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Real-time metagenomics-based diagnosis of community-acquired meningitis: A prospective series, southern France

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Summary

Background Point-Of-Care (POC) diagnosis of life-threatening community-acquired meningitis currently relies on multiplexed RT-PCR assays, that lack genotyping and antibiotic susceptibility profiling. We assessed the usefulness of real-time metagenomics (RTM) directly applied to the cerebrospinal fluid (CSF) for the identification, typing and susceptibility profiling of pathogens responsible for community-acquired meningitis.

Methods A series of 52 CSF samples from patients suspected of having community-acquired meningitis, were investigated at POC by direct RTM in parallel to routine real-time multiplex PCR (RT-PCR) and bacterial culture, for the detection of pathogens. RTM-generated sequences were blasted in real-time against an in-house database incorporating the panel of 12 most prevalent pathogens and against NCBI using EPI2ME online software, for pathogen identification. *In-silico* antibiogram and genotype prediction were determined using the ResFinder bio-tool and MLST online software.

Findings Over eight months, routine multiplex RT-PCR yielded 49/52 positive CSFs, including 21 *Streptococcus pneumoniae*, nine *Neisseria meningitidis*, eight *Haemophilus influenzae*, three *Streptococcus agalactiae*, three *Herpesvirus-1*, two *Listeria monocytogenes*, and one each of *Escherichia coli*, *Staphylococcus aureus* and *Varicella-Zoster Virus*. Parallel RTM agreed with the results of 47/52 CSFs and revealed two discordant multiplex RT-PCR false positives, one *H. influenzae* and one *S. pneumoniae*. Both multiplex RT-PCR and RTM agreed on the negativity of three CSFs. While multiplex RT-PCR routinely took 90 min, RTM took 120 min, although the pipeline analysis detected the pathogen genome after 20 min of sequencing in 33 CSF samples; and after two hours in 14 additional CSFs; yielding > 50% genome coverage in 19 CSFs. RTM identified 14 pathogen genotypes, including a majority of *H. influenzae* b, *N. meningitidis* B and *S. pneumoniae* 11A and 3A. In all 16 susceptible cultured bacteria, the *in-silico* antibiogram agreed with the *in-vitro* antibiogram in 10 cases, available within 48 h in routine bacteriology.

Interpretation In addition to pathogen detection, RTM applied to CSF samples offered supplementary information on bacterial profiling and genotyping. These data provide the proof-of-concept that RTM could be implemented in a POC laboratory for one-shot diagnostic and genomic surveillance of pathogens responsible for life-threatening meningitis.

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Keywords: Community-acquired meningitis; Real-time metagenomics; Point-of-care (POC) laboratory diagnosis; Antibiotic resistance; Genotyping

Research in context

Evidence before this study

Community-acquired bacterial meningitis is a life-threatening infection that can progress to mortality within 48 h. The emergency diagnosis of infectious meningitis is currently based on multiplex real-time amplification using a syndromic panel limited by the most frequent microorganisms which lead to a central nervous system prognosis. Bacterial characterisation and drug resistance profiling require additional *in-vitro* investigations which take over 48 h, delaying pathogen-targeted treatment. Genomic surveillance and antibiotic resistance testing are based only on bacteria isolated from cerebrospinal fluid, failing in 60% of cases. We searched PubMed up to 30 November 2021 for research articles published in English, using the following search terms “real-time metagenomics sequencing”, “meningitis”, and “direct diagnosis”. Several articles had been published testing the Oxford Nanopore technologies sequencing on CSF, but no investigations were found into the direct diagnosis of CSF series. When the three terms were used together, only two articles previously published by us were found. As previously reported, we implemented real-time metagenomic sequencing (RTM) at the POC laboratory for the diagnosis of life-threatening infectious meningitis, in addition to the BioFire FilmArray® investigation. In light of its simplicity, rapidity and additional information collected, we propose RTM as a powerful diagnostic tool for the investigation of prospective series of CSF samples collected from patients with meningitis.

Added value of this study

In this study, we diagnosed a series of community-acquired meningitis cases by RTM directly from CSF samples. Over eight months, 52 CSFs were investigated directly by RTM using a four-hour workflow. Thirty-three CSF samples (63.5%) were diagnosed as positive after a 20-min sequencing run and an additional 14 were diagnosed as positive after two hours. The pipeline analysis of antibiotic resistance and bacteria genotyping was provided *in-silico* at the same time as sequencing, in contrast to conventional diagnostics. In addition, uncultured bacteria were successfully profiled *in-silico*, basing on pathogen genome analysis, independently of genome coverage.

Implications of all the available evidence

Despite the limited sample size in this study, using a four-hours workflow, RTM proved successful in diagnosing, genotyping, and profiling bacteria directly from CSF

samples. At two discordances with conventional multiplex RT-PCR, RTM is a suitable method for the diagnosis of life-threatening meningitis at the POC laboratory.

Introduction

The rapid diagnosis of life-threatening, community-acquired meningitis (CAM) remains challenging in point-of-care (POC) laboratories.¹ Bacterial meningitis charged with a 24-h mortality of 8–15%,² results in an estimated 290,000 deaths every year, causing more than 50% deaths annually from all meningitis causes, and leaving one in five people who recover with chronic handicap.³ Community-acquired bacterial meningitis around the world is mainly due to *Streptococcus pneumoniae* (*S. pneumoniae*), *Neisseria meningitidis* (*N. meningitidis*), *Haemophilus influenzae* (*H. influenzae*) and *Streptococcus agalactiae* (*S. agalactiae*).⁴ In Europe, *S. pneumoniae* and *N. meningitidis* are the most common causes of bacterial meningitis,² usually affecting children ≤ 5 years in 22.5% and 47% of cases, respectively.² Current POC diagnosis of bacterial meningitis is based on real-time multiplex PCR (RT-PCR) assays incorporating a syndromic meningitis and encephalitis panel,^{5–10} targeting small specific pieces of the pathogen genome.^{5,6} These approaches overlook serotype/genotype diversity, a major limitation for the microbiological diagnosis of bacterial meningitis, do not provide sufficient information for pathogen genotyping, and require bacterial culture to characterise different serotypes and antimicrobial resistance.^{5,6} Accordingly, additional specific PCRs have to be performed for genotyping *N. meningitidis* B and C serotypes,^{11–13} and *H. influenzae* b serotype associated to invasive diseases.^{2,14}

Real-time metagenomics sequencing (RTM) could, theoretically, overcome this limitation, identifying the causative agent of meningitis,¹⁵ as well as its genotype/serotype directly from the cerebrospinal fluid (CSF) based on pathogen genome sequence.^{14,16} Indeed, we and others have already published evidence that RTM could be implemented in a POC laboratory, for one-shot diagnostic, genotyping as well as *in-silico* antibiotic resistance prediction, which is competitive in time and cost with commercial multiplex RT-PCR.^{14–18}

In this study, we prospectively diagnosed a series of cases of community-acquired meningitis, directly using RTM on left-over CSF samples in a POC laboratory.

