Molecular characterisation and antibiotic susceptibility of meningococcal isolates from healthy men who have sex with men

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ABSTRACT

Objectives To evaluate and characterise meningococcal carriage among healthy men who have sex with men (MSM) within a screening programme for Neisseria gonorrhoeae infection at the San Gallicano Dermatological Institute, Italy.

Methods A total of 441 MSM attending the STI/HIV Centre of the San Gallicano Dermatological Institute, Rome, Italy, in 2016 were routinely screened for N. gonorrhoeae infection by pharyngeal and rectal swabs. N. meningitidis isolates were evaluated for antibiotic susceptibility and characterised by whole genome sequencing. Genetic relationships among the meningococcal carriage isolates were determined using core genome multilocus sequence typing analysis. The soluble domain of AniA (sAniA) protein expression by western blotting was also evaluated.

Results A total of 62 (14.1%, 95% CI 11.1 to 17.6) carriage meningococci were found among 441 MSM. Forty-three viable N. meningitidis isolates were cultivated (42 from pharyngeal and 1 from rectal swabs). All the viable isolates were susceptible to ceftaxime, ceftriaxone, ciprofloxacin and rifampicin. Four isolates were penicillin G-resistant, and 73% of those penicillin G-susceptible showed a minimum inhibitory concentration from 0.064 μg/mL to 0.25 μg/mL. Serogroup B was the most frequent (44.2%), followed by Z (16.3%), E (9.3%), and Y and W (2.3%), respectively. Multilocus sequence typing analysis identified 29 sequence types belonging to 12 clonal complexes. The sAniA protein was expressed in 8 out of 28 (29%) screened meningococcal carriage isolates.

Conclusions Serogroup B meningococcal carriage identified from oral and anal specimens among healthy MSM was the most frequent serogroup identified in this study. Molecular evaluation revealed a degree of similarity among strains belonging to the same clonal complex.

INTRODUCTION

Neisseria meningitidis (Gram-negative diplococcus) is responsible globally for invasive meningococcal disease (IMD), in particular septicaemia and meningitis. Humans are the only hosts of N. meningitidis and can carry them harmlessly in the pharynx. 1 N. meningitidis can be transmitted through respiratory droplets from patients with IMD or from carriers. 1

N. meningitidis is transmitted from person to person, and close contact with a healthy carrier can facilitate the spread of the bacteria to susceptible individuals. 2 Meningococcal carriage was estimated to occur in 10% of the general population, with most people becoming a carrier multiple times in their lifetime and may carry the bacteria for several months. 3 Meningococcal carriage increases in teenagers, with peak in young adults at 24%–25%. 4 Carriage status increases in smokers, closed and semiclosed populations, such as military recruits and university students, and in household contacts of a case of meningococcal disease. 4–5

Meningococci can cross the epithelial barrier of the pharynx and gain access to the bloodstream, causing septicaemia. Moreover, it can penetrate the blood–brain barrier, giving rise to meningitis. 2 The human pharynx remains the site preferentially colonised by N. meningitidis. Although N. meningitidis is an aerobic bacterium, it can respond to oxygen limitation condition by inducing activation of the alternative pathway of denitrification, which is initiated by the copper-containing nitrite reductase (AniA). 6 Therefore, occasionally urogenital and rectal meningococcal colonisation has been reported, with evidence of transmission between sexual partners involving the pharynx, urogenital tract and rectum. 7 Molecular characterisation studies have been carried out to understand the epidemiology and the spread of this invasive meningococcal population. 8–9 Outbreaks of IMD among men who have sex with men (MSM) have been reported 10–11; however, few data are available on meningococcal carriage in MSM. 12–15

Based on a screening programme for N. gonorrhoeae conducted during 2016 at the San Gallicano Institute, Rome, Italy, we evaluated the prevalence of N. meningitidis carriage isolates among healthy MSM, and the antimicrobial susceptibility and the molecular characteristics, by whole genome sequencing (WGS), and the expression of AniA protein among the analysed isolates.

MATERIALS AND METHODS

Study population and sample management

MSM attending, from January to December 2016, the STI/HIV Centre of the San Gallicano Dermatological Institute Hospital, Rome, Italy, were
screened for *N. gonorrhoeae* infection by pharyngeal and rectal swabs. *N. meningitidis* isolates from the same plates used for *N. gonorrhoeae* growth were identified. All meningococcal isolates, with a pseudoanonymous identification (ID) code, were sent to the National Reference Laboratory (NRL) for IMD at the Istituto Superiore di Sanità for further characterisation.

