

Molecular Diversity of *Staphylococcus aureus* Colonizing the Upper Respiratory Tract of Residents and Staff in A Nursing Home

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Abstract

Elderly people living in nursing homes are a high-risk population for *Staphylococcus aureus* infection. Multiple comorbidities, a weakened immune system, inadequate hygienic conditions, and crowding might increase the prevalence rates of this opportunistic pathogen. However, the epidemiological aspects, genetic diversity, and transmission of *S. aureus* in nursing homes are still poorly understood, especially in Poland. This study aimed to determine the genetic relatedness and prevalence of colonization of *S. aureus* isolated from the anterior nares and the throat of residents and staff in a nursing home located in Lublin, Poland. The study showed a high *S. aureus* prevalence rate among participants (46.1%), yet there was a low frequency of MRSA strains among residents (1.7%) and staff (0%). The multiple-locus variable-number tandem-repeat fingerprinting (MLVF) analysis demonstrated a high degree of genetic diversity of *S. aureus* strains colonizing the anterior nares and the throat of the participants. The occurrence of simultaneous colonization with more than one unique *S. aureus* strain in any one individual as well as the incidence of colonization with the same genetic variant of *S. aureus* in different individuals was observed. These findings suggest that inter-participant *S. aureus* transmission might contribute to the development of cross-infections.

Key words: *Staphylococcus aureus*, colonization, elderly, genotyping, nursing home

Introduction

Staphylococcus aureus is an opportunistic human pathogen known for its increasing prevalence in both hospitals and the general community. An incremental rise in drug-resistance and high disease-causing capabilities has also been observed (Zhang et al. 2015; Zurita et al. 2016). The most common anatomical location for the commensal carriage of *S. aureus* is the anterior nares (Kluytmans et al. 1997; Pollitt et al. 2018), although recent epidemiological studies highlight the role of pharyngeal carriage in *S. aureus* transmission (Nilsson and Ripa 2006; Hamdan-Partida et al. 2010). The complexity of host-colonizer interactions has led researchers to investigate alternative contexts in pursuit of the mechanism for colonization. Long term care facilities (e.g. nursing homes) constitute communities in which crowding and inadequate hygienic conditions might contribute to higher MSSA (methicillin-sensitive *S. aureus*) and MRSA (methicillin-resistant *S. aureus*) prevalence rates. Chronic diseases, immune system

disorders, and frequent hospitalization make elderly people vulnerable to *S. aureus* infection (Ledell et al. 2003; Chuang et al. 2015; Peters et al. 2017).

S. aureus is also one of the most genetically diverse bacteria and the mechanisms by which the predominant strains are selected within each biological niche depends on the geographic location as well as the studied human host (Zurita et al. 2016). MSSA strains are more prevalent than MRSA strains and have a higher genetic diversity, whereas MRSA outbreaks are usually caused by only a few different MRSA clones (Goering et al. 2008). Therefore, analyzing the genetic relatedness of colonizing MSSA strains in nursing homes might result in a better understanding of *S. aureus* epidemiology.

We aimed at determining the genetic diversity and relatedness of MSSA and MRSA strains colonizing the upper respiratory tract of nursing home residents and staff using the MLVF (multiple-locus variable-number tandem-repeat fingerprinting) method. We hypothesized that there would be the same genetic MLVF

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banding patterns of *S. aureus* retrieved from different individuals, and the presence of genetic homogeneity might indicate that bacterial transmission has occurred.

Experimental

Materials and Methods

Bacterial strains. Among 102 participants enrolled in our study, more than three-quarters were female and the mean age of the residents (81.5 years old) was almost twice as high as that of the staff members (43 years old) (Table I). Two anatomical locations (the anterior nares and the throat) were swabbed using sterile cotton swabs (Medlab, Poland) premoistened with sterile 0.9% saline solution (POCH, Poland). The samples were immediately transported to the laboratory and processed within two hours. The swabs were inoculated on 5% sheep blood agar plates (BioRad, USA) and mannitol salt agar plates (BioMaxima, Poland) at 35.5°C for 24 h. The β -hemolytic or/and mannitol-positive colonies were tested for the presence of coagulase (Biomed, Poland) and finally identified as *S. aureus* using the Vitek 2 Compact automated system and GP cards (Biomérieux, France).