Methods

Ethics

As per French legislation, no specific patient consent was required. The analysis of biological samples obtained in the medical care context was considered as non-interventional research (article L1221-1.1 of the French Public Health Code), requiring only the non-opposition of the patient during sampling (article L1211-2 of the French Public Health Code). All data were generated as part of routine laboratory work at the Assistance Publique-Hôpitaux de Marseille and Nîmes university hospital, in the context of the routine clinical management of patients suspected of having community-acquired meningitis. No specific clinical sampling was performed for this study and RTM was applied to anonymised left-over CSF samples for which the age and sex of patient were anonymously collected, following a standard routine laboratory protocol including multiplex real-time PCR, which was carried out in full respect of the French law regarding clinical research. Accordingly, this study was approved by IHU Méditerranée Infection Ethics Committees under number: 2021-004 before the study began in Marseille; and further approval was granted by “Interface Recherche Bioéthique Institutional Review Board” Ethics Committee of Nîmes CHU under the following number: 21.0016 before the study began in Nîmes.

Routine microbial diagnosis

In the POC laboratory, all CSF samples were routinely examined to count white and red blood cells directly using NucleoView NC-3000 equipment and NucleoView/ChemoMetec software (ChemoMetec NucleoCounter, Allerød, Denmark). In parallel, 200 µL samples of CSF were used for multiplex RT-PCR diagnosis (BioFire FilmArray®, bioMérieux, Marcy-l'Étoile, France) as previously described.⁵ Further in the core laboratory, 200 µL of CSF was inoculated on chocolate agar PolyViteX (bioMérieux) and Columbia agar enriched with a 5% sheep's blood (bioMérieux) medium incubated at 37 °C under 5% CO₂ for five days, and on Columbia agar enriched with 5% sheep blood (bioMérieux) under anaerobic conditions for ten days at 37 °C for bacterial culture and *in-vitro* antibiogram in the standard bacteriology laboratories. For any isolate, antibiograms were validated according to the antibiotic panel approved by the French Antibiogram Comity of Microbiology Society (CA-SFM, version V1 2020) (Appendix 1).

Installation of the RTM platform in the POC

We updated the equipment available in our POC laboratories by setting up MinION sequencers (Oxford Nanopore Technologies, Oxford, UK) (Figure 1, Appendix 2). As an example, in the Marseille POC laboratory, an

RTM bench was set up on a surface area of 210 × 70 cm², the atmosphere was stabilised at one atm and the temperature was controlled by a central air conditioning at 20 °C ± 2 °C. The RTM bench was equipped with a Biocap® hood (Erlab, Val-de-Reuil, France), a clean area for DNA preparation, Qubit® for DNA quantification, a thermal cycler (ThermoFisher, Illkirch, France), an incubator at 20 °C with agitation for the different incubation steps, a vortex for mixing reagents and buffers, magnetic rack, a tube ice rack for enzyme storage during the manipulation, a mini centrifuge at 12,000 g, micro-pipettes with different volumes and a biological waste container (Figure 1, Appendix 2). Metagenomic handling was performed in an 1800 cm² workspace. For library sequencing, four MinION instruments were attached in parallel to a powerful computer equipped with minimum Windows 10 or a Linux version (16.04 LTS) operating system, an i7 processor, RAM ≥ 8Gb, USB 3 port, with and enough disk space to store the data (~ 1 Tb) (Lenovo, China), and an internet connection. MinION-Sequencer reading, and data storage were performed using Minknow Oxford Nanopore software version (8.3.1). In addition, Oxford Nanopore EPI2ME software was installed for real-time data analysis.

RTM procedure

Total DNA was extracted from 200 µL of left-over CSF samples using an EZ1 DNA Tissue Kit (Qiagen, Courtaboeuf, France), after 15 min of incubation at 56 °C with 20 µL proteinase K (Qiagen), then eluted in a 50-µL volume. For the real-time next-generation sequencing, the Oxford Nanopore library preparation was performed in a 75-µL final volume as previously described.¹⁴ Briefly, 48 µL DNA was prepared and end-repaired in 60 µL containing 3.5 µL of NEBNext FFPE DNA Repair buffer and 3.5 µL of Ultra II End-prep reaction buffer, 2 µL of NEBNext FFPE DNA Repair mix (New England Biolabs, Evry-Courcouronnes, France) and 3 µL of Ultra II End-prep enzyme mix (New England BioLabs). The repair reaction Master Mix was incubated for five minutes at 20 °C followed by a five-minute incubation at 65 °C on a GeneAmp PCR System Thermal Cycler (Applied Biosystems, Foster City, CA, USA). Repaired DNA was purified using equal volumes of Agencourt Ampure XP beads (Beckman Coulter, Villepinte, France), and eluted in 25 µL of sterile water after incubation for five minutes at room temperature and two washes with 70% ethanol. A barcoding step was added to the standard Oxford Nanopore protocol to avoid any cross-contamination and to reduce the cost of the tests. A 22.5 µL volume of repaired DNA was barcoded in 50 µL containing 2.5 µL of native barcoding and 25 µL of Blunt/TA Ligase Master Mix (BioLabs), incubated for ten minutes at room temperature. Barcoded DNA was purified using 50 µL of Agencourt Ampure XP and