**Antimicrobial susceptibility test**

Susceptibility to cefotaxime, ceftriaxone, ciprofloxacin, penicillin G and rifampicin was determined on all viable meningococcal carriage isolates using the minimum inhibitory concentration (MIC) test strip method (Liofilchem, Italy) on Mueller-Hinton agar (Thermo Scientific, USA), supplemented with 5% sheep blood. The breakpoints were those recommended by the European Committee on Antimicrobial Susceptibility Testing version 11.0 (1 January 2021; http://eucast.org/).

**Beta-lactamase test**
The presence of the bacterial enzyme beta-lactamase was determined on all viable *N. meningitidis* isolates by rapid acid metric stick method (Oxoid, Italy) on IsoVitaleX 2% Thayer-Martin agar plate incubated overnight in 5% carbon dioxide (CO₂) atmosphere at 37°C.

**Molecular characterisation**

**Genomic DNA extraction and WGS**

Genomic DNA from cultured meningococcal isolates (n=43) was extracted using QIAamp DNA Mini Kit (Qiagen, Hilden, Germany) according to the manufacturer’s instructions and was sent to the Army Medical Center Scientific Department for WGS. For each isolate, 1 ng of DNA was used to prepare the sequencing libraries, following the Nextera XT DNA protocol, according to the manufacturer’s instructions (Illumina, California, USA). WGS was performed on meningococcal carriage isolates using Illumina MiSeq Sequencer Kit (V3; 600 cycles). On average, 1.9 million paired-end reads were obtained for each sample. A first quality check of the raw sequence data was performed using FastQC. Reads were trimmed to keep high-quality bases (Q score >25) using the Sickle V.1.33 software. Sickle is a sliding-window, adaptive, quality-based trimming tool for FastQ files (V.1.33 software; available at https://github.com/najoshi/sickle). De novo assembly was carried out with the ABYSS V.1.5.2 software (k parameter=63). Contigs longer than 500 bp were selected using an ad hoc script and kept for further analysis. The final assembly ranged from 92 to 847 (median=211) contigs per sample covering ~2.2 Mb of the *N. meningitidis* genome.

**Genome analysis**

Of the 43 genomes from carriage isolates, only 32 were uploaded to the PubMLST Neisseria database (http://pubmlst.org/neisseria/). Genogroup definition, PorA and FetA typing, and multilocus sequence typing (MLST) were determined through the meningococcal typing website (http://pubmlst.org/neisseria/). Such information defined the finetype and genotype as follows: genogroup: PorA(1.)VR1,VR2:FetA(F) VR:ST(cc).

WGS was used to identify penA and rpoB alleles, serogroup B vaccine antigen variants, and ania allele. Genomes were compared by means of the Bacterial Isolate Genome Sequence Database (BIGsdB) Genome Comparator Tool within the PubMLST Neisseria database through gene-by-gene analysis. Phylogenetic analysis was performed using the *N. meningitidis* core genome MLST (cgMLST) scheme (1605 loci) on 32 carriage isolates of this study, plus 7 invasive meningococci isolated during the study period (2016) in Rome and collected by the NRL within the National Surveillance System for IMD. These invasive meningococci were two serogroup B/cc1572 (ID41657 and ID52942), one serogroup B/cc32 (ID41658), one serogroup B/cc213 (ID52946), two serogroup C/cc11 (ID52933 and ID52934), and one serogroup C/cc334 (ID52961).

Incomplete loci were removed automatically from the distance matrix calculation for the neighbour-net graphs. The resulting output distance matrix was used to generate neighbour-net networks using SplitsTree4 software (V.4.13.1).

**Western blotting**

To determine the expression of AniA protein, western blotting assay was performed on a subsample of 28 meningococcal carriage isolates, according to previously described method, with minor modifications. A polyclonal antiserum (Bio-Fab Research, Italy) raised against the soluble domain of AniA protein (sAniA) of *N. meningitidis* serogroup B reference strain MC58 was used for experiments.