Table I
Basic demographic characteristics of participants and the prevalence of *S. aureus* colonization.

	Total (n = 102)	Residents (n = 60)	Staff (n = 42)
Age (years)			
Median (range)	68 (26–97)	81.5 (56–97)	43 (26–67)
Gender n (%)			
Female	78 (76.5)	43 (71.7)	35 (83.3)
Male	24 (23.5)	17 (29.3)	7 (16.7)
<i>S. aureus</i> colonization n (%)			
Nares	17 (36.2)	10 (35.7)	7 (36.8)
Throat	11 (23.4)	6 (21.4)	5 (26.4)
Nares and throat	19 (40.4)	12 (42.9)	7 (36.8)

Antimicrobial susceptibility testing was conducted using the Vitek 2 automated system and AST-P644 cards (BioMérieux, France) containing the following antibiotics: ceftaroline, ciprofloxacin, clindamycin, daptomycin, erythromycin, gentamicin, levofloxacin, linezolid, oxacillin, rifampicin, teicoplanin, tetracycline, tigecycline, trimethoprim/sulfamethoxazole, and vancomycin. Additionally, resistance to cefoxitin (30 μ g), tobramycin (10 μ g), fusidic acid (10 μ g) and mupirocin (200 μ g) was determined using disc diffusion method. D-test was conducted to determine the mechanism

of macrolide, lincosamide and streptogramin resistance (MLS_B). All discs were obtained from the same manufacturer (Becton Dickinson, USA). Disc-diffusion tests were performed and interpreted according to EUCAST recommendations (EUCAST 2018). All *S. aureus* isolates have been stored in freezers (–70°C) for further analysis using a medium of tryptic soy broth (TSB; BTL, Poland) and glycerol (POCH, Poland) in equal proportions (1:1).

Multiple-locus variable-number tandem-repeat fingerprinting (MLVF). The MLVF genotyping was performed to analyze the genetic diversity and relatedness of all 66 identified *S. aureus* strains. The MLVF method involved a multiplex-PCR reaction targeting five variable number of tandem repeats (VNTRs) in the following loci: *sspA*, *spa*, *sdr*, *clfA*, and *clfB*. These five VNTRs are located within the genes that encode the following proteins: serine proteinase V8, protein A, *sdrC*, *sdrD*, *sdrE* fibrinogen binding proteins, clumping factor A, and clumping factor B (Sabat et al. 2003).

Genomic DNA was extracted from the 24-h TSB culture prepared from a single bacterial colony. DNA extraction and purification were conducted according to the manufacturer's instruction (Genomic Mini, A&A Biotechnology, Poland), with the minor alteration of replacing lysozyme with an equal volume of lysostaphin (A&A Biotechnology, Poland). The quality and quantity of the DNA were assessed using BioPhotometer (Eppendorf, Germany) by measuring the absorbance at the wavelengths of 260 and 280 nm.

The PCR program consisted of predenaturation at 94°C for 5 min, 25 cycles of denaturation at 94°C for 30 s, annealing at 55°C for 30 s, extension at 72°C for 60 s, followed by a final extension at 72°C for 10 min (Biometra, Germany) (Grzegorzczak and Malm 2014). The PCR products and DNA molecular size marker (100 bp DNA Ladder Plus, ThermoScientific, USA) were resolved in 2% agarose electrophoresis gels (Sigma-Aldrich, USA) stained with SimplySafe dye (EURx, Poland). The banding patterns were photographed, exported to BioGene (Vilber Lourmat, France), and analyzed. The dendrogram was constructed using a UPGMA algorithm (unweighted pair group method with arithmetic mean) by applying the Dice Similarity Ratio. A tolerance level of 1% was used in comparing the banding patterns.

