx
87 spread to normally sterile sites such as the lung, blood, and central nervous system. Depending on the
88 infected organ, *S. pneumoniae* can cause two types of infection: (i) non-invasive (mucosal)
89 pneumococcal diseases such as otitis media and sinusitis, and (ii) invasive pneumococcal diseases (IPD)
90 such as bacteremia and meningitis. IPD incidence is highest among infants, the elderly, and
91 immunosuppressed people, most likely due to their less efficient immune systems (5).

92 Pneumococci possess several virulence factors, including the polysaccharide capsule, surface proteins,
93 and enzymes (6)(7). The polysaccharide capsule is the most important virulence factor as it aids the
94 *pneumococcus* in evading the immune response during colonization and invasion [8]. Its biosynthesis is
95 regulated by a cluster of genes in the capsular polysaccharide (*cps*) locus (8)(9). Pneumococcal serotypes
96 are defined by the type and order of monosaccharides that compose the capsule structure. To date, one
97 hundred pneumococcal serotypes have been identified (10). Each strain has a set of capsular synthesis
98 genes in the *cps* locus that determine its serotype. Immunogenic properties of the capsular
99 polysaccharide were utilized to develop all pneumococcal vaccines in use, including PCV7, PCV10, and
100 PCV13 that cover 7, 10, and 13 serotypes, respectively. Although the introduction of PCVs has

101 significantly reduced the burden of disease caused by vaccine types (VTs), serotype replacement has
102 increased the non-vaccine types (NVTs) carriage rate and IPD incidence (11)(12).

103 In November 2011, PCV13 was introduced in Malawi, which markedly decreased the health system
104 burden and rates of severe childhood pneumonia (13). PCV13 includes serotypes 1, 3, 4, 5, 6A, 6B, 7F,
105 9V, 14, 18C, 19A, 19F, and 23F. A case-control study in Malawi showed vaccine effectiveness against VT-
106 IPD of 80·7% (14). A nasopharyngeal carriage survey conducted in the Karonga district showed that
107 although the vaccine reduced the VT colonization rate, a moderate level of serotype replacement was
108 observed among carriers (15). Moreover, the emergence of antibiotic-resistant pneumococci due to the
109 overuse of antibiotics is a global concern in the 21st century (16). To develop new, more effective
110 vaccines against the vaccine-escape clones and design effective drugs against antibiotic-resistant strains,
111 it is critical to understand the functions of genes involved in pneumococcal colonization and
112 pathogenesis. During the past decade, the evolution of high-throughput sequencing technologies has
113 generated enormous amounts of genomic data that have enabled researchers to perform large-scale
114 genomic analysis. A well-known example is pan-genome studies. The pan-genome is the entire gene set
115 in a collection of closely related strains within a specie (17). Determining the genetic drivers is an active
116 and promising area of research that can provide insights into pneumococcal disease prevention and
117 treatment to reduce mortality rates. The pan-genome is useful for analyzing recombinogenic pathogens
118 such as *pneumococcus*(18). The recombination level is high in thirteen pneumococcal genomic loci
119 known as regions of diversity (RDs) numbered from RD1 to RD13, some of which are involved in
120 virulence (19)(20).

121 In this study, we used whole-genome sequencing (WGS) of 1477 pneumococcal samples recovered from
122 people residing in Blantyre, Karonga, and Lilongwe in Malawi to address three complementary issues: (i)
123 identifying the serotype distribution, (ii) understanding the structure of the pneumococcal population,
124 and (iii) identifying potential invasive driver genes and their biological functions.

125 # Materials and methods

126 **Study cohort and pneumococcal isolates**

127 Pneumococcal isolates used in the study were from patients at three central hospitals in Malawi: Queen
128 Elizabeth Central Hospital in Blantyre, Karonga Central Hospital in Karonga, and Kamuzu Central Hospital
129 in Lilongwe. Samples were collected from people visiting Queen Elizabeth Central Hospital during 1997-
130 2015 (53%) and from households in the Karonga district during 2009-2014 as part of two household
131 studies (44%) (21). The remainder of the samples were collected from Kamuzu Central Hospital. Isolates
132 were obtained either from the nasopharynx of carriers or the blood and cerebrospinal fluid (CSF) of
133 patients and archived at the Malawi-Liverpool Wellcome Trust Clinical Research Programme Centre. In
134 this study, the term "sterile sites" refers to blood and CSF, and "invasive samples" are those obtained
135 from the sterile sites.

136 ## Whole-genome sequencing and quality control

137 Archived samples were sequenced under the Global Pneumococcal Sequencing project and
138 Pneumococcal African Genomic Consortium at Wellcome Trust Sanger Institute in the United Kingdom.
139 The bacterial DNA was extracted using QIAamp DNA mini kit and QIAgen Biorobot manufactured by
140 QIAGEN. The whole genome sequencing was accomplished using the Illumina Genome Analyzer II and
141 HiSeq platforms, generating 125 nucleotides paired-end reads. The quality of the reads was evaluated
142 by Fastqc (22).

143 ## In-silico serotyping, sequence typing, and quantification of serotype invasiveness

144 SeroBA version 1.23.4 was used to infer the serotype of samples (23). SeroBA applies a k-mer method to
145 determine serotypes directly from the paired-end reads in FASTQ format. Any serotype with a relative
146 frequency greater than 5% was considered as an abundant serotype. Fisher's exact test was applied to
147 identify serotypes with a significant presence in the nasopharynx and sterile sites. P-values were

148 adjusted by the Benjamini-Hochberg method, and serotypes with an adjusted p-value less than 0.01
149 were considered significant. The odds ratio (OR) was calculated as follows OR = (*ad*)/(*bc*), where for
150 serotype *k*, a is the number of invasive serotype *k*, *b* is the number of carriage serotype *k*, *c* is the
151 number of invasive non-serotype *k,* and *d* is the number of carriage non-serotype *k*. For OR calculation,
152 zero values were replaced by 0.5. Any abundant serotype with a significant presence in the nasopharynx
153 was considered to have low invasiveness, and high invasiveness was assigned to abundant serotypes
154 with a significant presence in the sterile sites. Fisher's exact test was also applied to identify serotypes
155 whose frequencies changed significantly after introducing PCV13 in 2011 (adjusted p-value < 0.01).

156 **Genome assembly and annotation**

157 Genomes were assembled using Velvet Optimiser version 2.2.5 (24). The assembler was set to produce
158 contigs longer than 500 base pairs. The starting hash value was 61, and the end hash value was 119. The
159 quality of the assembled genomes was evaluated by Quast version 5.2(25). The assembled genomes
160 were annotated using Prokka version 13.1 (26).

161 **Pan-genome construction**

162 The pan-genome for samples was generated using Roary version 3.12.0 (27). Roary was run to perform
163 the core gene alignments with MAFFT version 7.313 (28). Genes in the pan-genome were divided into
164 three groups based on their abundance among samples. Genes that existed in 100% of samples were
165 considered as core genes, those that were not core but present in more than 95% of samples were
166 regarded as a soft-core genes, and the rest were considered as accessory genes.

167 **Analysis of the population structure**

168 Since core genes are conserved between all samples, diversity in the core genome exists in the form of
169 small-scale variations such as single nucleotide polymorphisms (SNPs) and short indels. SNPs and indels
170 distribution in the core-genome was analyzed to gain insight into the population structure. The genetic

171 separation between samples was illustrated by the phylogenetic tree built from SNPs and indels in the
172 alignment of core genes as the phylogeny markers. The core genome alignment was used as input into
173 the IQ-TREE version 2 (29) to draw a phylogenetic tree visualized by iTol version 3 (30).

174 Diversity in the accessory genome exists in the form of large-scale gene presence-absence variations.
175 The R package Nonnegative Matrix Factorization (31) visualized the pan-genome matrix as a gene
176 presence-absence heatmap. Furthermore, to determine which factors, including isolation sites
177 (nasopharynx, blood, and CSF), serotypes, geographical locations, and vaccination eras, have the
178 greatest impact on the gene presence-absence profile of samples, a PCA of gene distribution was
179 accomplished by the R package MixOmics version 6.20.0 (32).

180 Gene presence-absence statistical analysis

181 The phenotypic trait of samples was assigned based on the population structure and isolates'
182 invasiveness. The gene presence-absence statistical test was performed using Scoary version 1.6.1643
183 (33). This tool scores the components of the pan-genome for associations with observed phenotypic
184 traits while accounting for population stratification. The test was conducted between samples from
185 different sources and serotypes to find putative virulence factors. Every gene with a Bonferroni
186 corrected p-value less than 0.05 was reported as a significant gene.

187 Functional and gene ontology (GO) enrichment analysis

188 The list of significant genes was submitted to STRING webtool version 11.5 (34) for the functional
189 enrichment analysis. The STRING database is a network that integrates information from all publicly
190 available protein-protein interaction databases. The STRING's computational pipeline can predict direct
191 (physical) and indirect (functional) interactions between proteins. Interactions in STRING are derived
192 from 5 sources, including (i) genomic context predictions, (ii) high-throughput lab experiments, (iii) co-
193 expression, (iv) automated text mining, and (v) previous knowledge in databases. Functional enrichment
194 analysis in STRING uses information from classification systems such as the Kyoto Encyclopedia of Genes

195 and Genomes (KEGG) (35) and the Protein families database (Pfam) (36). The tool entitled "Multiple
196 Sequences" was selected, and the *S. pneumoniae TIGR4* was chosen as the reference organism. STRING
197 reports the associated enriched pathways with a false discovery rate of less than 0.05.

198 Results

199 In total, 825 isolates from the nasopharynx of healthy carriers, 368 isolates from the blood of
200 bacteremia patients, and 284 isolates from the CSF of meningitis patients were sequenced. The
201 demographics of the samples are shown in Table 1.

Table 1. Demographics of 1477 pneumococcal isolates collected in Malawi.

Characteristics	**Categories**	**Nasopharynx**	**Blood**	**CSF**
Age (in years)	< 5	538	165	141
	5-19	109	42	60
	20-40	60	67	50
	> 40	7	24	7
	Unknown	111	70	26
Sex	Female	401	131	111
	Male	313	122	122
	Unknown	111	115	51
City	Blantyre	169	357	259
	Karonga	656	0	0
	Lilongwe	0	0	23
	Unknown	0	11	2
Sampling period		2009-2014	1997-2015	2000-2015

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203 **Serotypes 1, 5, and 12F had the highest invasiveness, likely with a short period of**
204 **nasopharyngeal colonization**

205 In total, isolates were assigned to 56 serotypes. Regardless of the isolation sites, serotypes 1 (8.7%), 5
206 (7.8%), 6B (6.6%), 23F (6.3%), and 19F (5.5%) were abundant, with a relative frequency greater than 5%
207 of the entire cohort. 66% of samples were obtained before the PCV13 rollout in 2011, and 27% were
208 from the post-PCV13 era (S1 Fig).