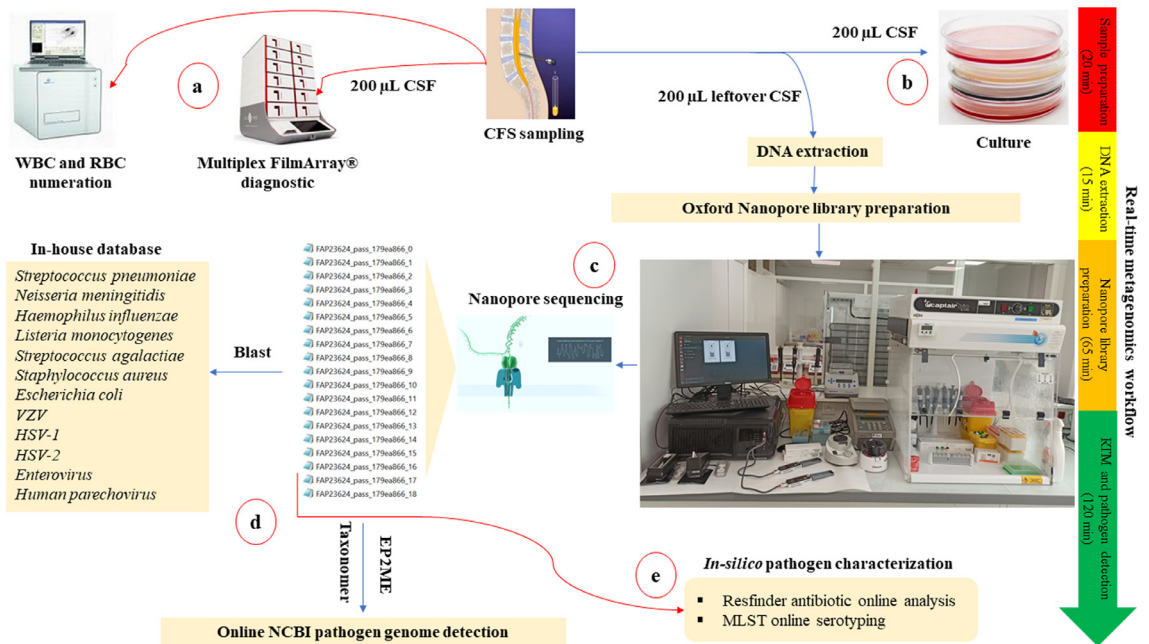


Figure 1. CSF workflow for the diagnosis of community-acquired meningitis in the POC laboratory for the 52 CSF series prospective investigation. When the CSF sample was received at the POC laboratory, several tests were performed to detect the meningitis causative agent. a) Systematically, the emergency multiplex BioFire FilmArray[®] assay performed using 200 µL CSF when the sample was received, followed by quantification of the blood cells. b) As per routine diagnosis, all CSF samples received at the POC were routinely cultured. c) The RTM diagnosis was performed in a total workflow that did not exceed four hours. Sample preparation and DNA extraction from left over CSF samples took 35 min, Oxford Nanopore library preparation took 65 min, and the MinION library sequencing took two hours. d) Real-time genome identification was performed directly by blast of the MinION generated data against an in-house database, then against the NCBI database using EPI2ME and Taxonomer online software. e) Antibiotic resistance and pathogen genotyping were *in-silico* predicted using the ResFinder online database (<https://cge.cbs.dtu.dk/services/ResFinder-4.1/>), and MLST online database (<https://pubmlst.org/organisms>).

Abbreviations: CSF: cerebrospinal fluid. WBC: white blood cell. RBC: red blood cell. RTM: real-time metagenomics. MLST: Multi-Locus Sequence Typing.

eluted in 65 µL sterile water, after incubation for five minutes at room temperature and two washes with 70% ethanol. A 65-µL volume of the barcoded DNA was indexed in 100 µL containing 20 µL NEBNext Quick Ligation Reaction Buffer (5X) buffer, 5 µL of Adapter Mix II (AMII) and 10 µL of T4 DNA Ligase and incubated for ten minutes at room temperature. 60 µL of Agencourt Ampure XP beads were then added to the ligation master mix and incubated for five minutes at room temperature. Two washes were performed using an LFB buffer, then eluted in a 15-µL volume and incubated for ten minutes at room temperature. Finally, 12 µL of the eluted library were added to 37.5 µL sequencing buffer and 25.5 µL loading beads and sequenced for up to two hours on a MinION sequencer (Oxford Nanopore, Oxford Science Park, UK) (Figure 1). For rapid pathogen genome identification, the output fastq_pass was generated every 1500 reads per file to favour real time analysis, using Minknow specific parameters before starting sequencing run.

In-silico data analysis

Pathogen genome identification. Pathogen genome sequences were detected in real-time using an in-house pipeline. First, total MinION data were aligned against an in-house database constructed in reference to Biofire FilmArray[®] panel (Appendix 6), including complete genome sequences for each one of *N. meningitidis*, *S. pneumoniae*, *H. influenzae*, *S. agalactiae*, *Escherichia coli*, *Listeria monocytogenes* (100 sequences per pathogen), VZV (148 sequences), HSV-1 (51 sequences), HSV-2 (33 sequences), Parechovirus (200 sequences) and Enterovirus (300 sequences) using a blast nucleotide command line with default parameters on the IHU server. This analysis was interpreted as positive when the number of sequences for any specific pathogen was ≥ 2. Further, detection was queried against NCBI GenBank database to increase probability pathogen genome detection using Taxonomer (<https://www.taxonomer.com>) and Oxford Nanopore EPI2ME

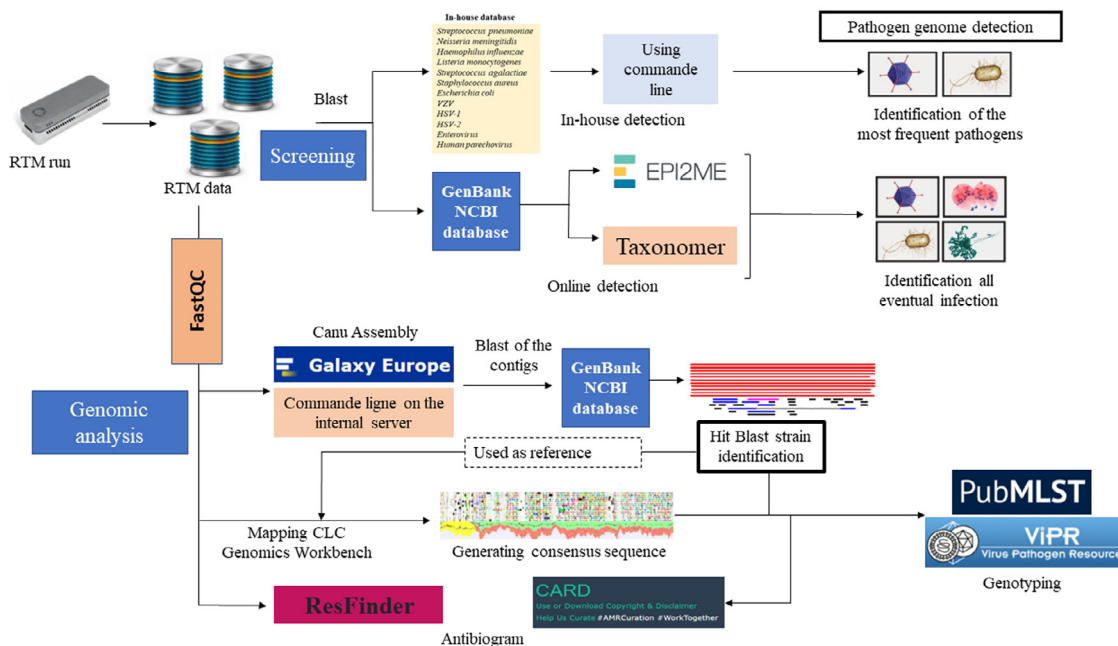


Figure 2. Bioinformatic pipeline. Rapid pathogen genome identification performed by direct alignment of MinION reads with an in-house database using blastn command line on the IHU server, further against NCBI GenBank by Taxonomer and EPI2ME online software. The analysis was interpreted as positive when ≥ 2 pathogen-specific reads were identified. To confirm pathogen identification, the quality of MinION reads was controlled by FastQC before assembly by “Canu assembler” on Galaxy Europe online software (<https://usegalaxy.eu/>). Hit-blast strains were identified by Blastn of the generated contigs against GenBank database, then used as reference genome for mapping the total MinION reads by CLC Genomics Workbench software version 21.0.3 (Qiagen). The consensus genomes were extracted in fasta files for pathogens genotyping on MLST on PubMLST (<https://pubmlst.org/organisms>) for bacteria genotyping and on ViPR online database (<https://www.viprbrc.org/>) for virus genotyping. The *in-silico* antibiogram was predicted on ResFinder (<https://cge.cbs.dtu.dk/services/ResFinder/>) and Resistance Gene Identification (<https://card.mcmaster.ca/analyze/rgi>) platforms using the total MinION reads and the generated fasta sequences with default settings.