Meningococcal isolates were cultured in standard growth conditions at 37°C with 5% CO₂ on IsoVitaleX 2% Thayer-Martin plates. Whole cell lysates and supernatants were prepared with 5% Laemmlı buffer. Sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) was performed on 12% polyacrylamide gel. Polyclonal rabbit antiserum prepared with 5% Laemmli buffer. Sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) was performed on 12% polyacrylamide gel. Polyclonal rabbit antiserum against AniA protein was used as primary antibody at 1:1000 dilution, and the anti-rabbit IgG conjugated to alkaline phosphatase (Merck Life Science, Italy) was used as secondary antibody at 1:15 000 dilution.

Finally, 5-bromo-4-chloro-3-indolyl-phosphate/nitro blue tetrazolium (BCIP/NBT) Alkaline Phosphatase Substrate (Merck Life Science) was used to visualise the sAniA protein. *N. meningitidis* MC58 strain was used as the control strain in all experiments.

**RESULTS**

**Demographic and clinical data**

Sixty-two (14.1%, 95% CI 11.1 to 17.6%) *N. meningitidis* isolates were identified from 441 MSM. Among them, 43 viable *N. meningitidis* isolates, 42 from pharyngeal and 1 from rectal swabs, were analysed. The median age of the subjects was 36 years (IQR: 23–52 years). All except for two subjects were Italians and confirmed as negative for other STIs.

**Antimicrobial susceptibility**

Antimicrobial susceptibility was performed on 37 out of 43 *N. meningitidis* carriage isolates. All of them were susceptible to cefotaxime, ceftriaxone, ciprofloxacin and rifampicin. Among the isolates susceptible to penicillin G (97%, 33 of 37), 27 showed MIC for penicillin G higher than 0.064 μg/mL (0.064 μg/mL < MIC <0.25 μg/mL) and 4 were penicillin G-resistant (PenR), with MIC values ranging from 0.38 μg/mL to 0.50 μg/mL. All four penicillin-resistant isolates were beta-lactamase-negative.

**Molecular characterisation**

The meningococcal genogroup was identified in the 43 meningococcal carriage isolates, with 19 (44%) serogroup B (MenB), 7 (16%) MenZ, 4 (9%) MenE, 1 MenY and 1 MenW. A total of seven isolates (16%) were cwl and four (9%) were non-groupable (NG).
MLST identified 29 sequence types (STs) belonging to 12 clonal complexes (ccs), of which cc162 (n=5, 42%) and cc461 (n=3, 25%) were the most frequent. Twelve sequence types, identified in 12 isolates (4 MenB, 4 MenZ, 2 NG, 1 MenE and 1 cnl), were not designed to any known clonal complex and were classified as ‘not assigned’ (na) (table 1).

Among the MenB isolates, cc461 (n=3), cc1572 (n=3), cc32 (n=2), cc41/cc44 (n=2), cc4821 (n=2), cc162 (n=1), cc213 (n=1) and cc269 (n=1) were found. Only one of the four MenE isolates belonged to cc1157; the MenW isolate belonged to cc11 and the MenY isolate belonged to cc174.

Among the cnl isolates, cc162 (n=3), cc1136 (n=2) and cc53 (n=1) were identified (table 1). A total of 29 PorA types and 16 FetA variants were identified. The most frequent PorA types were P1.7-12,14 (3 of 43, 7%) and P1.21-19,16-16 (3 of 43, 7%). The predominant FetA variant was F1-62 (6 of 43, 14%) and F5-7 (6 of 43, 14%). One carriage isolate (1 of 43, 2.3%) gave negative PCR results for fetA gene (table 1).

### Meningococcal serogroup B vaccine antigen variants

4CMenB variant family 2/Pfizer subfamily A (fHbp-2/A) was the most represented (49%, 21 of 43), followed by variant family 1/Pfizer subfamily B (fHbp-1/B) (21%, 9 of 43) and variant family 3/subfamily A (fHbp-3/A) (7%, 3 of 43).

#### Table 1 Molecular characteristics of 43 meningococcal carriage isolates

<table>
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<tr>
<th>Serogroup</th>
<th>Identification (<a href="http://pubmlst.org/Neisseria">http://pubmlst.org/Neisseria</a>)</th>
<th>Clonal complex</th>
<th>Sequence type</th>
<th>Finetype</th>
<th>Number of isolates</th>
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</table>

The table was created by the authors.