209 Amongst the carriage isolates, serotypes 19F (7.88%), 6B (7.27%), 16F (6.79%), and 23F (5.21%) were
210 abundant. Serotype 5 (20.38%) followed by 1 (16.58%), 23F (8.42%), and 6B (5.98%) dominated the
211 blood samples, while serotypes 1 (21.13%), 5 (8.1%), 12F (7.04%), 23F (6.69%), 6A (5.63%), and 6B
212 (5.28%) were predominant in the CSF. When aggregating blood and CSF groups, the most common
213 serotypes were 1 (18.56%), 5 (15.03%), 23F (7.67%), 6B (5.67%), and 6A (5%).

214 As illustrated in Fig 1 and S1 Table, of the abundant serotypes in the blood or CSF, serotypes 1 (p =
215 1.96E-34), 5 (p = 3.97E-19), and 12F (p=5.29E-06) had a significant presence among patients with a low
216 frequency in the nasopharynx including serotype 1 (1%), serotype 5 (2%) and serotype 12F (0.6%). Given
217 that the colonization phase is a pre-requisite for infection, serotypes 1, 5, and 12F may have a short
218 period of nasopharyngeal colonization before infecting the sterile sites. Since serotypes 1 and 5 were
219 also the most frequent in the entire cohort, they could be considered as the most prevalent serotypes
220 with the highest invasiveness. In this study, the terms “significant invasive serotypes” or "hyper-invasive
221 serotypes" were assigned to serotypes 1, 5, and 12F. Comparatively, abundant serotypes 16F and 19F in
222 the nasopharynx had a significant presence among carriers suggesting that they have a lower potential
223 to cause invasive disease. Other frequently detected serotypes, such as 6A, 6B, and 23F, were abundant
224 and equally distributed among carriers and patients. They may need to colonize the nasopharynx longer
225 than the hyper-invasive serotypes before infecting the sterile sites. Serotypes 6A, 6B, and 23F were
226 previously reported as abundant and common serotypes between non-invasive carriers and invasive
227 infection (37)(38).

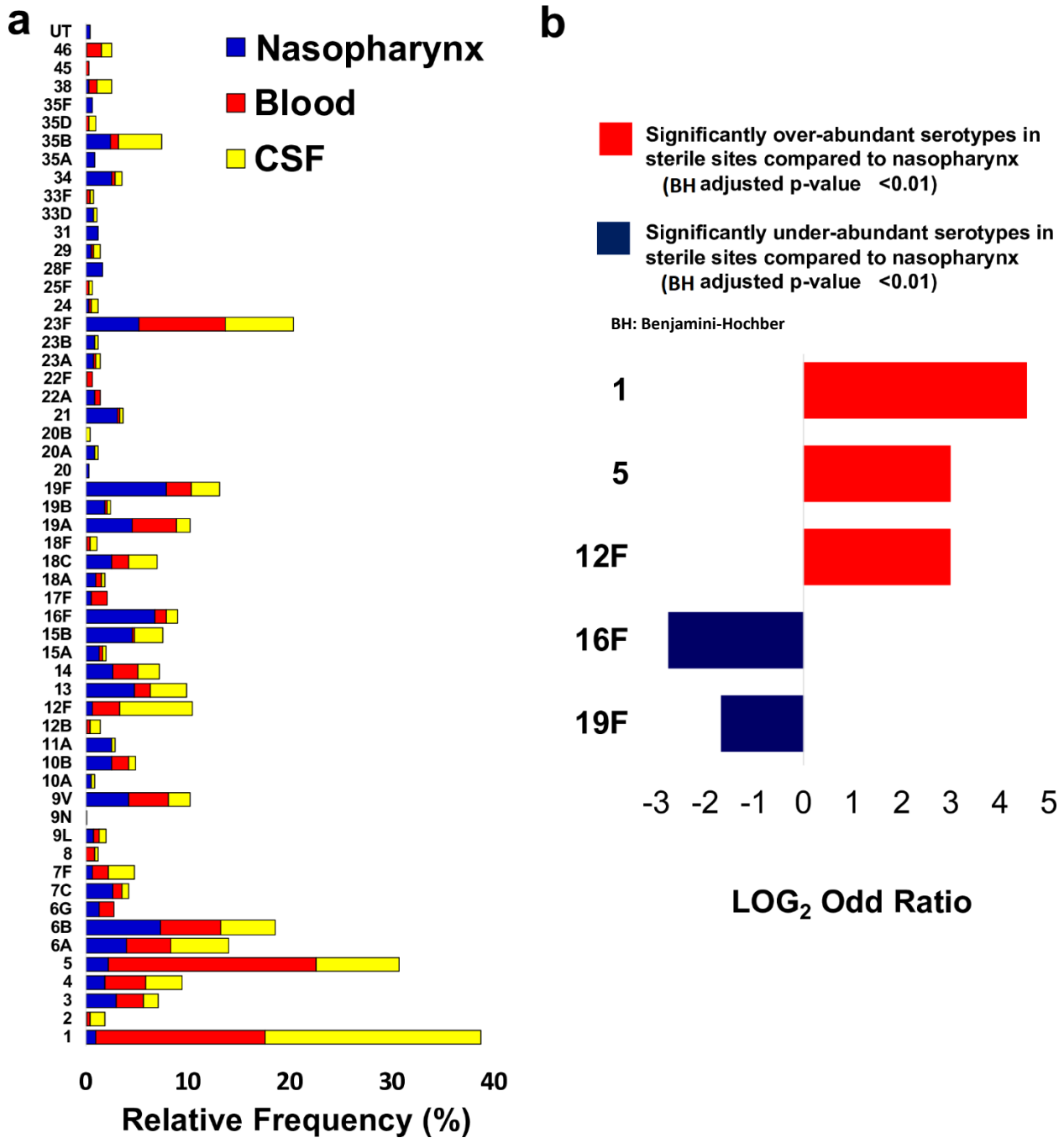

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Fig 1. The distribution of the 56 pneumococcal serotypes assigned to 1477 samples from Malawi. (a) The relative frequency of each serotype in the nasopharynx of carriers, the blood of bacteremia patients, and the CSF of meningitis patients is shown in blue, red, and yellow, respectively (UT: Un-Typeable). (b) The log-transformed odds ratio of the significantly over- and under-abundant serotypes in the sterile sites (blood and CSF). Fisher's exact test was applied to identify serotypes with a significant differential abundance among carriers and patients (nasopharynx and sterile sites) at the significance level of the Benjamini-Hochberg adjusted p-value < 0.01 (BH: Benjamini-Hochberg).

229 The temporal distribution of the hyper-invasive serotypes with respect to the vaccine rollout time was of
230 note. The relative frequency of serotype 1 significantly increased (pre-PVC13=8.6% and post-
231 PCV13=11.3%), serotype 5 significantly decreased (pre-PVC13=11% and post-PCV13=1.5%), and serotype
232 12F significantly increased (pre-PVC13=1.1% and post-PCV13=5.7%) after the vaccination program in
233 Malawi. Vaccination might be effective against serotype 5 but did not reduce the burden of IPDs caused
234 by serotype 1. Serotype 12F, not included in PCV13, may be an invasive strain emerging in the post-
235 PCV13 era. However, a more extensive dataset with more recent samples is needed to characterize the
236 long-term PCV13 effects.

237 **High diversity in the pneumococcal pan-genome**

238 For the genome assembly, the average optimized assembly hash value and the average N50 were 96 and
239 113,986, respectively. The mean assembled genome size was estimated at 2,116,779 nucleotides with a
240 standard deviation of 106,481. This length is in the range of previously reported *S. pneumoniae* genome
241 size (39). The length of the pan-genome was 5,178,167 base pairs, composed of 6,803 genes, including
242 729 core-(10.7%), 820 soft-core- (12.1%), and 5,254 accessory genes (77.2%). The pan-genome was open
243 as it did not reach sampling saturation as the number of genes increased as the sample size grew (S2
244 Fig). In Fig 2, the pan-genome is demonstrated as a gene presence-absence heatmap that shows
245 serotypes as the strongest determinant of the gene distribution in the pan-genome. The distinction was
246 evident for the hyper-invasive serotypes 1, 5, and 12F since they clustered in unique clades. The sets of
247 genes specifically present in different serotypes appear as small blue blocks in the heatmap.

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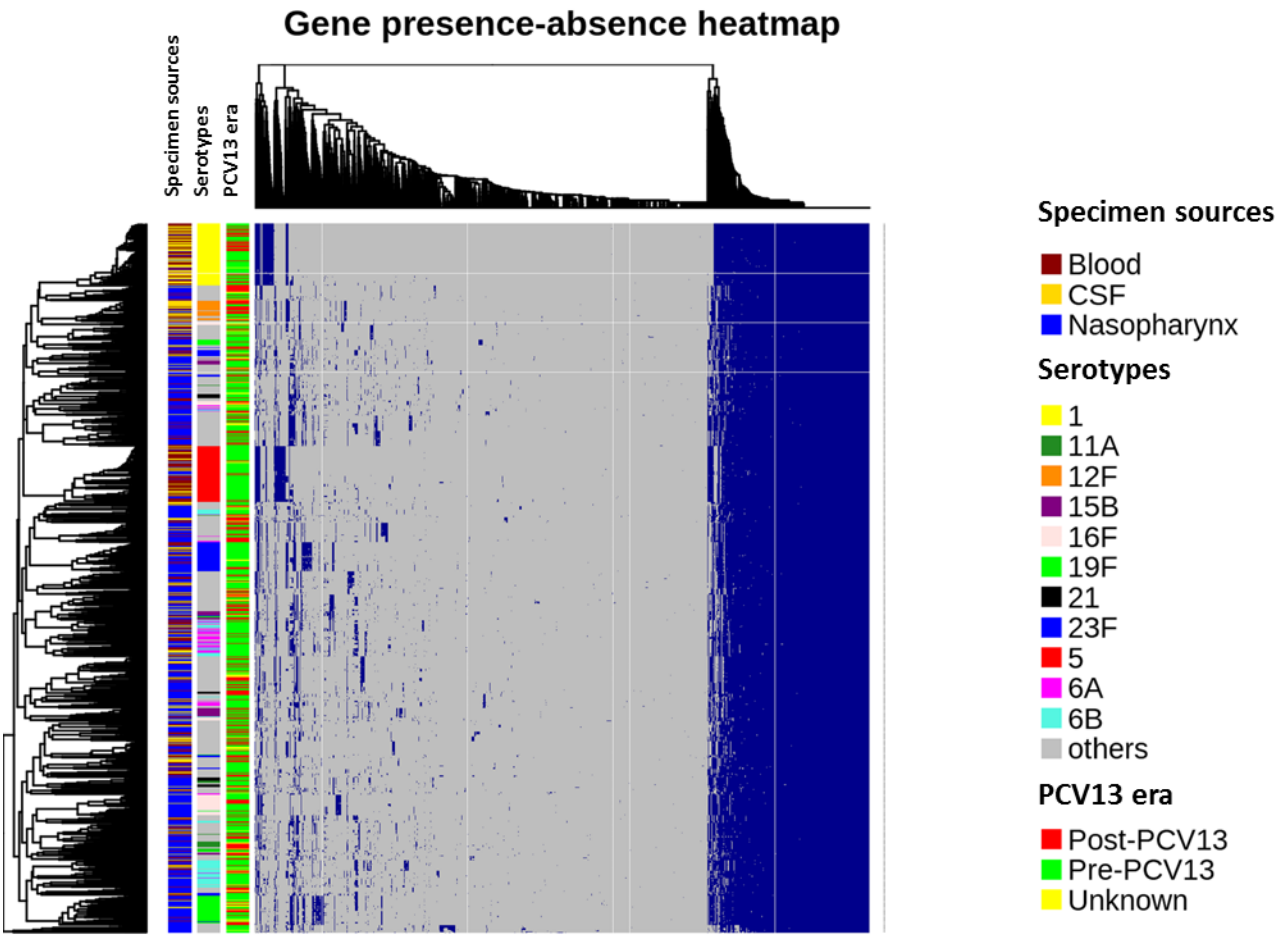