**In-silico* antibiogram and genotype were derived from genome sequence only in the case of $> 1\%$ genome coverage.

online software (Figures 1, 2). To confirm real-time identification, reads quality control was performed using FastQC online platform and assembled by “Canu assembler” tool (Version 2.1.1) on Galaxy Europe online software (<https://usegalaxy.eu/>) and generated contigs were then aligned by Blastn against NCBI GenBank database. The identified hit-blast strain, defined by maximum sequence similarity, was used as a reference sequence for mapping of the total MinION reads using CLC Genomics Workbench software version 21.0.3 (Qiagen) with default parameters. Consensus sequences were extracted in fasta files for further analysis (Figure 2). The *in-silico* prediction of antibiotic resistance-encoding genes was carried out using ResFinder (<https://cge.cbs.dtu.dk/services/ResFinder/>) and Resistance Gene Identification (<https://card.mcmaster.ca/analyze/rgi>) online software using total pathogen reads and hit-blast strains identified by Blastn against NCBI GenBank database after assembly of the MinION reads, in reference to antibiotics routinely assayed *in-vitro* (Appendix 1). *In-silico* prediction of antibiotic resistance was performed only in the case genome coverage was $\geq 1\%$.

Pathogen genotyping. Microbial genotypes were predicted using multi-locus sequence typing tools (MLST) on total specific sequences data and hit-blast stains identified by blast against NCBI GenBank database. Virus genotyping was performed directly on ViPR online database (<https://www.viprbrc.org/>) (Figure 2). Genotyping was *in-silico* predicted only in the case genome coverage was $\geq 1\%$.

Graphic representation of data. Graphical representation was performed using R software (version 4.0.3). Pie and donut charts were created using the PieDonut function in the webr package (<https://www.R-project.org/> and <https://CRAN.R-project.org/package=webr>).

Cost analysis. We compared estimations of global cost for Biofire FilmArray[®]-based diagnosis of CAM at the POC laboratory with that of RTM-based diagnosis. Noteworthy, these estimations incorporated mean cost by

sample for RTM (144€) and FilmArray® (114€), based on values calculated with reference prices for materials and reagents, in our laboratories. We estimated that 30 RTM assays costed 4320 € including pathogen identification, genotyping, and *in-silico* antibiogram; while we estimated that 30 FilmArray® assays costed 3411€ for pathogen identification only; eventually increased by cost of additional pathogen-specific PCRs, in case of negative FilmArray® assay; and that of additional *in-vitro* investigations for pathogen characterization. Moreover, relative cost obviously decreased in case of series of CSF samples tested at the same time: as an example, RTM cost decreased to less than 70 € / CSF sample for series of 12 samples tested with the same MinION flow-cell, while FilmArray® cost was independent of the number of tested CSF samples, being series-insensitive (Table 3, Appendix 4).

Role of funding source. The funders did not have any role in the study design, data collection, data analyses, interpretation, or writing of the report.

Results

General data

CSF samples collected from 52 patients prospectively investigated in this study included 24 CSF samples at the IHU Méditerranée Infection POC laboratory in Marseille and 28 CSF samples at the bacteriology and hygiene laboratory at Nîmes University Hospital. These 52 CSF samples were collected from 24 female patients and 28 male patients, aged between 0 and 90 years (median, 38 years old), investigated between December 2020 and July 2021 (Table 1, Appendix 3).

Routine investigations

Routine FilmArray® assays detected a microorganism in 49 CSF samples. Bacterial pathogens found in 45 CSF samples included 21 *S. pneumoniae*, nine *N. meningitidis*, eight *H. influenzae*, three *S. agalactiae*, two *L. monocytogenes* and one each of *E. coli* and *S. aureus*. In addition, three *Herpesvirus-1* and one *Varicella-zoster virus* (VZV) were detected. All viral cases resulted from PCR, while 20 bacterial meningitis cases were confirmed by culture and RT-PCR, 12 by RT-PCR only, including one case (sample 31) of *S. pneumoniae* > 35 Ct which was interpreted as negative in routine POC diagnostic, 12 cases with FilmArray® only, and PCR and culture failed to identify one case of *H. influenzae* (sample 20). The 20 culture positive CSFs grew 12 *S. pneumoniae*, three *N. meningitidis*, two *H. influenzae*, two *L. monocytogenes* and one *S. aureus*, while the other 26 CSFs detected positive for bacteria by RT-PCR were culture-negative. *In-vitro* antibiotic investigation yielded 16 susceptible bacteria (nine *S. pneumoniae*, three *N.*

meningitidis, two *H. influenzae*, and one each of *L. monocytogenes* and *S. aureus*), three *S. pneumoniae* which were resistant to erythromycin, clindamycin, pristinamycine, doxycycline, one *N. meningitidis* which was resistant to amoxicillin and rifampicin, and one *L. monocytogenes* which was resistant to trimethoprim and clindamycin (Table 1).

RTM investigations

In total, RTM detected pathogen genomes in 47 leftover CSF samples. Bacteria detected in 43 CSF samples included 20 *S. pneumoniae*, nine *N. meningitidis*, seven *H. influenzae*, three *S. agalactiae*, two *L. monocytogenes*, and one each of *E. coli* and *Staphylococcus aureus*. Viral pathogens detected in four CSF samples included three *Herpesvirus-1* and one *Varicella-Zoster Virus*. In addition, RTM yielded five negative CSF samples. In 63.5% CSF samples, pathogen genome detection in a 20-min sequencing run (Figure 3, Table 2) yielded 16 *S. pneumoniae*, seven *N. meningitidis*, six *H. influenzae*, one *S. agalactiae* and one VZV (median number of reads = 23), resulting from a blast analysis of MinION data against the in-house database and EPI2ME online analysis. A total of 47/52 CSF samples were detected as positive after two hours (median number of reads of 456.5). Genomic data analysis showed 19/47 (40.4%) of positive cases with >50% genome coverage including eight *S. pneumoniae*, five *N. meningitidis*, four *H. influenzae*, one each of *S. agalactiae* and VZV (Figure 1, Table 2). Viral cases were identified directly by blast against the in-house database and EPI2ME online software and confirmed by specific RT-PCR. False negative *S. pneumoniae* (sample 31) was confirmed by negative Illumina pair-end metagenomics, as previously described,¹⁶ may be due to failed DNA extraction and/or the limitation of RTM to detect low pathogen levels in the CSF (>35 Ct). The *in-silico* antibiogram analysis yielded 24/43 susceptible bacteria, 10 resistant bacteria and not realized in nine bacteria due to the low quantity of pathogens sequences (< 1% genome coverage) generated by MinION (Table 1).