‘–’ indicates no genome available.

‘na’ indicates not assigned to any known clonal complex.

cnl, capsule null locus; NG, non-groupable; ΔF, fetA gene deletion.
Remaining isolates (35 of 43, 81%) resulted negative to the \textit{nadA} gene.

Twenty-one PorAVR2 peptides were identified in 34 isolates. The most frequent PorAVR2 peptide was P1.14 (n=6), followed by P1.16-6 (n=3). Six PorAVR2 peptides (P1.16-75, P1.23, P1.30, P1.13-1, P1.13-9, P1.16-11) occurred twice and the remaining 15 PorAVR2 peptides occurred once.

\textit{penA} and \textit{rpoB} alleles

Identification of \textit{penA} and \textit{rpoB} alleles was performed on 43 meningococcal carriage isolates. A total of 16 different \textit{penA} alleles were found. Four \textit{penA} alleles were identified in the four isolates resistant to penicillin G, of which \textit{penA179}, \textit{penA209} and \textit{penA295} were identified in the isolate with MIC of 0.38 µg/mL and \textit{penA327} in the isolate with MIC of 0.5 µg/mL.

The two prevalent \textit{rpoB} alleles were \textit{rpoB21} (10 of 43, 23%) and \textit{rpoB28} (9 of 43, 21%), followed by \textit{rpoB2} (5 of 43, 12%), \textit{rpoB5} (4 of 43, 9%), \textit{rpoB1} (3 of 43, 7%), \textit{rpoB4} (3 of 43, 7%), \textit{rpoB207} (3 of 43, 7%), \textit{rpoB18} (3 of 43, 7%) and \textit{rpoB83} (2 of 43, 5%). \textit{rpoB8}, \textit{rpoB9}, \textit{rpoB34} and \textit{rpoB42} were detected each one in one isolate.

Core genome multilocus sequence typing

Thirty-two genomes of 43 meningococcal carriage isolates and 7 invasive meningococcal genomes were analysed using cgMLST to characterise the phylogenetic correlations. The cgMLST analysis included 1269 of the 1605 core genome loci, excluding 336 incomplete loci. All meningococcal carriage and invasive genomes were distributed based on clonal complex or sequence type (figure 3). The genome comparison revealed similarities and close relationships among genomes from meningococcal carriage isolates. The two MenZ/ST-13741 (ID63038 and ID61182) had no different loci. Moreover, these two genomes differed by 15 loci with ID61178 (MenZ/ST-13741) and by 55 loci with ID61174 (MenZ/ST-13741). Other two MenZ ID61164 (ST-12943) and ID61169 (ST-13746) showed a distance of 17 loci. The three MenE/ST-13740 genomes showed close relationships: between ID63037 and ID61180, isolated from the pharynx and rectal swabs of the same patient, respectively, a distance of 41 different loci were found. Moreover, ID61180 showed only three different loci with ID61181.

The two cc162 genomes from carriage isolates (ID61160 and ID61168) revealed a small genetic distance of 19 loci.
We observed that meningococcal carriage and invasive genomes, belonging to the same clonal complex, grouped together (figure 3). In particular, the two MenB/cc213 genomes (ID63041 from carriage isolate and ID52946 from invasive isolate) clustered together with a distance of 37 loci.

**Western blotting**

As shown in table 2, 8 out of 28 (29%) screened meningococcal carriage isolates on western blot assay for the expression of sAniA showed the ~55 kDa sAniA form. These isolates were MenE (n=3), MenB (n=2), MenY (n=1), MenZ (n=1) and cnl (n=1). The molecular characterisation evidenced six aniA alleles: 6, 11, 53, 229, 673 and 675. The aniA allele 675 (n=3) was found in three MenE/ST-13740 genomes (na).

AniA protein was not expressed in the remaining 20 isolates (71%). In four meningococcal isolates, the aniA gene was missing. In one isolate belonging to the genotype W:P1.5,2:F1-1:ST-11 (cc11), a premature stop codon was found in the aniA gene (aniA allele 1) (table 2). The remaining 15 isolates harboured an intact aniA gene but did not express the protein.

**DISCUSSION**

Isolation of invasive *N. meningitidis* from MSM has been already reported in several studies,11–13 and an increasing number of outbreaks of meningococcal disease has been reported in Europe and in the USA in the same population.14–15 22 23

In this study, a cohort of MSM participants at a screening programme for *N. gonorrhoeae* infection were considered for the presence of *N. meningitidis* in the oropharynx and urogenital tracts.