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Fig 2. The pan-genome matrix of 1477 pneumococcal isolates from Malawi. The pan-genome is visualized as a gene presence-absence heatmap representing the hierarchical unsupervised clustering of samples based on the distribution of genes in the pan-genome. Each row is a sample, and each column is a gene. A blue dot denotes the presence of each gene. On the right side of the heatmap, the large blue block shows core genes present in all samples. The left side of the heatmap represents the accessory genome along with the clustering bands. In addition to the significant serotypes 1, 5, 12F, 16F, and 19F, other abundant serotypes, including 6A, 6B, and 23F, as well as serotypes with source-based p-value < 0.05, including 21, 11A, and 15B, are also highlighted on the heatmap.

250 **The significant invasive serotypes showed the highest distinction in the core- and accessory-**

251 **genome**

252 The maximum-likelihood tree characterized by the distribution of the small-scale variants (SNPs and

253 indels) in the core-genome identified the hyper-invasive serotypes 1, 5, and 12F as monophyletic

254 clusters (Fig 3). The distinction of hyper-invasive serotypes that were separated as unique clusters was

255 higher than other abundant serotypes, such as 6B, 19F, and 23F, which appeared as multiple clusters on

256 the phylogenetic tree (Fig 3).

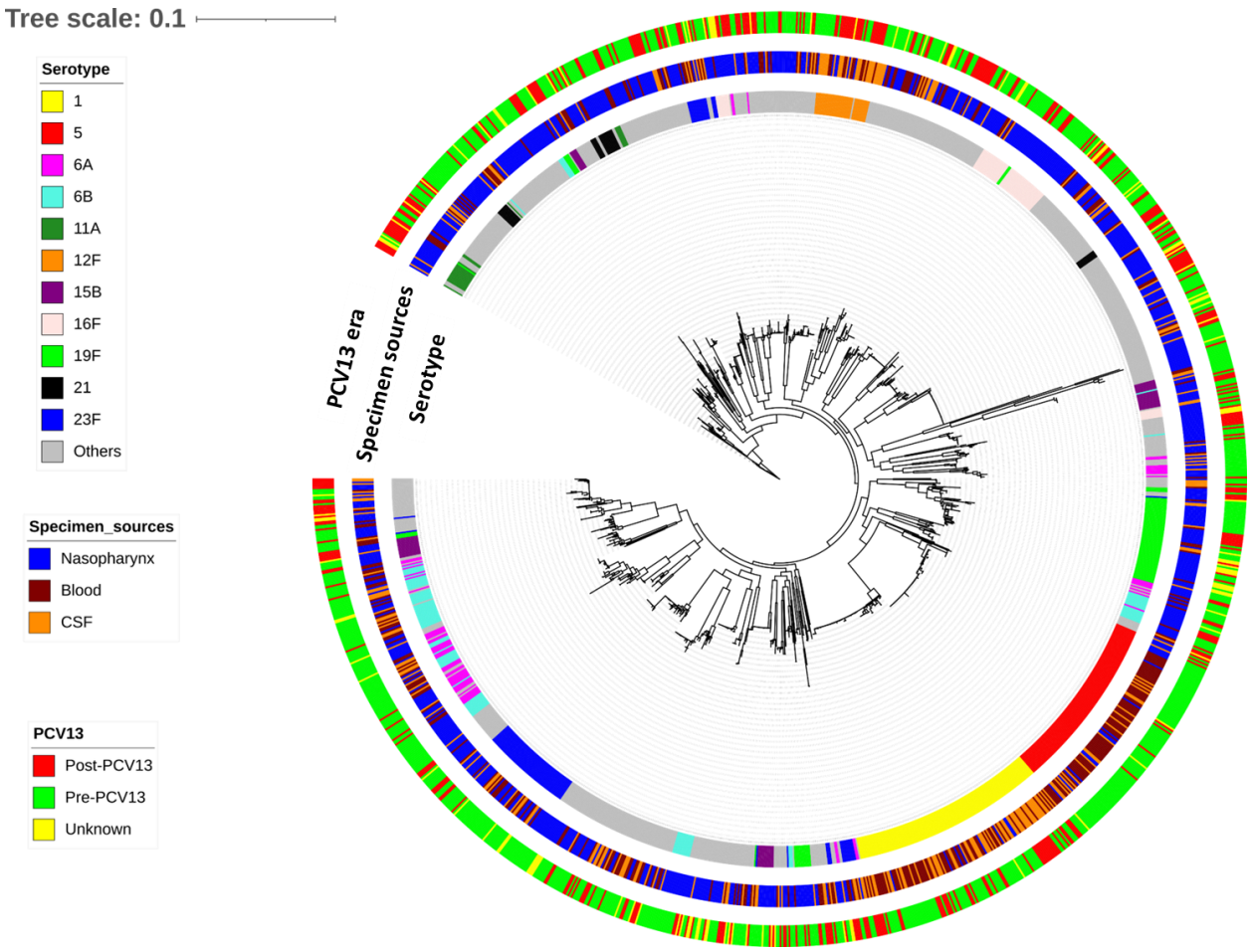

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Fig 3. The phylogenetic population structure of 1477 pneumococcal samples from Malawi. The phylogenetic tree was built based on the multiple sequence alignment of the core genome using the maximum likelihood method. Colors on the loops show the serotypes, specimen sources (isolation sites), and PCV13 eras. In addition to the significant serotypes 1, 5, 12F, 16F, and 19F, other abundant serotypes, including 6A, 6B, and 23F, as well as serotypes with source-based p-value < 0.05, including 21, 11A, and 15B, are also highlighted on the tree.

258 The PCA of large-scale variants distribution in the accessory-genome (gene presence/absence) showed

259 serotypes 1 and 5 clustered distantly from other strains (Fig 4). Meanwhile, a moderate level of

260 separation was observed for serotypes 12F, 19F, and 23F. This highlighted the distinction of serotypes 1

261 and 5 in their gene content as observed in their SNPs profiles demonstrated by the phylogenetic tree.

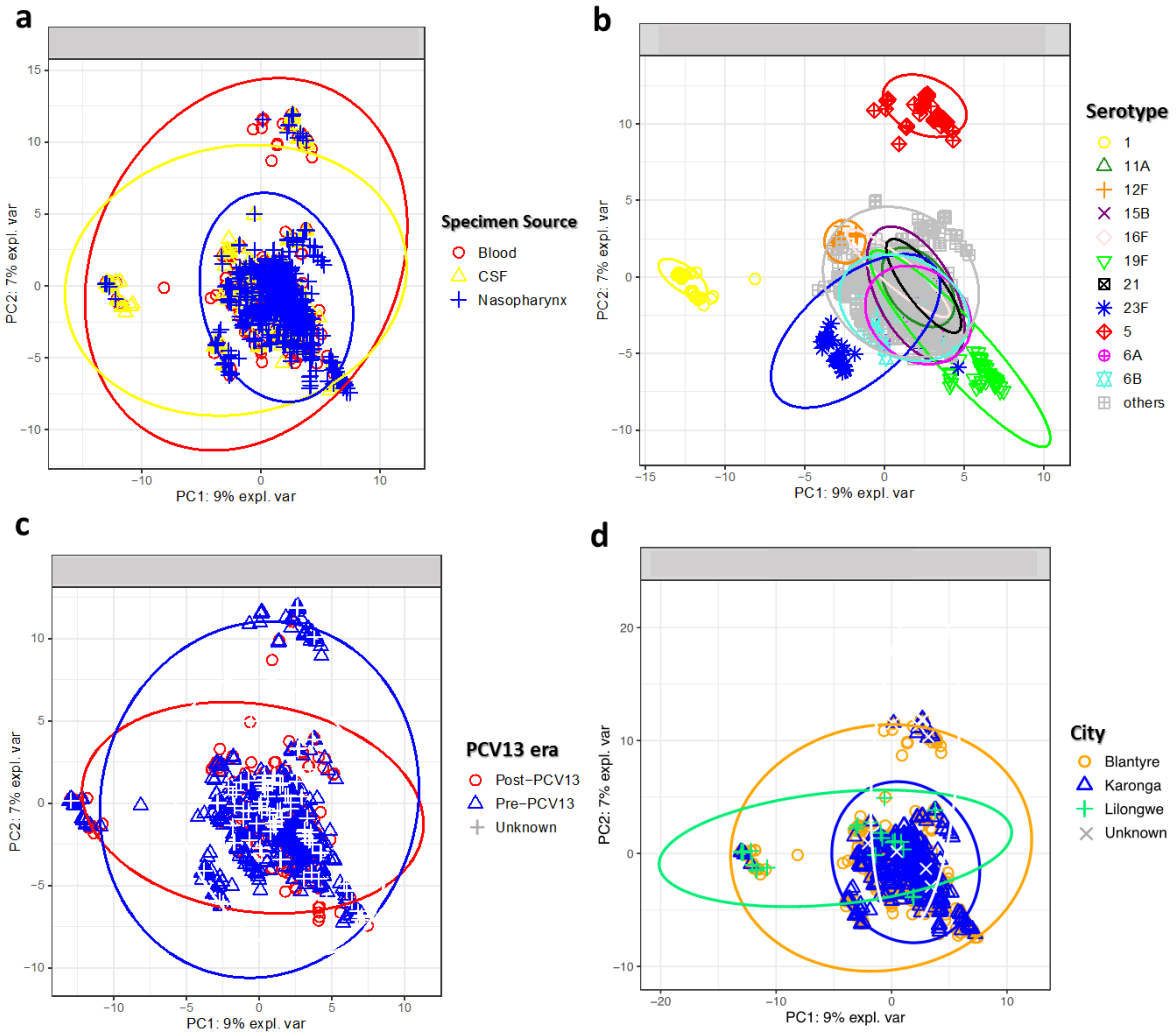

Fig 4. The PCA of the gene distribution in the pan-genome of pneumococcal isolates from 1477 Malawians. The PCA of variants (gene presence-absence) in the accessory-genome indicates the influence of (a) specimen sources (isolation sites), (b) serotypes, (c) PCV13 (vaccination) era, and (d) geographical locations on the gene presence-absence profile of pneumococcal isolates in Malawi. Serotypes 1 and 5 were clearly separated from other samples.