Comparison between RTM and routine investigations

Two discordances in pathogen detection were noted by comparing RTM and routine investigation data. One case of *H. influenzae* (sample 20) detected by FilmArray® but not by RTM was eventually interpreted as a false-positive of multiplex RT-PCR,⁵ in agreement with a negative specific RT-PCR and culture. One case of *S. pneumoniae* (sample 31) was detected by multiplex RT-PCR but not by RTM. Further control by routine specific RT-PCR yielded a >35 Ct and culture remained negative. A total of 23/43 (53.5%) positive cases of bacterial meningitis diagnosed by BioFire FilmArray® failed in culture, and it was not possible to carry out an *in-vitro*

Patient	Age	Gender	Final diagnostic	Biofire FilmArray	MinION	Culture	Specific RT-PCR	In-vitro antibiogram	In-silico antibiogram
Sample 1	90	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Not realized	Macrolides and related groups (erythromycin, clindamycin, streptogramin b), tetracycline, trimethoprim	Macrolides and related groups (erythromycin, clindamycin, lincomycin, quinupristin, pristinamycine, virginiamycin), tetracycline, rifamycin, fluoroquinolone, aminoglycoside, phenicol
Sample 2	70	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Not realized	Macrolides and related groups (erythromycin, clindamycin, streptogramin b), tetracycline, trimethoprim	Macrolides and related groups (erythromycin, clindamycin, lincomycin), tetracycline, phenicol
Sample 3	25	M	<i>Neisseria meningitidis</i>	Positive	Positive	Positive	Not realized	Susceptible	Susceptible
Sample 4	80	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Not realized	Not realized	Not realized
Sample 5	84	M	<i>Listeria monocytogenes</i>	Positive	Positive	Positive	Not realized	Clindamycin, Trimethoprim	Not realized
Sample 6	66	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Not realized	Macrolides and related groups (erythromycin, clindamycin, streptogramin b), tetracycline	Macrolides and related groups (erythromycin, clindamycin, streptogramin b, quinupristin, pristinamycine, virginiamycin), tetracycline
Sample 7	65	F	<i>Haemophilus influenzae</i>	Positive	Positive	Positive	Not realized	Susceptible	Amoxicillin/ampicillin, piperacillin, ticarcillin, cephalothin
Sample 8	64	F	<i>Neisseria meningitidis</i>	Positive	Positive	Negative	Not realized	Not realized	Not realized
Sample 9	30	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Not realized	Susceptible	Not realized
Sample 10	57	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 11	2	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Not realized	Susceptible	Susceptible
Sample 12	45	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 13	0	F	<i>Streptococcus agalactiae</i>	Positive	Positive	Negative	Not realized	Not realized	Macrolides and related groups (erythromycin, spiramycin, azithromycin)
Sample 14	7	F	<i>Neisseria meningitidis</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 15	58	M	<i>Neisseria meningitidis</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 16	0	F	<i>Streptococcus agalactiae</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 17	33	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Not realized	Susceptible	Macrolides and related groups (erythromycin, clindamycin, streptogramin b, quinupristin, pristinamycine, virginiamycin)

Table 1 (Continued)

Patient	Age	Gender	Final diagnostic	Biofire FilmArray	MinION	Culture	Specific RT-PCR	In-vitro antibiogram	In-silico antibiogram
Sample 18	0	M	<i>Haemophilus influenzae</i>	Positive	Positive	Negative	Not realized	Not realized	Amoxicillin/ampicillin, piperacillin, ticarcillin, cephalothin
Sample 19	32	M	Negative	Negative	Negative	Negative	Not realized	Not realized	Not realized
Sample 20	28	F	Negative	<i>Haemophilus influenzae</i>	Negative	Negative	Negative	Not realized	Not realized
Sample 21	68	M	Herpes Simplex Virus 1	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 22	14	F	Herpes Simplex Virus 1	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 23	77	F	Herpes Simplex Virus 1	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 24	29	F	Varicella Zoster Virus	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 25	89	F	<i>Haemophilus influenzae</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 26	74	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Not realized	Not realized	Not realized
Sample 27	49	F	<i>Haemophilus influenzae</i>	Positive	Positive	Negative	Not realized	Not realized	Susceptible
Sample 28	22	M	<i>Listeria monocytogenese</i>	Positive	Positive	Positive	Not realized	Susceptible	Not realized
Sample 29	0	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible
Sample 30	78	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 31	11	M	<i>Streptococcus pneumoniae</i>	Positive	Negative	Negative	Positive > 35Ct	Negative	Not realized
Sample 32	22	M	<i>Neisseria meningitidis</i>	Positive	Positive	Negative	Positive	Not realized	Penicillin A
Sample 33	18	M	<i>Neisseria meningitidis</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 34	42	M	<i>Neisseria meningitidis</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 35	0	M	<i>Haemophilus influenzae</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible
Sample 36	0	M	<i>Haemophilus influenzae</i>	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 37	40	M	<i>Haemophilus influenzae</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 38	57	M	<i>Staphylococcus aureus</i>	Positive	Positive	Positive	Positive	Susceptible	Cefoxitin (mecA)
Sample 39	40	M	Negative	Negative	Negative	Negative	Negative	Not realized	Not realized
Sample 40	59	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 41	33	F	<i>Neisseria meningitidis</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible
Sample 42	18	F	<i>Streptococcus agalactiae</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 43	0	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible
Sample 44	20	F	<i>Neisseria meningitidis</i>	Positive	Positive	Positive	Positive	Susceptible	Penicillin A
Sample 45	23	F	Negative	Negative	Negative	Negative	Negative	Not realized	Not realized
Sample 46	51	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible
Sample 47	55	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Positive	Not realized	Susceptible
Sample 48	64	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible
Sample 49	0	M	<i>Escherichia coli</i>	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 50	6	M	<i>Streptococcus pneumoniae</i>	Positive	Positive	Negative	Positive	Not realized	Not realized
Sample 51	85	F	<i>Streptococcus pneumoniae</i>	Positive	Positive	Positive	Positive	Susceptible	Susceptible

Table 1: Concordance and discordance of routine diagnosis with RTM, in-vitro and in-silico antibiogram. Antibiotics not used in the routine are in green characters. The final diagnosis is the diagnosis validated in routine by clinicians.