Results and Discussion

Prevalence of *S. aureus* colonization. Among all participants, 46.1% (47/102) were colonized with *S. aureus* (Table I). The pattern of colonization was divided into three categories based on the anatomical location. The category that registered the highest number of individuals was the combination of the anterior

nares and the throat (40.4% of colonized individuals), followed by the anterior nares only (36.2%), and the throat only (23.4%).

S. aureus colonizes multiple anatomical locations and the frequency of colonization in various niches might vary depending on the methodology and the context of the study. Our study demonstrated that the anterior nares are colonized more frequently than the throat, which is consistent with the findings of previous studies on the prevalence of both MSSA and MRSA occurring together (Roghmann et al. 2015; Zhang et al. 2015; Peters et al. 2017). Alternatively, some researchers have reported that colonization only in the throat is the most prevalent pattern (Nilsson and Ripa 2006; Hamdan-Partida et al. 2010) or have observed the high prevalence of extra-nasal colonization as well, i.e., in the groin and the perianal regions (Mody et al. 2008).

In the current study, 31.6% (6/19) of the participants colonized with two *S. aureus* strains (one in the anterior nares and one in the throat) registered dissimilar MLVF banding patterns, implying that the two *S. aureus* strains were unrelated. A similar observation has been made by Mongkolrattanothai et al. (2011), who noted that simultaneously carriage of two or more unrelated *S. aureus* isolates was observed in 30.4% of positive swab samples collected from children. Colonization with more than one strain of bacteria entails a risk of heteroresistance, i.e., a situation in which an antibacterial agent is effective against one strain, but not against another strain of the same bacteria. Truly, in cases of co-colonization with MSSA and MRSA both together, heteroresistance might pose a therapeutic challenge.

Antimicrobial resistance profile of *S. aureus*.

Table II shows the resistance profiles of *S. aureus* strains with the information about their location in the dendrogram. Among all *S. aureus* strains tested, 10.6% (7/66) were resistant to tetracycline; 6.1% (4/66) to macrolides, lincosamide, and streptogramin; 4.5% (3/66) to tobramycin and/or gentamicin, and 1.5% (1/66) to ciprofloxacin, levofloxacin, and daptomycin. We observed the presence of the MLS_B -constitutive resistance only. The share of *S. aureus* isolates susceptible to all of the antimicrobial agents tested was high (77.3%, 51/66), although the presence of MDR (multi-drug resistant) MRSA strain (Table II) suggests the need of conducting longitudinal studies investigating its transmission and long-term colonization patterns (Sun et al. 2017). The antimicrobial susceptibility testing identified 98.5% (65/66) of *S. aureus* strains as MSSA, whereas only one isolate was identified as MRSA. Consequently, the MRSA prevalence rate in our studies was low when compared to similar studies: 1% among all participants and 1.7% specifically among the residents. The prevalence of MRSA colonization among residents of long term care facilities across the world

Table II
The antimicrobial resistance profiles of *S. aureus* strains isolated from residents and staff in a nursing home.

Main cluster	Sub-cluster	<i>S. aureus</i> strain	Resistance profile
A	IV	R23T	TET
	IV	S123T, S123N	TET
	IV	S133T, S133N	TET
	V	S116T	TOB, GEN
B	VII	R38T	FOX, TOB, CIP, LEV, DAP
	IX	R42T, R42N	TET
	IX	S139N	E, CC
	X	R2T	E, CC
	XI	R33T, R33N	E, CC
	XII	S134N	E, CC
-	-	R41N	TET, TOB, GEN

Note. *S. aureus* strains marked with the same number were retrieved from the anterior nares (N) and the throat (T) of one resident (R) or staff member (S) and had 100% genetic homology according to the MLVF analysis. Tetracycline (TET), tobramycin (TOB), gentamicin (GEN), cefoxitin (FOX), ciprofloxacin (CIP), levofloxacin (LEV), daptomycin (DAP), erythromycin (E), clindamycin (CC).

varies within a wide range: from 0% in nine Swedish nursing homes (Andersson et al. 2012), 3.7% in nine nursing homes in Brasil (da Silveira et al. 2018), 7.2% in 19 long term care facilities in Luxemburg (Mossong et al. 2013), 20.4% in 36 residential care homes for the elderly in Hong Kong (Chuang et al. 2015), to 28% in 13 nursing homes in Michigan and Maryland (Roghmann et al. 2015).