The proportion of meningococcal carriage (14.1%) was consistent with that already described by Carlin et al.14 While invasive meningococcal isolates from MSM found in Europe were predominantly serogroup C,11 this serogroup was absent among our carriage isolates. Majority of the carried meningococci were of serogroup B, one of the two most frequent serogroups of invasive meningococcal isolates circulating in Italy in the same period (https://www.iss.it/sn-mbi-rapporti-iss).

The clonal complexes of the carriage meningococci analysed accounted for the most frequently clonal complexes reported in healthy carriers.3 4 Only three carriage isolates belonged to the hypervirulent clonal complexes (cc41, cc44 and cc11), mainly associated with invasive meningococci circulating in the country.24

Phylogenetic analysis to investigate the genetic relationships among these carriages and invasive meningococci, isolated in Rome in the same period, revealed clusters including isolates belonging to the same clonal complexes regardless of the invasive-ness. We can hypothesise that carriage meningococci belonging to a hypervirulent clonal complex may be widely transmitted in the MSM community and may determine an invasive disease. Moreover, the high degree of genetic similarity for carriage isolates belonging to the same clonal complex may suggest the presence of a cluster of those isolates among the screened MSM.

Evaluation of antibiotic susceptibility revealed a high proportion of carriage isolates with reduced susceptibility to penicillin G and resistance to penicillin G in four carriage isolates. Among the resistant carriage isolates to penicillin G, one *penA327* allele was already described in the meningococcal isolates from the outbreak of IMD among MSM.25 This allele is identical to the allele *penAXXXIV*, associated with resistance to third-generation cephalosporin and reduced susceptibility to cefotaxime in *N. gonorrhoeae*.25 However, *penA327* allele was associated with resistance to penicillin G but not with resistance to cephalosporin and reduced susceptibility to cefotaxime. These results correlate with the possibility of a genetic exchange among...
The ability of *N. meningitidis* to grow under oxygen limitation through denitrification pathway has facilitated its adaptation to environments where oxygen availability is variable. While this pathway is crucial to the survival of *N. gonorrhoeae*, the denitrification pathway is not essential in the survival of *N. meningitidis* in the nasopharynx. Hereby, 29% of the carriage meningococci expressed the AniA protein except for 5 isolates, lacking the gene or with a premature stop codon in the first half of the nucleotide sequence, while the other 15 meningococcal isolates were unable to express the protein despite the presence of the entire *aniA* gene. Since the expression of meningococcal AniA protein is under the control of a complex regulatory network involving multiple transcriptional regulators, it is possible to hypothesise that nucleotide polymorphisms in the putative regulatory sites explain the inability of these meningococci to produce AniA.

A report published by Taha et al revealed that meningococcal isolates collected from the urethra harboured a functional AniA, but were defective for fHbp due to a premature stop codon. This phenotype (AniA+, fHbp−) could reflect an adaptation and selection of meningococci which colonised uncommon sites, such as the urethra and the rectum. This result differs from the data here presented, with majority of the isolates from the pharynx demonstrating meningococcal carriage on September 10, 2022 by guest. Protected by copyright.http://sti.bmj.com/ Sex Transm Infect: first published as 10.1136/sextrans-2021-055173 on 17 November 2021. Downloaded from http://sti.bmj.com/ Sex Transm Infect: first published as 10.1136/sextrans-2021-055173 on 17 November 2021. Downloaded from...
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It is important to identify the meningococcal serogroup to assess how and if these strains may be preventable by vaccination. Information on serogroups could be essential in the implementation of public health control measures in case of an outbreak, including vaccination, which has been reported to manage an outbreak of IMD of serogroup C in MSM in New York City.1,2

Key messages

⇒ Serogroup B was the main serogroup identified among the meningococci isolated from healthy men who have sex with men (MSM) in this study.

⇒ The acquisition of penA327 allele in one penicillin G-resistant meningococcal carriage isolate highlights the possibility of exchange, in the pharynx, of genetic materials among meningococci and gonococci.

⇒ This study underlines the importance of pharyngeal and rectal swab screening to detect Neisseria meningitidis among MSM.

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