Fig 3 and Fig 4 show that factors such as time, locations, and isolation sites (specimen sources) could not explain the small- and large-scale genetic variants in the pan-genome. Indeed, the serotype of samples was the best driver of the population structure. Of importance was that the hyper-invasive serotypes had the maximum core and accessory distances from other strains, which means they are genetically different from other serotypes. Since these serotypes were associated with disease, their high genetic distinction may be associated with their invasive potential. Nonetheless, the analysis must rigorously

269 account for population structure and carefully curate any hits to show statistical association with the
270 disease across the population. The first thing to point out is that the genetic distinction of serotype 1
271 and 5 is not observed in 12F. This may be due to the lower number of 12F (n=35) compared to 1 (n=129)
272 and 5 (n=116). Moreover, the near-perfect separation between serotypes 1 and 5 and the other
273 serotypes may be just because these strains were hardly found in the carriage group. Therefore, these
274 patterns may primarily reflect biases in sampling rather than genuine genetic phenomena. To address
275 these issues, ten samples of each hyper-invasive serotype and the vaccine types involved in PCV13 were
276 randomly selected from the nasopharynx, blood, and CSF. The PCA of the gene distribution was
277 repeated for the downsampled dataset and identified the stratification so that each of serotypes 1, 5,
278 and 12F formed a distinct group clustered distantly from other strains (S3 Fig).

279 Typically, serotypes are separated from each other since they have different serotype-defining capsule
280 genes. The issue that must be addressed is why hyper-invasive serotypes cluster more distantly in
281 contrast with other serotypes. The hypothesis we explored is that serotypes 1, 5, and 12F gained or lost
282 a list of genes that caused their quick invasiveness. There are other abundant serotypes in the blood and
283 CSF (such as 6B and 23F), however, they have a similar prevalence in the nasopharynx, which means
284 they may stay in the nasopharynx longer than the hyper-invasive serotypes.

285 **The gene presence-absence statistical analysis**

286 The following issues could skew the gene presence-absence analysis:

287 a) The batch effect introduced by geographical locations: 85% of carriage samples were from
288 Karonga, whereas 95% of disease samples were from Blantyre (Table 1). A comparison between
289 carriage and disease groups may only identify the difference between pneumococcal genomes
290 from two geographical locations rather than between the non-invasive and invasive groups.
291 b) Study limitation: The likely presence of invasive serotypes in the carriage group made it unclear
292 which nasopharyngeal samples progressed to disease after collection. Indeed, the carriage

293 population likely contained invasive serotypes that could bias the test between carriage and
294 disease samples to identify potential virulence genes.
295 c) Population structure: The significant abundance of the hyper-invasive serotypes in the patient
296 group and their highest genetic distinction would skew the test between the carriage and
297 patient groups. The difference between carriage and disease groups would actually be the
298 difference between the carriage and hyper-invasive serotypes (and not all strains in the blood
299 and CSF).

300 To measure the geographical batch effect, the carriage isolates from Karonga were compared to
301 carriage isolates from Blantyre. The test did not detect any significant gene in either Blantyre or Karonga
302 groups, implying that the gene content of the carriage samples from Blantyre and Karonga was similar.
303 The serotype distributions in the carriage group from Karonga and Blantyre were also similar. Therefore,
304 the effect of the geographical locations on the pneumococcal genomes was not remarkable. This was
305 expected as the distance between Karonga and Blantyre is about 830 kilometers, and the race of people
306 in these cities is similar.

307 Despite the possible presence of the invasive serotypes in the carriage group (e.g., 6A, 6B, and 23F that
308 were prevalent among the carriers and patients), comparing the whole carriage and patient groups
309 could still make sense since an invasive serotype in the nasopharynx may change its genome before
310 entering the sterile sites. Indeed, the potential invasive serotypes in the nasopharynx were isolated
311 during their colonization phase when they could exchange genes with other pneumococci or bacterial
312 species through recombination and horizontal transfer of genes. Thus, the genome of an invasive
313 serotype in the nasopharynx is not necessarily identical to its genome in the blood and CSF. The
314 association test was applied to compare the whole carriage and disease group, while the hyper-invasive
315 serotypes were excluded (location-based analysis). The exclusion of the hyper-invasive serotypes
316 prevented them from skewing the difference between the carriage and patient groups. The location-

317 based analysis identified 27 significant genes including 11 genes significantly present in the blood and

318 CSF and 16 genes in the nasopharynx (Table 2 and 3, and S2 Table).

Table 2. Significant genes (p-value < 0.05) in the blood and CSF (hyper-invasive serotypes were excluded).

ID	Annotation	P-value	Odds ratio
SP_0360	Capsular polysaccharide biosynthesis protein (from RD3)	8.17E-05	4.897059
SP_0358	Capsular polysaccharide biosynthesis protein (from RD3)	8.17E-05	4.897059
SP_0351	Capsular polysaccharide biosynthesis protein (from RD3)	8.17E-05	4.897059
SP_0359	Capsular polysaccharide biosynthesis protein (from RD3)	8.17E-05	4.897059
SP_0357	Capsular polysaccharide biosynthesis protein	0.000175	4.758421
SP_1037	Type II restriction endonuclease BcgI	0.001074	2.878307
SP_1953	Bacteriocin/lantibiotic secretion ABC transporter permease protein	0.001864	4.631892
SP_0535	Putative immunity protein	0.004674	1.988523
SP_1056	Relaxase/Mobilisation nuclease domain (From RD6)	0.012604	2.932153
SP_1656	Hypothetical protein	0.014050	1.963212
SP_0347	Capsular polysaccharide biosynthesis protein (from RD3)	0.020560	1.808997

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Table 3. Significant genes (p-value < 0.05) in the nasopharynx (hyper-invasive serotypes were excluded).

ID	Annotation	P-value	Odds ratio
SP_1770	Glycosyl transferase, glyB (from RD10)	0.000132	0.502513
SP_1771	Glycosyl transferase, family 2/family 8 (from RD10)	0.012770	0.560236
SP_1763	Preprotein translocase secY family protein (from RD10)	0.017254	0.563256
SP_1765	Glycosyl transferase, glyF (from RD10)	0.018150	0.564542
SP_0939	Hypothetical protein	0.020740	0.483266
SP_1766	Glycosyl transferase, glyE (from RD10)	0.020740	0.483266
SP_1767	Glycosyl transferase, glyD (from RD10)	0.023196	0.568879
SP_1762	Accessory secretory protein asp1 (from RD10)	0.023196	0.568879
SP_1761	Accessory secretory protein asp2 (from RD10)	0.023196	0.568879
SP_1760	Accessory secretory protein asp3 (from RD10)	0.023196	0.568879
SP_1755	Hypothetical protein	0.031054	0.558554
SP_1757	Glycosyl transferase, glyB (from RD10)	0.031054	0.558554
SP_1764	Glycosyl transferase, glyG (from RD10)	0.023196	0.568879
SP_1768	Conserved hypothetical protein (from RD10)	0.031054	0.558553
SP_1758	Poly(glycerol-phosphate) alpha-glucosyltransferase, tagE (from RD10)	0.031122	0.571941
SP_1759	Preprotein translocase, secA subunit (from RD10)	0.031321	0.574557

329

330 The most significant genes in the blood and CSF were from *cps* locus (RD3) may be due to a higher level
331 of encapsulation during the disease. Other significant genes, such as SP_1953, SP_0535, SP_1037, and
332 SP_1056, contribute to toxic secretion and recombination. SP_1056 is from a region known as
333 pneumococcal pathogenicity island 1 (PPI1), located in RD6. This gene encodes a mobilization protein
334 required for the horizontal transfer of genes and plasmids through bacterial conjugation. SP_1056 forms
335 the relaxation complex or relaxosome with the help of other enzymes (40).

336 Significant genes in the nasopharynx (absent from samples in the blood and CSF) were from RD10, which
337 is known as SecY2A2 island and is responsible for the secretion of pneumococcal serine-rich repeat
338 protein or PsrP (S4 Fig)(41). RD10 contains several glycosyltransferases and secretory components that
339 were significantly absent from the genome of samples collected from blood and CSF.

340 To explore the divergence of the hyper-invasive serotypes, they were compared to serotypes 16F and
341 19F, which were significantly present in the nasopharynx (serotype-based analysis). Serotypes 16F and
342 19F could represent non-invasive strains better than the whole nasopharyngeal population that
343 potentially contained some invasive serotypes. Indeed, it was a test between serotypes with the highest
344 and lowest invasiveness to characterize the genomes of serotypes 1, 5, and 12F. The gene gain and loss
345 profiles in the hyper-invasive serotypes may include components contributing to their virulence and
346 short colonization period. The serotype-based analysis identified 184, 157, and 186 significant genes in
347 serotypes 1, 5, and 12F, respectively (S3 Table, S4 Table, and S5 Table) that were much larger than the
348 number of significant genes identified by the location-based analysis. The functional enrichment analysis
349 identified the phosphotransferase system (PTS, KEGG ID: spn02060) as over-represented and oxidative
350 phosphorylation (KEGG ID: spn00190) as under-represented pathways in the hyper-invasive serotypes
351 (p-value < 0.05).

352 In total, there were 18 significant genes jointly present in the hyper-invasive serotypes (Fig 5.a),
353 including elements of the PTSs that transport sucrose and lactose across the membrane (SP_0302,
354 SP_0303, SP_0304, SP_0305, SP_0306, SP_0308, SP_0309, and SP_0310), bacteriocins (SP_0544 and
355 SP_1051) and a permease protein (SP_1527). The over-represented pathway (spn02060) was associated
356 with significant genes that code for PTS transporters involved in carbohydrate metabolism. Seven genes
357 were unannotated. The PTS transporters genes were also present in a high proportion of abundant
358 serotypes in sterile sites such as 6B (67%) and 23F (86%).