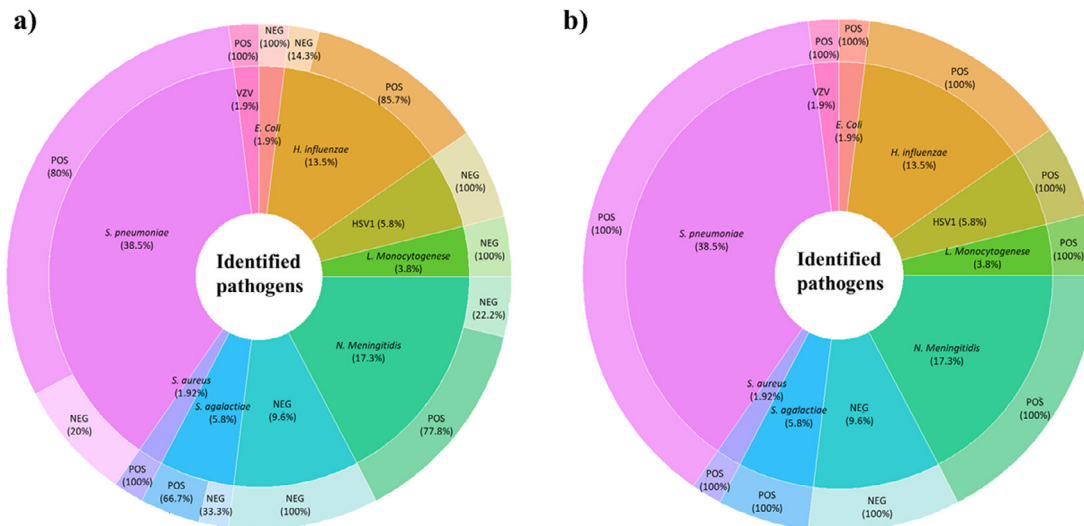


Figure 3. Real-time metagenomics data analysis and pathogen characterisation. a) Real-time data analysis and pathogen identification at 20 min sequencing. Thirty-one CSF samples were diagnosed as positive after a 20-min sequencing run, including 16 *S. pneumoniae*, seven *N. meningitidis*, six *H. influenzae*, one *S. agalactiae*, and one VZV. b) Total generated data and pathogen identification after a two-hour sequencing run. A total of 47/52 CSFs diagnosed positive after two hours RTM, included 20 *S. pneumoniae*, nine *N. meningitidis*, seven *H. influenzae*, three *S. agalactiae*, three *Herpes Simplex Virus*, two *L. monocytogenes*, one case each of *E. coli* and *S. aureus*, and one VZV.

antibiogram (Table 1). From the 16 susceptible bacteria which were identified in routine bacteriology, antibiotic susceptibility testing of cultured bacteria yielded 10/16 concordant *in-vitro* and *in-silico* antibiograms, while four bacteria (*H. influenzae*, *S. pneumoniae*, *N. meningitidis* and *S. aureus*) were *in-silico* resistant for beta-lactams, macrolides and related antibiotics, penicillin A, and cefoxitin respectively (Table 1), probably due to the absence of expression of resistance despite the possession of the antibiotic resistance encoding genes. In two cultured *S. pneumoniae* and *L. monocytogenes* the *in-silico* antibiogram had been not realised faced to the low quantity of data <1% (Table 1). Furthermore, partial discordance was observed between *in silico* and *in vitro* antibiogram for CSF sample “1” and “2”. For both CSFs, *in vitro* antibiogram identified a resistant *S. pneumoniae* for trimethoprim and streptogramin b. However, *in silico* antibiogram yielded a resistant *S. pneumoniae* to lincomycin, quinupristin, pristinamycine, rifamycin, fluoroquinolone, aminoglycoside and phenicol for CSF sample “1”, and a resistant *S. pneumoniae* to lincomycin and phenicol in CSF sample “2” (Table 1). In addition, an *in-silico* antibiogram was successfully performed in 22 uncultured bacteria, including 14 susceptible bacteria (five *S. pneumoniae*, four *N. meningitidis*, three *H. influenzae*, and two *S. agalactiae*), one *S. agalactiae* was *in-silico* predicted resistant to erythromycin, azithromycin and spiramycin; one *H. influenzae* *in-silico* predicted resistant for amoxicillin, ampicillin, cephalothin, piperacillin, ticarcillin and one *S. pneumoniae* *in-silico*

predicted resistant for erythromycin, streptogramin B, chloramphenicol and lincomycin. In addition, *in-silico* antibiogram could not be derived from genome sequencing following low quantity of genomic data, in five additional uncultured bacteria.

Bacteria genotyping

Genome-sequence-derived-MLST analysis yielded 13 bacterial serotypes derived from 34 MLST profiles. Sixteen genotyped *S. pneumoniae* yielded six serotypes (six 3A serotype, five 11A, two 6B, one 16F, one 5A and one 12F serotype), three *N. meningitidis* serotypes (five B serotype, two C serotype and one A serotype), three *H. influenzae* serotypes (three B serotype, two F serotype and one non-typable), three cases of *S. agalactiae* V serotype (Figure 4, Table 2). In addition, 50% of patients positive for *S. pneumoniae* have more than 55 years-old, with a dominance of 3A and 6B serotypes. Six out of nine *N. meningitidis* cases were aged 30 or under, with a dominance of the B-serotype. Four out of seven *H. influenzae* cases were aged 25 years or under and three were over the age of 45 years. The b-serotype was identified in patients aged 0, 45 and 65 years old, the F-serotype was identified in two patients aged 25 and 90 years old. One non-typable *H. influenzae* was identified in a 42-year-old patient. There was no association between bacteria serotypes, age, and gender of the patients. According to the low number of viral cases diagnosed in this series, virus genotyping was not performed.