Multiple-locus variable-number tandem repeat fingerprinting of *S. aureus*. Using molecular typing methods to determine the genetic diversity of human pathogens is crucial in identifying the reservoirs and routes of transmission (Grzegorzczak and Malm 2014). Unlike other bacterial genotyping methods, such as PFGE (pulsed-field gel electrophoresis), MLST (multi-locus sequence typing) or spa typing, MLVF has a comparable resolution, but is cheaper, less time- and labour-consuming (Luczak-Kadlubowska et al. 2008; Holmes et al. 2010; Kosecka-Strojek et al. 2016). The high discriminatory power of MLVF is its main advantage, enabling precise detection of single events of transmission between patients (Karynski et al. 2008).

The MLVF of the 66 *S. aureus* strains isolated from 47 individuals resulted in 51 banding patterns: 36 unique banding patterns were represented by only one strain and 15 banding patterns consisted of two identical *S. aureus* strains each. In accordance with our results, a high genetic diversity of MSSA has been reported in other studies (Emaneyni et al. 2011; Karynski et al. 2008; Zurita et al. 2016). Zurita et al. (2016) observed 69 MLVA (multiple-locus variable-number

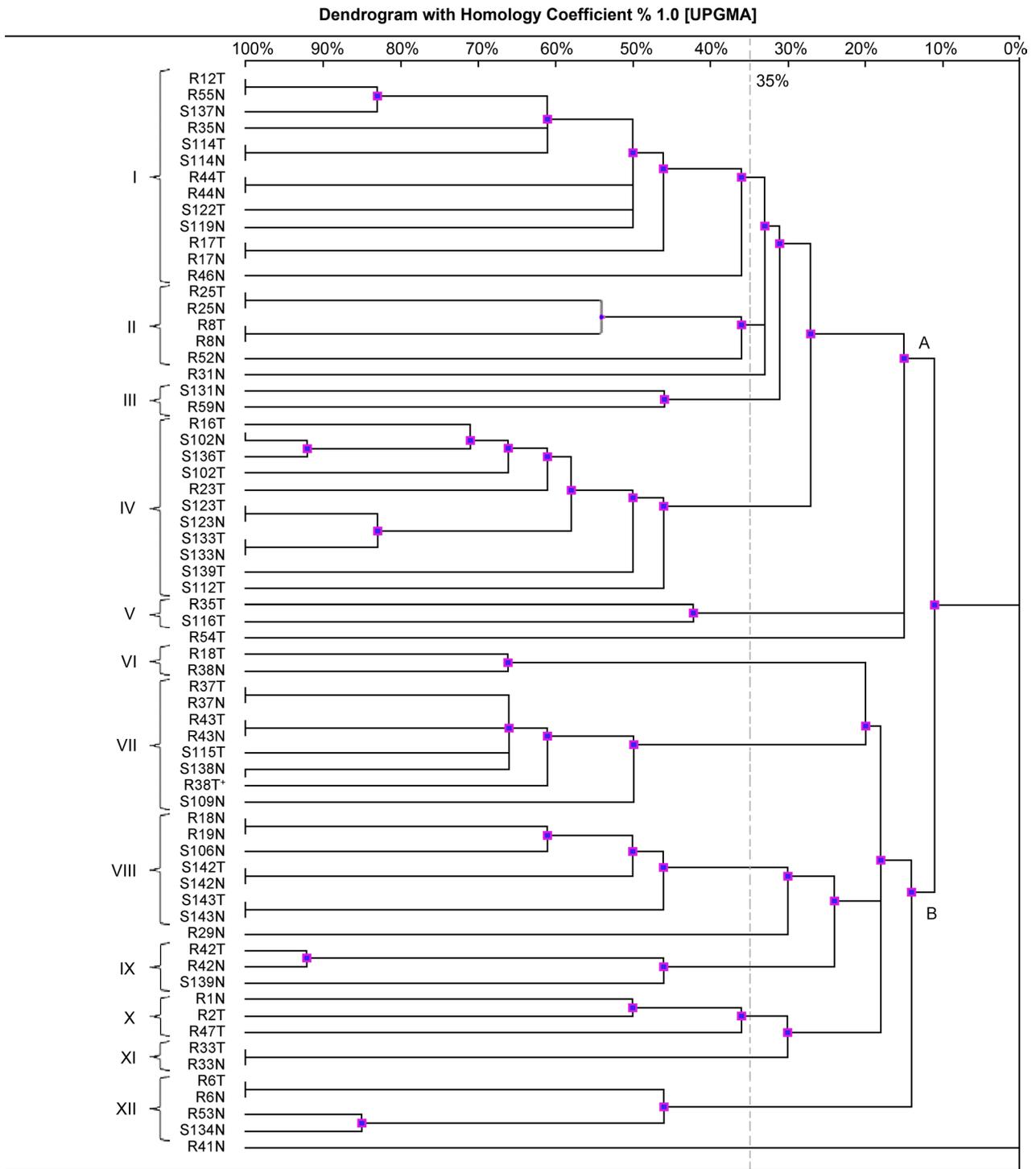


Fig. 1. MLVF dendrogram of *S. aureus* strains isolated from the anterior nares and the throat of residents and staff in a nursing home.

Note. The name of *S. aureus* strain contains the following information: R – collected from resident, S – collected from staff member, number characteristic for each participant, N – nasal swab, T – throat swab. Two main clusters (A and B) and twelve subclusters (I–XII; cut-off > 35%) are marked in the dendrogram. MRSA strain (R38T) is marked with an asterisk.

tandem-repeat analysis) genotypes among 70 tested MSSA strains from various populations.

The MLVF typing allowed us to divide the 66 *S. aureus* isolates into two main clusters, herein designated as A and B (Fig. 1). One *S. aureus* strain (R41N) was not compatible with any other banding pattern.

In the dendrogram, it represents a singleton with no association to the main clusters. The main cluster A was divided into five subclusters (herein designated I–V); the main cluster B was divided into seven subclusters (herein designated VI–XII). Altogether, the twelve subclusters included 94% (62/66) of all strains (cut-off