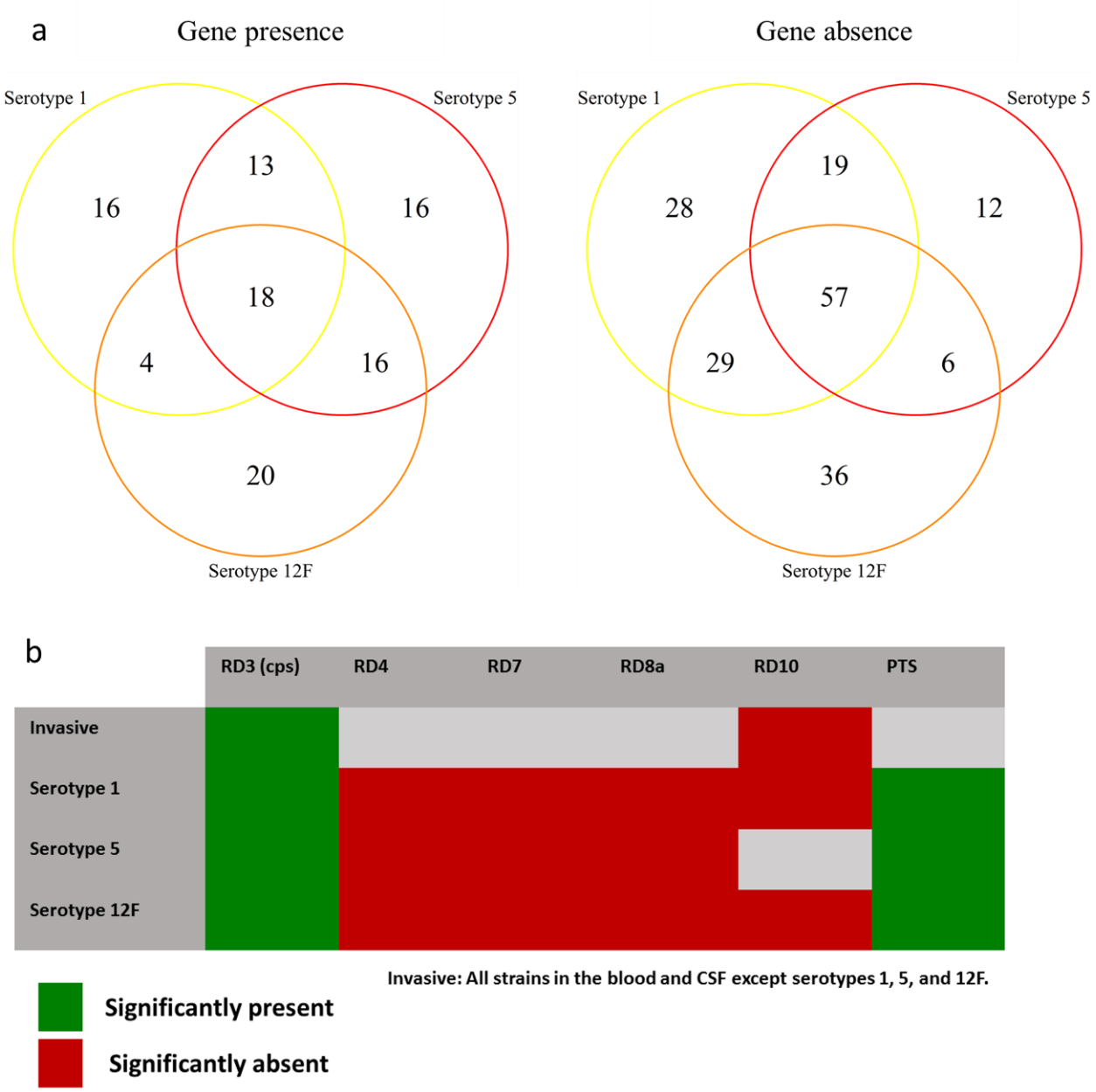
359 There were 57 significantly absent genes from serotypes 1, 5, and 12F (Fig 5.a). The most significant
360 ones were from RD8a, a region composed of two operons, RD8a1 (SP_1315-1324) and RD8a2 (SP_1325-
361 SP_1331) (S5 Fig). Genes in RD8a1 include eight *ntp* genes that code for the multiunit *V-type*
362 *proton/sodium ATP synthase* complex that produces ATP through oxidative phosphorylation in the
363 presence of a Na^+ gradient across the membrane (42). RD8a2 includes *neuraminidase*, *N-*
364 *acetylneuraminate lyase (nanA),* and *N-acetylmannosamine-6-phosphate epimerase (nanE).* These genes
365 cleave carbohydrates from the glycoproteins on the surface of epithelial cells. Other genes in RD8a2
366 encode the Sodium/solute symporter subunits that use Na^+ gradient to import the carbohydrates (43).
367 Symporter means the channel transports the solute (carbohydrates) and co-solute (Na^+) in the same
368 direction using the energy stored in an inwardly directed sodium gradient. In summary, genes in RD8a
369 operons work together to produce ATP, cleave carbohydrates from the host epithelial cells, and import
370 them into the bacterial cell. RD8a was not present in any samples assigned to serotypes 1, 5, and 12F,
371 whereas it was present in other abundant serotypes such as 6A, 6B, 16F, and 19F. The under-
372 represented pathway (sp00190) in the hyper-invasive serotypes was linked to genes in RD8a. The lack of
373 RD8a in the hyper-invasive serotypes may relate to their speed at entering the blood and CSF, where
374 they do not have access to free oxygen molecules that are required for oxidative phosphorylation (44).

375 Other significant genes absent from the hyper-invasive serotypes were from RD4 and RD7. RD4 is
376 composed of a group of sortase enzymes that catalyze the assembly of pilins into pili and the anchoring
377 of pili and other surface proteins to the cell wall (45)(46). The pilus is a hair-like structure associated
378 with bacterial adhesion and colonization (47). Due to the possible short colonization period of the
379 hyper-invasive serotypes, they may not benefit from genes in RD4 that assemble the pilus. Genes in RD7
380 have remained uncharacterized to date.

381 There were similarities between hyper-invasive serotypes and other strains in the sterile sites. As
382 observed for samples in the blood and CSF, capsule genes were significantly present in the hyper-
383 invasive serotypes, and RD10 was absent from serotypes 1 and 12F, however, this region was fully

384 conserved in serotype 5. RD10 was also conserved in 100% of serotypes 16F and 19F. The summary of

385 the gene present-absent analysis is illustrated in Fig 5.b.



386

Fig 5. The summary of the gene presence-absence analysis. (a) The number of significant genes present-absent in the hyper-invasive serotypes. The gene presence-absence analysis was applied using Scoary to compare the gene pools of the hyper-invasive serotypes and serotypes 16F and 19F. P-values were corrected by the Bonferroni method, and significant genes had an adjusted p-value of less than 0.05. (b) The significant presence and absence of RDs in samples from blood and CSF is shown as a presence-absence heatmap.

387 As mentioned, serotypes 6B and 23F were abundant in both carrier and patient groups, the test
388 between serotype 6B from the nasopharynx (n=60) and sterile sites (n=37) and serotype 23F from the
389 nasopharynx (n=43) and sterile sites (n=50) did not identify any significant gene. The gene content of
390 these two serotypes in the nasopharynx and sterile sites was similar.

391 To highlight genes that might assist pneumococci in crossing the blood-brain barrier, a test was
392 conducted between the samples from blood and CSF. The analysis between whole blood (n=368) and
393 CSF (n=284), serotype 1 samples from blood (n=60) and CSF (n=61), and serotype 5 samples from blood
394 (n=75) and CSF (n=23) did not identify any significant genes.

395 Discussion

396 The *S. pneumoniae* genome is highly diverse, and only a small fraction of genes is conserved between all
397 strains. In this species, the pan-genome is open and provides an unlimited gene repertoire for the highly
398 recombinogenic pneumococci. Changing the *S. pneumoniae* habitat could promote using different
399 combinations of genes by the organisms to diversify their genome and efficiently respond to
400 environmental stresses. In this study, the genetic diversity was identified to be very high, with only
401 10.7% of genes being core. These genes have been conserved in all samples over a long time, at least
402 from 1997 to 2015, and their presence may be vital for cell survival, making them potential targets for
403 drug design and vaccine development. In particular, core genes with no SNP in their structure would of
404 interest. Examples of the conserved core genes identified in this study included SP_1961 (*rpoB*, DNA-
405 dependent RNA polymerase), SP_0251 (formate acetyltransferase), SP_1891 (*amiA*, Oligopeptide
406 binding protein), and SP_0855 (*parC,* Topoisomerase IV). The conserved core genes are involved in DNA
407 transcription and translation.

408 Serotypes 1, 5, and 12F were significantly prevalent among patients and scarce in the carrier group. This
409 characteristic was interpreted as their high invasiveness likely with a short period of nasopharyngeal
410 colonization. On the other hand, serotypes 16F and 19F were significantly found among carriers,

411 implying that they dominate the nasopharynx with low invasiveness. Most of the other serotypes were
412 common between carriers and patients. Knowing that pneumococcal virulence strongly depends on the
413 serotype of isolates, we sought to address why several serotypes were shared across both nasopharynx
414 and sterile sites. Here, we discuss two possible scenarios that could justify the ubiquitous presence of
415 some serotypes in both nasopharynx and sterile sites.

416 The first scenario is related to the colonization of *S. pneumoniae*, which is known as a prerequisite for
417 virulence(4). Several samples from the carriage group may be actually the invasive serotypes collected
418 during their colonization phase. Abundant serotypes such as 6B and 23F that had a similar frequency
419 amongst carriers and patients need to colonize the upper respiratory tract longer than the hyper-
420 invasive serotypes before entering the sterile organs. In contrast, the hyper-invasive serotypes 1, 5, and
421 12F colonize the nasopharynx for a short period and quickly enter the sterile sites.

422 The second possible scenario relates to the differential gene expression pattern of shared genes in the
423 ubiquitous serotypes. Although the type-specific *cps* genes were identified in the isolates of both
424 nasopharynx and sterile sites, the expression pattern of these genes could vary within each strain, which
425 would contribute to the invasiveness of some strains. Several studies described a cycle of encapsulation
426 and un-encapsulation amongst pneumococcal strains. Isolates benefit from mutations in the *cps* locus to
427 either cease or re-start the capsule expression (48). The lack of a capsule at the epithelial surface
428 enables the bacterium to expose its surface proteins on the cell wall underneath the capsule and
429 promote adherence to the host epithelial cells. It has been estimated that 15% of isolates in the upper
430 respiratory tract are unencapsulated and adhere to the respiratory epithelial cells more efficiently than
431 encapsulated isolates (49)(50). Lack of capsule also facilitates acquiring virulence and resistance genes
432 from other isolates. The thick capsule prevents immunoglobulins from interacting with the pathogen
433 surface proteins during disease. Meanwhile, the negatively charged CPS interferes with the function of
434 the host phagocytes (51)(52). Taken together, the presence of the *cps* locus in the genome of isolates
435 assigned to the same serotype does not necessarily reflect the encapsulation of all cells.

436 Serotypes 1 and 5 can infect all age groups and cause severe IPDs (53). Serotype 1 is genetically distinct
437 between different geographical regions (54) and is known as the leading cause of pneumococcal
438 meningitis in Africa (55)(56). Our findings support previous research showing that serotypes 1, 5, and
439 12F are the major cause of IPDs in Malawi (57). Serotype 1 was persistently dominant in pre- and post-
440 PCV13 eras, serotype 5 was only predominant in pre-PCV3, and serotype 12F emerged after vaccination.
441 The study also characterized the high genetic distinction of the hyper-invasive serotypes in Malawi by
442 identifying significant genes that are present or absent in their genome structure compared to other
443 serotypes. Many of the significant genes present in the nasopharynx and sterile sites were homologous
444 and had the same function, and there were significant genes with an unknown function (hypothetical
445 proteins). However, of greatest interest, we did find several significant genes with specific functions that
446 could explain the difference between the biology of the hyper-invasive serotypes and nasopharyngeal
447 samples.