Samples	Run time	Identied pathogen	Specific reads at 20 min	Total reads at 2 h	Specific reads at 2 h	Specific reads (%)	Number of nucleotide	Genome coverage	Genotyping
Sample 1	2H	<i>Streptococcus pneumoniae</i>	5897	174,156	33,465	19.22	1,940,109	95.17	Serotype 16F
Sample 2	2H	<i>Streptococcus pneumoniae</i>	7457	125,520	35,055	27.93	1,959,603	87.48	Serotype 6B
Sample 3	2H	<i>Neisseria meningitidis</i>	45	116,752	905	0.78	762,285	34.01	Serotype B
Sample 4	2H	<i>Streptococcus pneumoniae</i>	0	9137	6	0.065667068	4183	0.02	Not realized
Sample 5	2H	<i>Listeria monocytogenes</i>	0	480,000	21	0.00004375	14,597	0.5	Not realized
Sample 6	2H	<i>Streptococcus pneumoniae</i>	7	151,187	135	0.089293392	73,893	3.36	Serotype 11A
Sample 7	2H	<i>Haemophilus influenzae</i>	3471	180,169	26,014	14.43866592	1,751,169	88.37	Serotype b
Sample 8	2H	<i>Neisseria meningitidis</i>	0	3830	3	0.078328982	2687	0.1	Not realized
Sample 9	2H	<i>Streptococcus pneumoniae</i>	0	34,860	12	0.034423408	4578	0.22	Not realized
Sample 10	2H	<i>Streptococcus pneumoniae</i>	5	11,260	132	1.172291297	65,775	3	Serotype A5
Sample 11	2H	<i>Streptococcus pneumoniae</i>	3451	30,015	17,074	56.88489089	1,994,683	93.65	Serotype 3A
Sample 12	2H	<i>Streptococcus pneumoniae</i>	245	131,943	1652	1.252055812	1,346,254	63.2	Serotype 3A
Sample 13	2H	<i>Streptococcus agalactiae</i>	123	206,130	1017	0.493377965	390,297	19.38	Serotype V
Sample 14	2H	<i>Neisseria meningitidis</i>	4513	60,134	18,965	31.53789869	2,023,530	93.96	Serotype C
Sample 15	2H	<i>Neisseria meningitidis</i>	411	57,227	3713	6.488196131	1,644,843	73.43	Serotype A
Sample 16	2H	<i>Streptococcus agalactiae</i>	341	125,815	2058	1.635735008	1,504,297	71.3	Serotype V
Sample 17	2H	<i>Streptococcus pneumoniae</i>	239	6501	1175	18.07414244	517,575	24.3	Serotype 11A
Sample 18	2H	<i>Haemophilus influenzae</i>	314	9283	1406	15.14596574	1,318,171	66.52	Serotype b
Sample 19	2H	Negative	0	39,810	0	0	0	0	Not realised
Sample 20	2H	Negative	0	8110	0	0	0	0	Not realised
Sample 21	2H	<i>Herpes Simplex Virus 1</i>	0	4760	5	0.105042017	501	0.3	Not realized
Sample 22	2H	<i>Herpes Simplex Virus 1</i>	0	14,110	2	0.014174344	271	0.2	Not realized
Sample 23	2H	<i>Herpes Simplex Virus 1</i>	0	7082	9	0.127082745	1519	1	Unkown
Sample 24	2H	<i>Varicella Zoster Virus</i>	41	251,411	608	0.241835083	86,736	69.38	Unkown
Sample 25	2H	<i>Haemophilus influenzae</i>	967	775,438	4352	0.005612312	1,555,463	83.8	Serotype f
Sample 26	2H	<i>Streptococcus pneumoniae</i>	2	74,278	31	0.000417351	18,086	0.8	Not realized
Sample 27	2H	<i>Haemophilus influenzae</i>	8	14,994	63	0.004201681	2,8751	1.45	Serotype f
Sample 28	2H	<i>Listeria monocytogenese</i>	0	511,274	6	1.17354E-05	9104	0.3	Not realized
Sample 29	2H	<i>Streptococcus pneumoniae</i>	29	305,696	2566	0.00839396	1,138,209	51.8	Serotype 6B
Sample 30	2H	<i>Streptococcus pneumoniae</i>	46	52,353	136	0.00259775	44,612	3	Serotype 3A
Sample 31	2H	Negative	0	52,000	0	0	0	0	Not realised
Sample 32	2H	<i>Neisseria meningitidis</i>	63	64,276	1107	0.017222603	1,918,857	85.6	Serotype B
Sample 33	2H	<i>Neisseria meningitidis</i>	51	92,000	591	0.006423913	796,395	37.12	Serotype C
Sample 34	2H	<i>Neisseria meningitidis</i>	4	228,118	238	0.00104332	359,034	16.73	Serotype B
Sample 35	2H	<i>Haemophilus influenzae</i>	192	84,000	2751	0.03275	1,869,540	99.06	Non Typable

Table 2 (Continued)

Samples	Run time	Identied pathogen	Specific reads at 20 min	Total reads at 2 h	Specific reads at 2 h	Specific reads (%)	Number of nucleotide	Genome coverage	Genotyping
Sample 36	2H	<i>Haemophilus influenzae</i>	23	287,552	561	0.001950951	378,275	20.66	Serotype b
Sample 37	2H	<i>Haemophilus influenzae</i>	0	69,418	5	7.20274E-05	1282	0.06	Non Typable
Sample 38	2H	<i>Staphylococcus aureus</i>	24	288,211	477	0.001655037	2,19,011	7.81	Not realised
Sample 39	2H	Negative	0	60,069	0	0	0	0	Not realised
Sample 40	2H	<i>Streptococcus pneumoniae</i>	97	88,376	1125	0.0127297	6,20,988	29.15	Serotype 3A
Sample 41	2H	<i>Neisseria meningitidis</i>	356	91,198	8726	0.095681923	2,054,566	91.29	Serotype B
Sample 42	2H	<i>Streptococcus agalactiae</i>	0	7409	5	0.000674855	573	0	Serotype V
Sample 43	2H	<i>Streptococcus pneumoniae</i>	0	74,260	8	0.00010773	27,083	1.3	Serotype 12F
Sample 44	2H	<i>Neisseria meningitidis</i>	0	7160	1638	0.22877095	1,161,716	51.16	Serotype B
Sample 45	2H	Negative	Negative	82,070	0	0	0	0	Not realised
Sample 46	2H	<i>Streptococcus pneumoniae</i>	562	199,270	6047	0.030345762	2,064,033	96.9	Serotype 11A
Sample 47	2H	<i>Streptococcus pneumoniae</i>	12	4238	456	0.107597924	1,011,157	47.5	Serotype 3A
Sample 48	2H	<i>Streptococcus pneumoniae</i>	4358	163,052	82,665	0.506985502	2,081,073	97.7	Serotype 11A
Sample 49	2H	<i>Escherichia coli</i>	0	2950	2	0.000677966	/	/	Not realised
Sample 50	2H	<i>Streptococcus pneumoniae</i>	2	12,370	14	0.00113177	4009	0.2	Not realized
Sample 51	2H	<i>Streptococcus pneumoniae</i>	0	3221	2,094	0.650108662	783,321	36.8	Serotype 11A
Sample 52	2H	<i>Streptococcus pneumoniae</i>	435	82,070	7632	0.092993786	2,028,482	95.23	Serotype 3A

Table 2: partial and total genomic data, RTM analysis and bacteria genotyping.

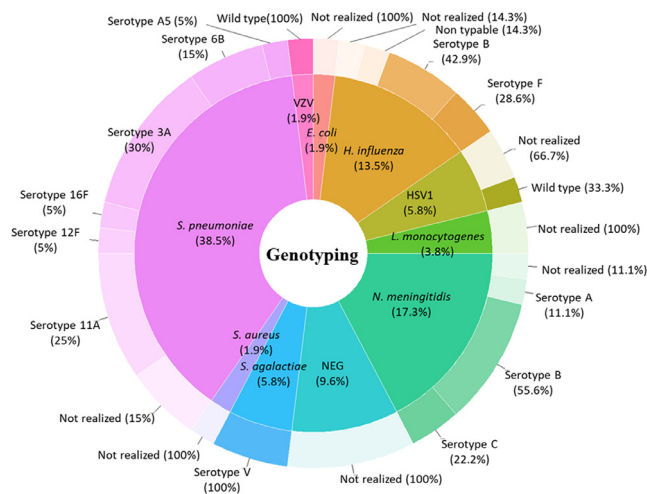


Figure 4. Pathogen genotyping and distribution according to the causative bacteria. Genotype investigation was performed on PubMLST database (<https://pubmlst.org/>) for bacteria pathogens and ViPR database (<https://www.viprbrc.org/>) for virus genotyping. Total of 13 bacterial serotypes identified from 34 MLST profiles: Six *S. pneumoniae* serotype (six 3A serotype, five 11A, four 6B, one 16F, one 5A and one 12F serotype), (six 3A serotype, five 11A, two 6B, one 16F, one 5A and one 12F serotype), three *N. meningitidis* serotypes (five B serotype, two C serotype and one A serotype), three *H. influenzae* serotypes (three B serotype, two F serotype and one non-typable), three cases of *S. agalactiae* V serotype. Whereas all viruses detected in this series were for wild-type serotype.

