Comparative profiling of agr locus, virulence, and biofilm-production genes of human and ovine non-aureus staphylococci

Elisa Azara1, Carla Maria Longheu1, Sonia Attene2, Silvana Sanna3, Marco Sale4, Maria Filippa Addis5 and Sebastiana Tola1*

Abstract

Background: In a collaboration between animal and human health care professionals, we assessed the genetic characteristics shared by non-aureus staphylococci (NAS) infecting humans and dairy ewes to investigate their relatedness in a region concentrating half of the total National sheep stock. We examined by PCR 125 ovine and 70 human NAS for biofilm production, pyrogenic toxins, adhesins, autolysins genes, and accessory gene regulator (agr) locus. The microtiter plate assay (MPA) was used for the phenotypic screening of biofilm production. Ovine NAS included S. epidermidis, S. chromogenes, S. haemolyticus, S. simulans, S. caprae, S. warneri, S. saprophyticus, S. intermedius, and S. muscae. Human NAS included S. haemolyticus, S. epidermidis, S. hominis, S. lugdunensis, S. capitis, S. warneri, S. xylosus, S. pasteuri, and S. saprophyticus subsp. bovis.

Results: Phenotypically, 41 (32.8%) ovine and 24 (34.3%) human isolates were characterized as biofilm producers. Of the ovine isolates, 12 were classified as biofilm-producing while the remaining 29 as weak biofilm-producing. All 24 human isolates were considered weak biofilm-producing. Few S. epidermidis isolates harbored the icaA/D genes coding for the polysaccharide intercellular adhesin (PIA), while the bhp, aap, and embp genes coding biofilm accumulation proteins were present in both non-producing and biofilm-producing isolates. Fifty-nine sheep NAS (all S. epidermidis, 1 S. chromogenes, and 1 S. haemolyticus) and 27 human NAS (all S. epidermidis and 1 S. warneri) were positive for the agr locus: agr-3se (57.8%) followed by agr-1se (36.8%) predominated in sheep, while agr-1se (65.4%), followed by agr-2se (34.6%) predominated in humans.

Concerning virulence genes, 40, 39.2, 47.2, 52.8, 80 and 43.2% of the sheep isolates carried atlE, aae, sdrF, sdrG, eno and epbS respectively, against 37.1, 42.8, 32.8, 60, 100 and 100% of human isolates. Enterotoxins and tsst were not detected.

Conclusions: Considerable variation in biofilm formation ability was observed among NAS isolates from ovine and human samples. S. epidermidis was the best biofilm producer with the highest prevalence of adhesin-encoding genes.

Keywords: Non-aureus staphylococci, Human, Ovine, Biofilm, Adhesins, Toxins, Quorum sensing
relies on dairy sheep farming, and controlling intramammary infections (IMI) is crucial.

Sheep mastitis prevalence is estimated to range from 5 to 30%, and several reports indicate that non-*aureus* staphylococci (NAS) are the most prevalent microorganisms causing subclinical disease in small ruminants [1–4]. Therefore, the exchange of colonizing and pathogenic microorganisms, with their antimicrobial-resistance and pathogenicity gene pools, can occur among sheep and farmers. NAS have recently gained attention as nosocomial agents causing frequent infections in debilitated or compromised patients, mainly associated with catheters and other indwelling medical devices [5]. NAS, and in particular *S. epidermidis*, can produce a multicellular biofilm that decreases the antibiotic concentration within the colony, promotes multiplication, and enhances the survival of invading bacteria [6]. Biofilm formation can be best assessed by the microtiter plate assay (MPA), as it produces a quantitative result by measuring the optical density of the stained biofilm [7, 8]. The main constituent of the NAS biofilm matrix is a linear 1,6-linked glycosidic polymer, also known as polysaccharide intercellular adhesin (PIA), synthesized by proteins encoded by the intercellular adhesion (ica) operon. Among the ica genes, icaA and icaD have an essential role in biofilm production [9]. The coexistence of both icaA/D genes leads to the full phenotypic expression of the capsular polysaccharide [9]. However, PIA-independent biofilms involving accumulation-associated protein (Aap), biofilm homologue protein (Bhp) and extracellular matrix-binding protein (Emb) have also been reported [10, 11].