448 Genes from RD8a were not present in the genome of serotypes 1, 5, and 12F. The presence of this
449 region has been reported to be associated with the virulence of serotypes 6B and 14 in the United States
450 (58). We found this region conserved in serotypes 13, 14, 16F, and 19F in Malawi. One may hypothesize
451 that RD8a may be required by serotypes that colonize the nasopharynx longer than serotypes 1, 5, and
452 12F. The functions of genes in RD8a strengthen the assumption to some extent that RD8a may be
453 essential for long nasopharyngeal colonization. RD8a harbors genes such as *neuraminidase*, *nanA,* and
454 *nanE* that cleave terminal sialic acid residues from mucoglycans and epithelial glycoconjugates and assist
455 the pathogen in penetrating the mucus layer and adhering to the epithelial cells in the nasopharynx. In
456 the meantime, because free carbohydrates are scarce in the upper respiratory tract (59), cleaved sialic
457 acid can be used as the carbon source for metabolism. During colonization, secretion of the
458 pneumococcal toxins elevates the level of sodium ions (Na^+) in the nasopharynx (60), the sodium-solute
459 symporter in RD8a imports a wide variety of substrates with the sodium ions into the cell (61).

460 Most importantly, the *ntp* gene cluster in RD8a encodes the *V-type sodium ATP synthase* that pumps the
461 extra sodium ions out of the cell (42) and uses the sodium-motive force for oxidative phosphorylation
462 and ATP synthesis (62). Oxidative phosphorylation is the final step of aerobic respiration that requires
463 free oxygen molecules for ATP synthesis. Pneumococci are facultative anaerobes that can either
464 perform aerobic or anaerobic respiration with or without oxygen. In the upper respiratory tract, they
465 access atmospheric oxygen molecules that can be used by *ntp* genes to perform oxidative
466 phosphorylation. However, genes in RD8a may not be beneficial for hyper-invasive serotypes that
467 supposedly do not stay in the nasopharynx for long. Thus, the level of aerobic ATP synthesis is
468 presumably higher in serotypes 13, 14, 16F, and 19F in contrast with serotypes 1, 5, and 12F, which lack
469 RD8a.

470 Pneumococci can ferment up to 30 types of carbohydrates, imported mainly by two types of membrane
471 transporters, including ATP-binding cassette (ABC) transporters and PTS transporters (63). The major
472 differences between ABC and PTS transporters are: (i) ABC transporters use energy from ATP, but PTS
473 transporters use energy from phosphoenolpyruvate, and (ii) ABC transporters do not modify the
474 imported substrate, but PTS transporters phosphorylate the incoming sugar upon transport. Generally,
475 ABC transporters require more energy than PTS transporters, albeit they can transport longer and more
476 complicated carbohydrates (64). Unlike isolates in the nasopharynx, serotypes 1, 5, and 12F have access
477 to more simple and free host dietary carbohydrates in the blood and the central nervous system. Due to
478 a potential lower ATP synthesis level in serotypes 1, 5, and 12F (due to the lack of RD8a), they may
479 prefer to use PTS transporters to uptake sugars such as fructose and lactose, and that is why genes that
480 encode PTS transporters are significantly more present in the hyper-invasive serotypes. In addition to
481 the sugar uptake, PTS transporters regulate several pathways in bacteria, such as gene expression and
482 communication between cells. Thus, the phenotypic effects of the PTS transporters should not be
483 limited just to their ability to import carbohydrates (65).

484 Genes from RD10 were absent from serotypes 1 and 12F. However, they were conserved in serotypes 5,
485 16F, and 19F. Moreover, the location-based gene presence-absence analysis showed that RD10 was
486 significantly present in the nasopharyngeal samples compared to samples from sterile sites. These genes
487 have homology to the general secretion pathway protein B sceA2/Y2 system components in
488 *Streptococcus gordonii* that contribute to the secretion of the general secretion pathway protein B
489 involved in infective endocarditis (66). In the *S. pneumoniae* genome, the homolog of general secretion
490 pathway protein B is the PsrP that is transported to the bacterial cell surface by the SecA2/Y2 system
491 encoded by genes in RD10. Studies conducted on *Streptococcus gordonii* showed that the presence of
492 SecA2/Y2 promotes the adhesion to epithelial cells in the nasopharynx and erythrocytes in the blood
493 (67)(68), which is why SecA/Y2 is significantly present in nasopharyngeal samples and serotype 5
494 (abundant in the blood). The presence of the secA2/Y2-like component should also facilitate the export
495 of pneumolysin that enhances adhesion to the host cell and survival in the blood (69)(70).

496 In conclusion, some of the genes specifically present (absent) in the hyper-invasive serotypes (1, 5, and
497 12F) may contribute to the invasiveness and low colonization rate. However, experimental work is
498 required to confirm the outcome of computational work. The serotype is the best driver of the
499 pneumococcal population structure, however, the study characterized the genetic divergence of
500 serotypes 1, 5, and 12F to be much greater than other serotypes. Their significant presence in the blood
501 and CSF accounted for the most pronounced genomic and functional differences observed between the
502 nasopharynx and sterile sites. The low frequency of serotypes 1, 5, and 12F among carriers may be due
503 to their short-term colonization before entering the sterile sites. These invasive serotypes harbor the
504 elements of PTS transporters while they lack genes from RD8a. In serotype 5, RD10 is highly conserved,
505 but this region was absent from serotypes 1 and 12F. The study demonstrated that isolation sites do not
506 significantly influence the genomic structure of pneumococcal isolates. Although a few genes were
507 associated with the virulence of commonly present serotypes in both nasopharynx and sterile sites, we
508 speculated that the differences between them could be captured by other high throughput techniques

509 such as gene expression analysis. In summary, the results shed light on the pneumococcal population
510 structure and serotypes in Malawi. The unique cluster of significant genes in the hyper-invasive
511 serotypes and highly conserved core genes could be potential therapeutic targets.

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527

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715 # Supporting information

716 **S1 Fig. Characteristics of the 1477 pneumococcal isolates used in the study.** (a) The relative frequency
717 of serotypes in the entire cohort, samples were assigned to 56 serotypes. For each sample, the in-silico
718 serotyping was accomplished by SeroBA. (b) Frequency of isolates in the pre- and post-PCV13 eras in
719 Malawi. (c) Frequency of isolates obtained from each specimen source.

720 **S2 Fig. The pan-genome of 1477 pneumococcal samples isolated in Malawi was obtained from 1997 to**
721 **2015.** The pan-genome is an open pan-genome, which means the number of total genes increases
722 unlimitedly when the sample size grows. The dashed line represents the number of total genes, and the
723 solid line represents the number of conserved genes in the pan-genome.

724 **S3 Fig. The three-dimensional PCA of the gene distribution in the vaccine types.** For each serotype and
725 for downsampling, 10 samples were randomly selected from the nasopharynx, blood, and CSF. The PCA
726 was conducted using the R package MixOmics. Hyper-invasive serotypes 1, 5, and 12F clustered
727 separately from other strains.

728 **S4 Fig. Genes in RD10 are absent from serotypes 1 and 12F but conserved in serotype 5, 16F, and 19F.**
729 Genes from RD10 encode the components of the secretory system SecA2/Y2 that transports
730 glycoproteins to the bacterial cell surface, which are required for binding to the human proteins on the
731 surface of epithelial cells and erythrocytes.

732 **S5 Fig. RD8a consists of two operons RD8a1 (SP_1315-1324) and RD8a2 (SP_1325-1331).** This region is
733 not detected in the significant invasive serotypes 1, 5, and 12F, but it is present in more than 80% of
734 serotype 16F and 19F that significantly dominates the nasopharynx. The important biological processes
735 carried out by these genes are the transport of ions across the membrane and the synthesis of ATP
736 molecules.

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740 **S1 Table. Statistical analysis of serotypes' prevalence across specimen sources.**

741 **S2 Table. Gene presence-absence analysis (Invasive vs Nasopharyngeal, serotypes 1, 5, and 12F were**

742 **excluded).**

743 **S3 Table. Gene presence-absence analysis (Serotype 1 vs 16F & 19F).**

744 **S4 Table. Gene presence-absence analysis (Serotype 5 vs 16F & 19F).**

745 **S5 Table. Gene presence-absence analysis (Serotype 12F vs 16F & 19F).**

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759 # Figure captions

760 **Fig 1. The distribution of the 56 pneumococcal serotypes assigned to 1477 samples from Malawi.** (a)
761 The relative frequency of each serotype in the nasopharynx of carriers, the blood of bacteremia
762 patients, and the CSF of meningitis patients is shown in blue, red, and yellow, respectively (UT: Un-
763 Typeable). (b) The log-transformed odds ratio of the significantly over- and under-abundant serotypes in
764 the sterile sites (blood and CSF). Fisher's exact test was applied to identify serotypes with a significant
765 differential abundance among carriers and patients (nasopharynx and sterile sites) at the significance
766 level of the Benjamini-Hochberg adjusted p-value < 0.01 (BH: Benjamini-Hochberg).

767 **Fig 2. The pan-genome matrix of 1477 pneumococcal isolates from Malawi.** The pan-genome is
768 visualized as a gene presence-absence heatmap that represents the hierarchical unsupervised clustering
769 of samples based on the distribution of genes in the pan-genome. Each row is a sample, and each
770 column is a gene. A blue dot denotes the presence of each gene. On the right side of the heatmap, the
771 large blue block shows core genes present in all samples. The left side of the heatmap represents the
772 accessory genome along with the clustering bands. In addition to the significant serotypes 1, 5, 12F, 16F,
773 and 19F, other abundant serotypes, including 6A, 6B, and 23F, as well as serotypes with source-based p-
774 value < 0.05, including 21, 11A, and 15B, are also highlighted on the heatmap.