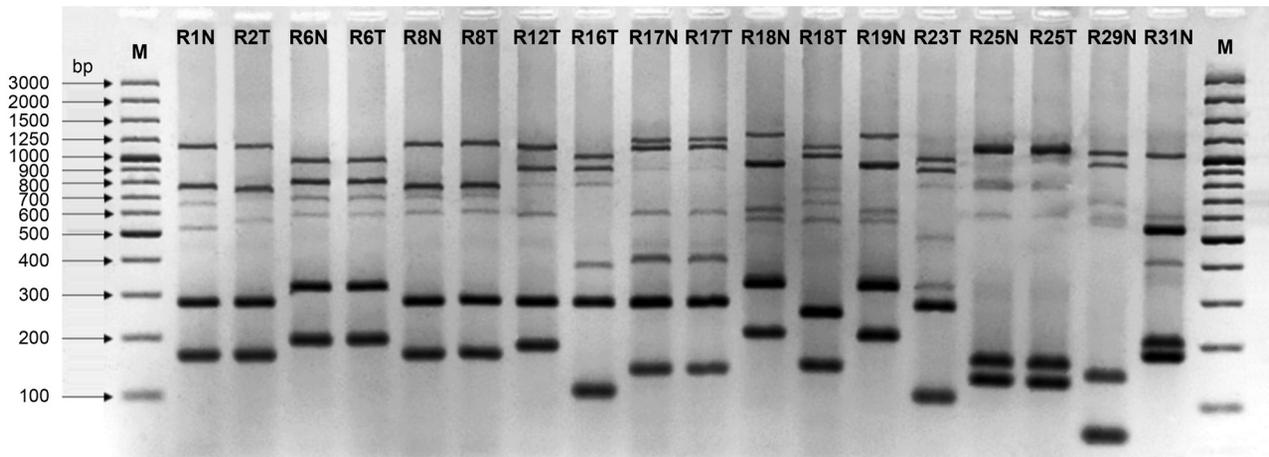


Fig. 2. MLVF of *S. aureus* strains isolated from the nose (N) and the throat (T) of thirteen residents living in a nursing home.

Note. Four residents were simultaneously colonized with the same genetic variant of *S. aureus* (strain R6N and R6T; strain R8N and R8T; strain R17N and R17T; strain R25N and R25T) and one resident was inhabited by two genetically different strains (R18N and R18T).
M – molecular marker (100–3000 bp).

> 35%). Comparing the subclusters concerning the size, the two largest (I and IV) both included nine banding patterns, and 13 and 11 *S. aureus* strains, respectively. The six smallest subclusters (III, V, VI, IX, X, and XI) each included between one and three banding patterns and two or three *S. aureus* strains.

Exemplary MLVF banding patterns for *S. aureus* strains are shown in Fig. 2. The number of MLVF bands depends on the variable expression of genes in the *sdr* region of the bacterial DNA. The expression of between one and three genes (*sdrC*, *sdrD* and/or *sdrE*) translates into a range between five and seven MLVF bands (Sabat et al. 2003). In our study, the six-band and seven-band profile were the most numerous, accounting for 56.0% and 36.4% of all isolates, respectively. The less common five-band profile captured only 7.6% of all isolates.

At the level of *S. aureus* isolates, we were able to group 26 isolates in even pairs based on perfect relatedness (one hundred percent homology). Without exception, each related pair of isolates was retrieved from the anterior nares and the throat of one individual, i.e., 13 individuals contributed one pair each. At the level of individuals, this meant that 27.6% (13/47) of the colonized individuals contributed a related pair of isolates. At the group level, i.e., the serendipitous detection of related isolates across the population, we were able to group four individuals in even couples, each couple sharing one bacterial strain. However, the anatomical location and pattern of co-colonization were inconsistent, rendering the similarities imperfect (for details, see subclusters I, VI and VIII in Fig. 1). These findings suggest that inter-participant transmission of *S. aureus* has occurred and might result in cross-infection – especially among immunocompromised individuals living in crowded conditions. Andersen et al. (2002) have

also reported that environmental factors, i.e., crowding and insufficient staffing, might increase the risk of *S. aureus* cross-infection.

Although our study provides valuable information about the genetic diversity, prevalence of colonization, and the occurrence of *S. aureus* transmission in a nursing home, it has several limitations. The MLVF method returns its most valid results when applied within a limited geographical area. As a result, making comparisons between studies from various countries or regions is difficult. Future research in the field of active microbial surveillance must include longitudinal aspects by subjecting the study populations to repeated sampling over extended periods. Understanding the temporal aspects of colonization might lead to new insights regarding the routes of bacterial transmission.

Author contributions

M.K. and A.G. collected the samples and performed the experiments. All authors conceived, designed the experiments and wrote the paper.

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Ethical approval

The study was approved by the Bioethics Committee at the Medical University of Lublin (KE 0254/59/2016). The samples were collected in January 2018 from residents and staff (n = 102) living and working in a nursing home (Lublin, Poland). Written informed consent was obtained from each individual before participation.

Conflict of interest

The authors do not report any financial or personal connections with other persons or organizations, which might negatively affect the contents of this publication and/or claim authorship rights to this publication.

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