Virology data

Four viral infections were diagnosed in immunocompetent patients in this series (Table 1). Only DNA viruses were detected here, including three reactivated HSV-1 (samples 21, 22, 23), diagnosed in women aged 14 and 77 years and one 68-year-old man, all diagnosed with meningoencephalitis. In addition, wild-type VZV (<https://www.viprbrc.org/>) was detected in only one CSF (sample 24) collected in a seemingly immunocompetent 29-year-old woman with a past medical history of childhood VZV infection, no further recent contact with the virus including no vaccination, and no clinical zona; all data suggestive of VZV-reactivation (Table 2, Figure 4).

Discussion

We investigated RTM directly applied to left over CSF samples for the diagnosis of community-acquired meningitis at the POC laboratories in two university hospitals in southern France. Multiplex RT-PCR assays currently used routinely in the POC laboratory for this purpose only detect pieces of pathogen genome, providing detection and identification.⁵ This study indicated that, in addition to detection and identification, an RTM diagnostic strategy using Oxford Nanopore sequencing performed well on the diagnosis of known and non-routinely detectable pathogens in CSF samples, the antibiotic susceptibility profile, as well as their genotype.^{14,16,19} Moreover, cross-contamination, shown to limit the interpretation of positive multiplex assay results in several original studies and resulting meta-

analysis,^{5,20,21} was removed by the addition of a barcoding step; as illustrated here in one case of false-positive BioFire filmArray[®] *H. influenzae*.

In contrast to multiplex RT-PCR approaches which require specific conditions and equipment,^{5,6,22} RTM can be implemented in a surface area of less than two square metres in a POC laboratory using simple materials and with no requirement for advanced bioinformatics knowledge. This makes RTM a useful POC diagnostic tool, based on the simplicity, rapidity and cost-effectiveness of the process.^{15,23,24} Additional pieces of information were added concerning bacterial genotype/serotype, which are not applicable using conventional methods limited by the pathogen genome detection and culture, failing in 22/43 CSF, which required a specific PCR target for all pathogens.^{5,6} *In-silico* antibiogram investigation allowed to detect the presence of genes encoding for antibiotic resistance, which was concordant with the *in-vitro* investigation in 10/16 (62.5%) of cultured CSFs. In addition, RTM detected the presence of genes encoding for antibiotic resistance further phenotypically detected in three culture-positive CSFs and four additional failed bacteria cultures, despite the low level of pathogen in the CSF. In addition *in-silico* analysis of RTM data relieved supplementary information about antibiotic resistance mechanism detected in one *N. meningitidis* by the presence of *farB* gene encoding for efflux pumps, which was *in-vitro* resistant to rifampicin.²⁵ In addition, we found significant pathogen genotype diversity mostly represented by *S. pneumoniae* 3A, 11A and 6B serotypes, followed by *N. meningitidis* B and C-serotypes. The

H. influenzae b-serotype was identified most often in this study, followed by the non-typable and *H. influenzae* A serotype. This enabled the real-time genomic fine and accurate surveillance of bacteria genotypes and variants circulating in southern France, based on pathogen genome sequences (Figure 4), for the definition of a new strategy of infectious disease control including vaccination, as previously described in a case of non-typable *H. influenzae* meningitis identified in a patient vaccinated with the b-serotype.¹⁴ In addition, this strategy successfully detected four wild type viral DNA samples in agreement with routine multiplex RT-PCR,²⁶ validating its application for the direct investigation of DNA pathogens.

The limits we encountered reflect ways in which the method can be improved. The failure of RTM in one CSF (sample 31) which was detected positive in routine multiplex-RT-PCR indicated the need to increase RTM sensitivity for RT-PCR-detected pathogens with Ct > 35, given the higher sensitivity of the BioFire FilmArray[®] assay based on nested multiplex-RT-PCR.^{5,7} Increased sensitivity could be achieved by improving DNA extraction through an adapted automatic library preparation protocol for low pathogen levels, including microbial genome enrichment and human genome depletion.^{7,23,27} Also, the enlargement of the microbial panel included in the in-house database and its combination into one protocol DNA and RNA RTM is needed. This enlargement would allow for the one-shot detection of most pathogens responsible for community-acquired meningitis and meningoencephalitis, especially RNA viruses, the most frequently encountered causative agents of meningitis²⁸ and non-routinely diagnosed bacterial meningitis at the POC laboratory.¹⁹

Conclusion

This study goes beyond a few previous reports^{14,16,19} which all indicated that RTM has the potential to complement current multiplex RT-PCR assays for the rapid detection, genotyping and *in silico* antibiotic resistance profiling of pathogens responsible for community-acquired meningitis. This technique is already competitive in terms of time with the routine multiplex-based diagnostics in POC laboratories. Implementation of RTM as a POC diagnostic tool for life-threatening meningitis may provide real-time genomic surveillance of meningitis causative pathogen variants circulating in the study area, to define a new strategy of epidemiological control and vaccination. The authors are in the way to implement RTM in routine POC in selected situations including potential multiplex-PCR failures, based on herein reported diagnosis results along with a preliminary cost analysis and preliminary formation course for residents in medical biology (Table 3, Appendix 4).

Approach	Blunt/TA Ligase Master Mix	NEBNext [®] FFPE DNA Repair Mix	NEBNext [®] FFPE DNA Repair Mix	NEBNext [®] Quick Ligation TM *	Flow Cell (R9)	Ligation Kit	Flow Cell Priming Kit*	Flow Cell Wash Kit	Native Barcoding (PCR-free) *	Biofire FilmArray cartridge	Verifiable cost by test
RTM	1,44	6 €	9 €	12,72	54	45 €	5 €	13 €	11 €	0 €	143,3€
Biofire FilmArray	0	0	0	0	0	0	0	0	0	114	114€

Table 3: Cost analysis and comparison of RTM and routine FilmArray[®] test.

Further developments may include the application of RTM on cases of undocumented meningitis and RNA virus cases to enrich the repertoire of meningitis causative pathogens, non-routinely diagnosed in CNS diseases.

Contributors

MM experimental design, implementation of RTM workbench at the POC, software, creation of database, bioinformatics data analysis, interpretation and writing the original draft paper. QK ensured sample collection, helped with implementation of the RTM workbench at the POC, and data analysis and writing. AB, RS sample collection and clinical data. FS statistical analysis. CR, LH, and PEF conceptualisation, reagents, implementation of RTM workbench at the POC and validation. JPL and MD writing of the original draft paper, critical reviewing of the paper, validation, management of the work and funding. All authors read and approved the final version of the manuscript

Data sharing statement

All the extracted scaffold and contigs Fasta sequences corresponding to different pathogens were submitted to GenBank NCBI, available through BioProject No. PRJEB49201.

Declaration of interests

The authors declare no conflicts of interest. In particular, the authors did not receive any contribution from any of the suppliers mentioned in this report.

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Supplementary materials

Supplementary material associated with this article can be found in the online version at doi:10.1016/j.ebiom.2022.104247.

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