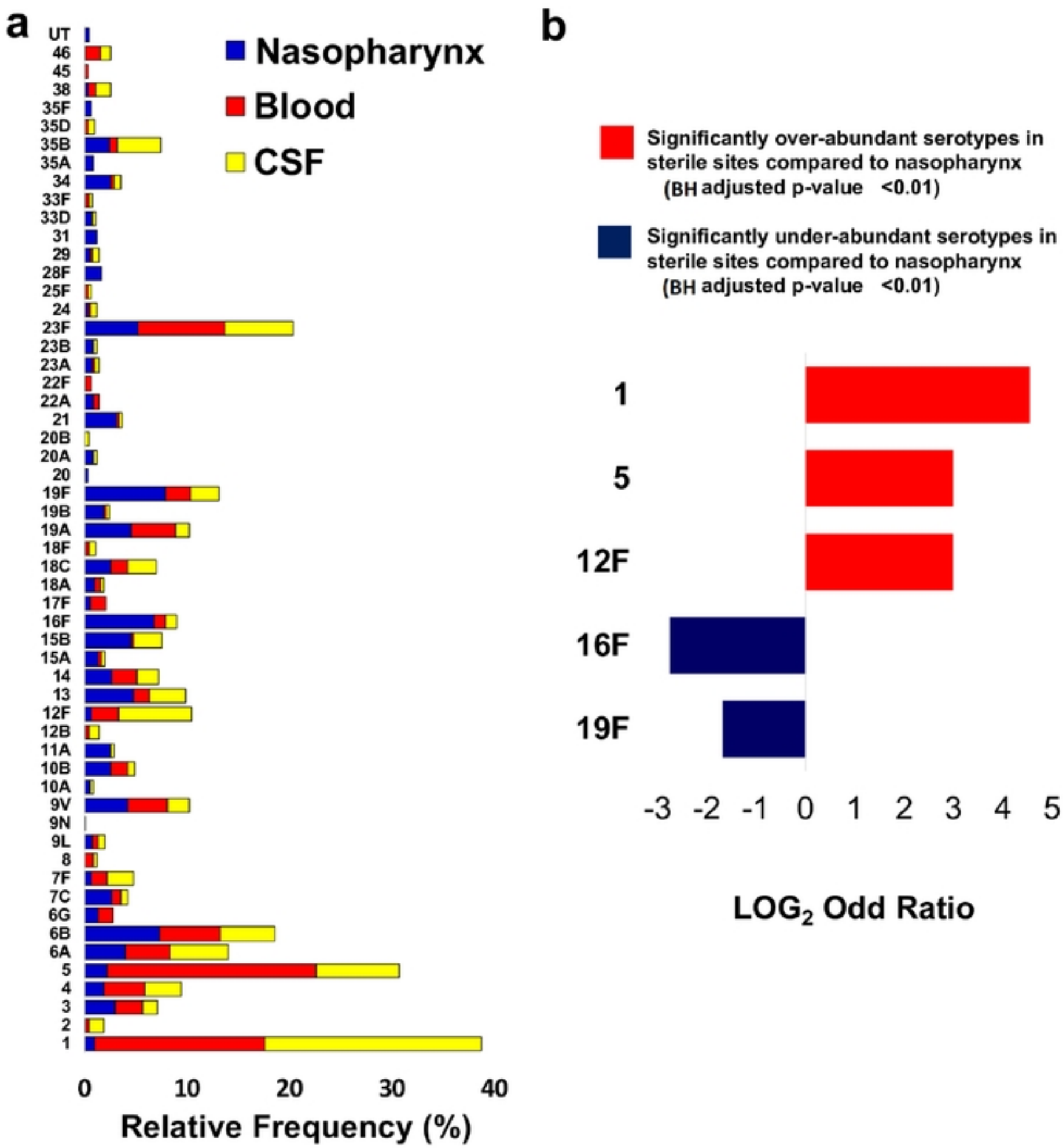
775

776 **Fig 3. The phylogenetic population structure of 1477 pneumococcal samples from Malawi.** The
777 phylogenetic tree was built based on the multiple sequence alignment of the core genome using the
778 maximum likelihood method. Colors on the loops show the serotypes, specimen sources (isolation sites),
779 and PCV13 eras. In addition to the significant serotypes 1, 5, 12F, 16F, and 19F, other abundant
780 serotypes, including 6A, 6B, and 23F, as well as serotypes with source-based p-value < 0.05, including 21,
781 11A, and 15B, are also highlighted on the tree.

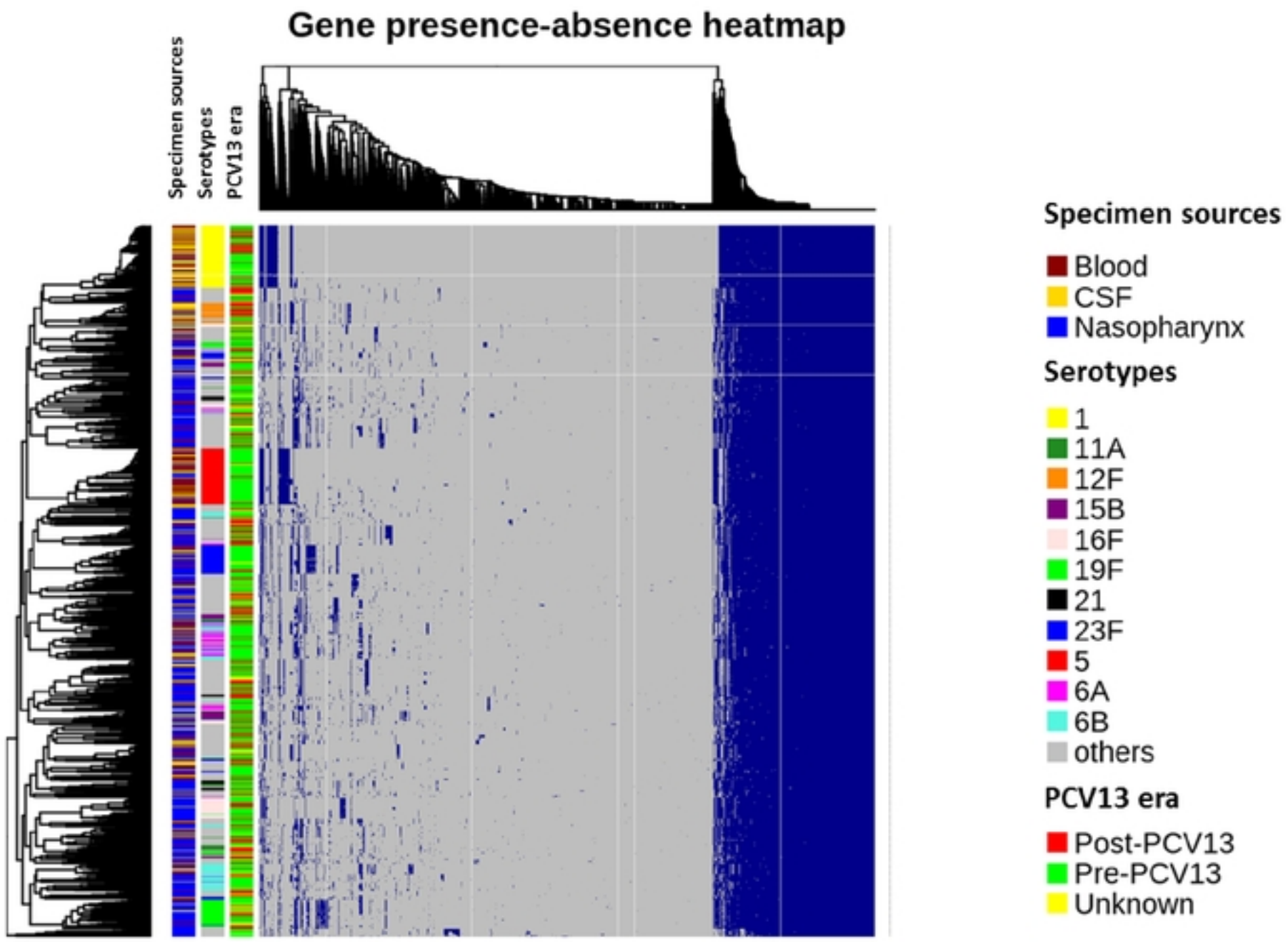
782 **Fig 4. The PCA of the gene distribution in the pan-genome of pneumococcal isolates from 1477**
783 **Malawians.** The PCA of variants (gene presence-absence) in the accessory-genome indicates the
784 influence of (a) specimen sources (isolation sites), (b) serotypes, (c) PCV13 (vaccination) era, and (d)
785 geographical locations on the gene presence-absence profile of pneumococcal isolates in Malawi.
786 Serotypes 1 and 5 were clearly separated from other samples.

787 **Fig 5. The summary of the gene presence-absence analysis. (a) The number of significant genes**
788 **present-absent in the hyper-invasive serotypes.** The gene presence-absence analysis was applied using
789 Scoary to compare the gene pools of the hyper-invasive serotypes and serotypes 16F and 19F. P-values
790 were corrected by the Bonferroni method, and significant genes had an adjusted p-value of less than
791 0.05. (b) The significant presence and absence of RDs in samples from blood and CSF is shown as a
792 presence-absence heatmap.

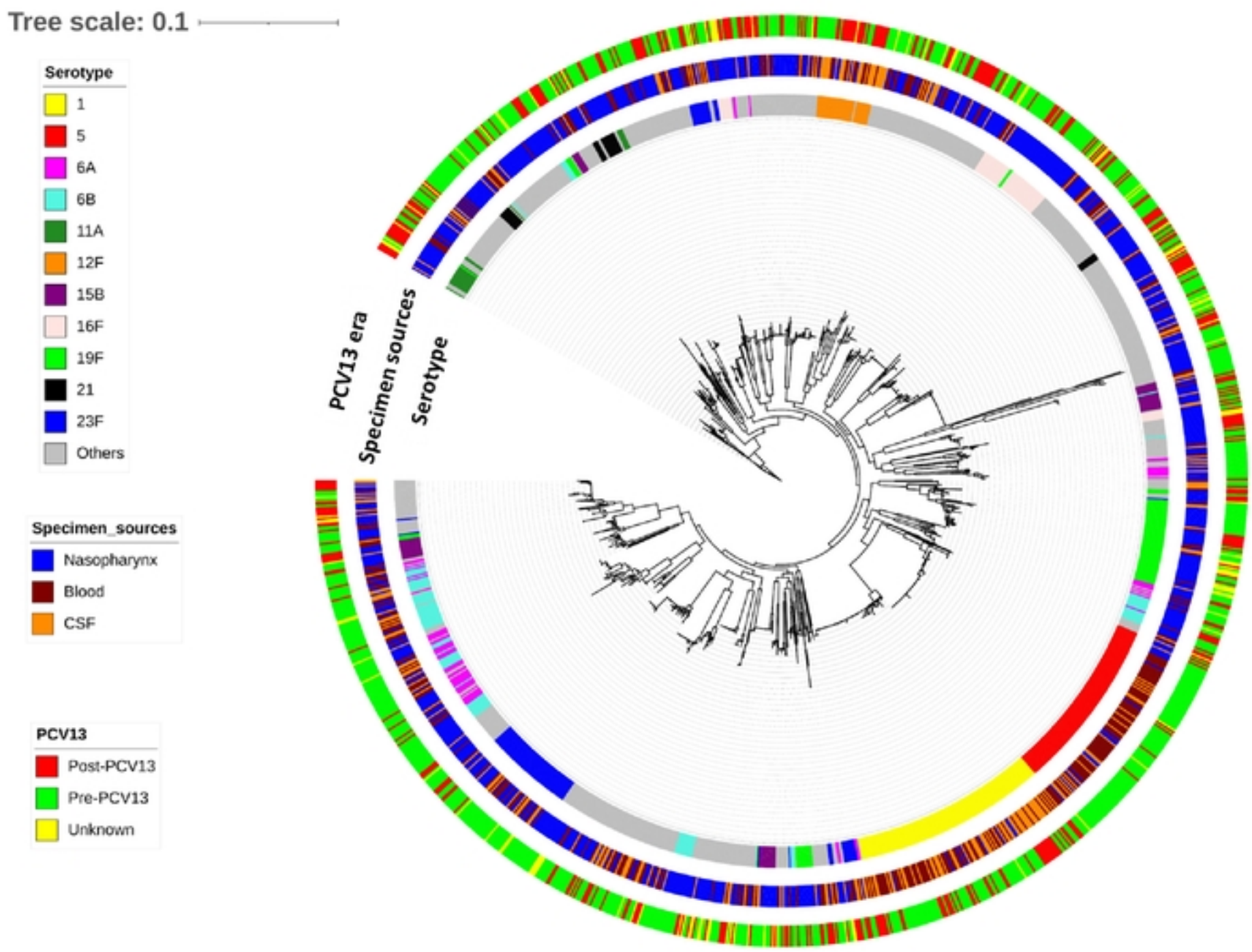
793


a
Nasopharynx
Blood
CSF
UT
46
45
38
35F
35D
35B
35A
34
33F
33D
31
29
28F
25F
24
23F
23B
23A
22F
22A
21
20B
20A
20
19F
19B
19A
18F
18C
18A
17F
16F
15B
15A
14
13
12F
12B
11A
10B
10A
9V
9N
9L
8
7F
7C
6G
6B
6A
5
4
3
2
1
0
10
20
30
40
Relative Frequency (%)
b
Significantly over-abundant serotypes in sterile sites compared to nasopharynx (BH adjusted p-value <0.01)
Significantly under-abundant serotypes in sterile sites compared to nasopharynx (BH adjusted p-value <0.01)
1
5
12F
16F
19F
-3
-2
-1
0
1
2
3
4
5
LOG2 Odd Ratio

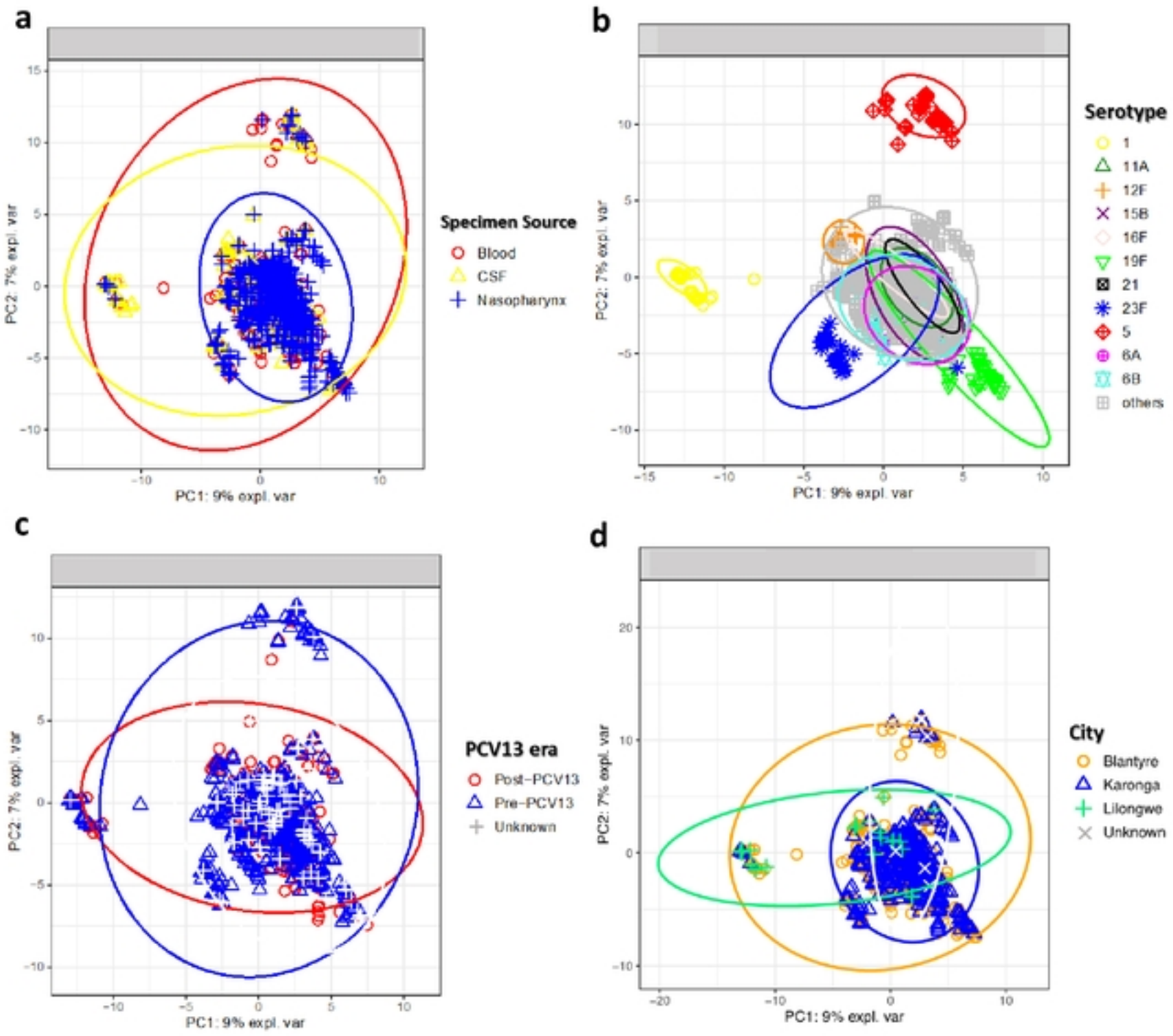
Figure


Gene presence-absence heatmap
Specimen sources
Serotypes
PCV13 era
Specimen sources
Blood
CSF
Nasopharynx
Serotypes
1
11A
12F
15B
16F
19F
21
23F
5
6A
6B
others
PCV13 era
Post-PCV13
Pre-PCV13
Unknown

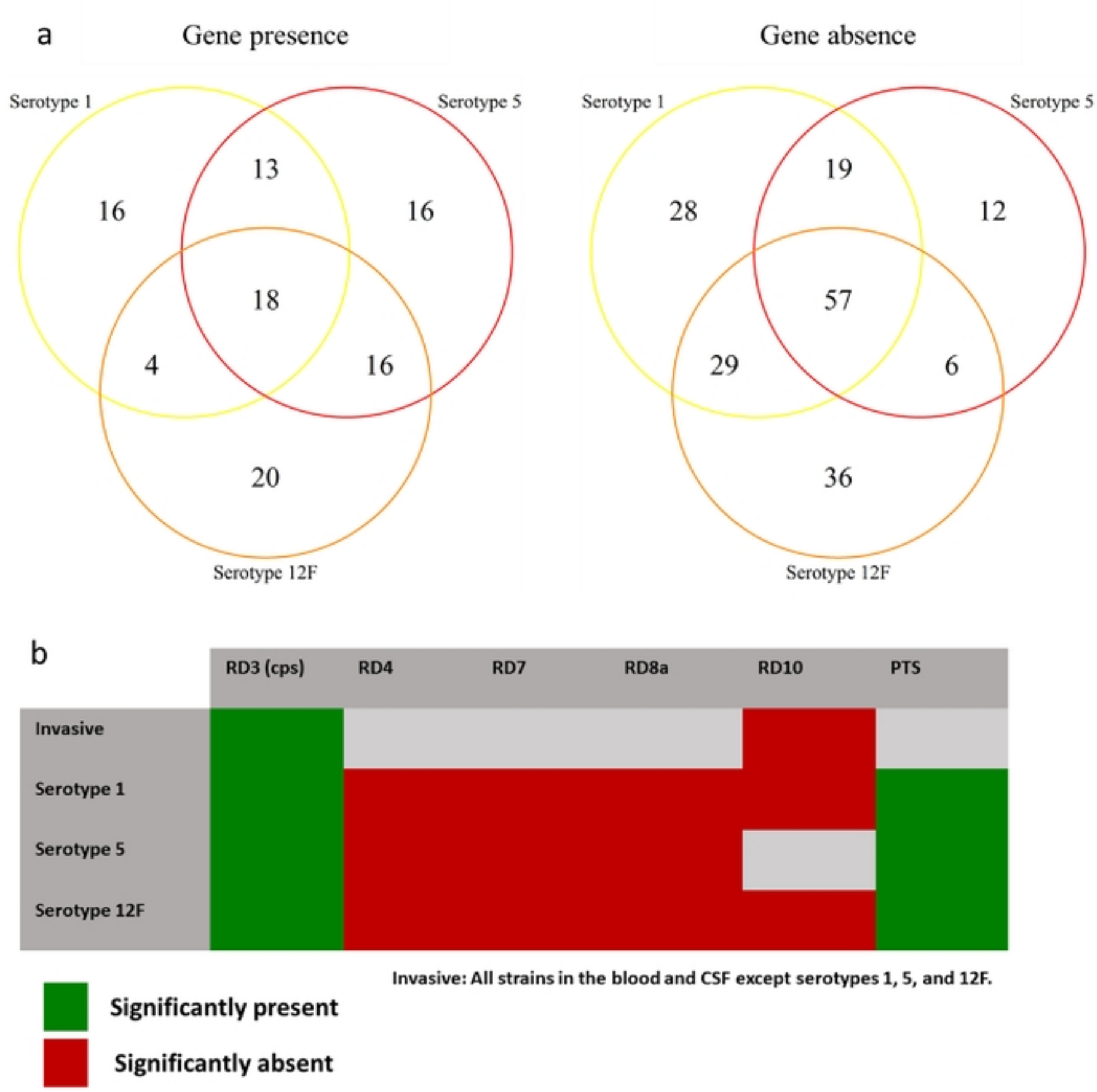
Figure


Tree scale: 0.1
Serotype
1
5
6A
6B
11A
12F
15B
16F
19F
21
23F
Others
Specimen_sources
Nasopharynx
Blood
CSF
PCV13
Post-PCV13
Pre-PCV13
Unknown
PCV13 era
Specimen sources
Serotype

Figure


a
b
c
d
PC1: 9% expl. var
PC2: 7% expl. var
Specimen Source
Blood
CSF
Nasopharynx
Serotype
1
11A
12F
15B
16F
19F
21
23F
5
6A
6B
others
PCV13 era
Post-PCV13
Pre-PCV13
Unknown
City
Blantyre
Karonga
Lilongwe
Unknown

Figure


a
Gene presence
Serotype 1
Serotype 5
Serotype 12F
16
13
16
18
4
16
20
Gene absence
Serotype 1
Serotype 5
Serotype 12F
28
19
12
57
29
6
36
b
RD3 (cps)
RD4
RD7
RD8a
RD10
PTS
Invasive
Serotype 1
Serotype 5
Serotype 12F
Invasive: All strains in the blood and CSF except serotypes 1, 5, and 12F.
Significantly present
Significantly absent

Figure