Generally, NAS can produce several virulence factors that contribute collectively to colonization and invasion of host cells and tissues, as well as evasion of immune responses [12]. Virulence factors include the autolysins AltE and Aae [13, 14], and microbial surface components recognizing adhesive matrix molecules (MSCRAMMs) that mediate initial adhesion to different surfaces and promote colonization and serum protein binding [15]. The best known *S. epidermidis* MSCRAMMs are the fibrinogen-binding protein SdrG [16], and the collagen-keratin-binding protein SdrF [17, 18].

Furthermore, the production of various toxins can also contribute to NAS virulence [19], including staphylococcal enterotoxins (SEs) and toxic shock syndrome toxin 1 (TSST-1) [20]. Five serological types of SEs are typically known (SEA to SEE), but new types of SEs (SEG to SE1V) have also been identified and characterized [21, 22]. The quorum-sensing system (QS) agr, i.e. accessory gene regulator [23, 24] regulates biofilm formation, intercellular communication, and numerous virulence factors including toxins and autolysins. Three distinct genetic groups (types 1, 2, and 3) based on the *agr* locus polymorphism have been described in *S. epidermidis* [25], but data on the genetic polymorphisms of the *agr* locus in different species of NAS were not available in the scientific literature at the beginning of this investigation.

In this study, we compared the molecular characteristics of NAS isolated from the milk of sheep with mastitis and human clinical specimens with the following aims: 1) assess the biofilm production characteristics by phenotypic and genotypic methods, 2) carry out genotypic screening for a set of MSCRAMMs, autolysins, enterotoxins and tsst-1 genes and 3) investigate the *agr* locus and its genetic polymorphism.

**Results**

**Ovine NAS**

We analyzed a total of 125 isolates, including *S. epidermidis* (*n* = 57), *S. chromogenes* (*n* = 29), *S. haemolyticus* (*n* = 17), *S. simulans* (*n* = 8), *S. caprae* (*n* = 6), *S. warneri* (*n* = 5), *S. saprophyticus* (*n* = 1), *S. intermedius* (*n* = 1) and *S. muscae* (*n* = 1). Table 1 reports the isolates included in the study, while Supplementary Table S1 reports the primers used for PCR amplifications.

Table 2 summarizes the biofilm formation results. Out of 125 isolates examined, 41 (32.8%) were classified as biofilm producers; of these 29 (23.2%) were classified as weak biofilm-producers (WBP) while 12 (9.6%) as strong biofilm producers. Only one isolate harbored both icaA and icaD genes while two had only the icaA gene. On the other hand, 37 (29.6%), 22 (17.6%) and 63 (50.4%) isolates harbored the bhp, aap and emb genes, respectively (Table 2). For autolysin genes, 50 (40%) isolates were PCR positive for *altE* while 49 (39.2%) were positive for *aee*. Concerning adhesion factors (MSCRAMMs), 59 (47.2%), 66 (52.8%), 100 (80%) and 54 (43.2%) isolates harbored *sdrF*, *sdrG*, *eno* and *epbS*, respectively. All isolates were negative for *clfA* (Table 3). Regarding the *agr* type, 21 (16.8%) isolates belonged to *agr*-1 while 33 (26.4%) to *agr*-3. None of the isolates belonged to type 2 (Table 4). No amplification was obtained for the toxin genes analyzed.

*S. epidermidis*  
*S. epidermidis* was the most represented ovine NAS. Out of 57 isolates, 17 (30%) were classified as WBP and 11 (19%) as BP; only one non-BP harbored both icaA/D genes. On the other hand, 54 (94.7%), 30 (52.6%) and 20 (16%) isolates harbored *emb*, *bhp* and *aap*, respectively. Concerning autolysin genes, 48 (84.2%) were PCR-positive for *altE* and 46 (80.7%) for *aee*. MSCRAMM genes were found in high percentages: 98% for *eno*, 91% for *epbS*, 87.7% for *sdrG* and 78.9% for *sdrF*. No amplification was obtained for *clfA*, *fnbA*, *bhp*, *cna* and *fib* (Table 3). All *S. epidermidis* isolates were positive for the...
agr locus: 33 (57.8%) belonged to agr-3, while 21 (36.8%) to agr-1 (Table 4). Three isolates were non-typeable.

S. chromogenes, S. haemolyticus, and minor ovine NAS
S. chromogenes (n = 29) and S. haemolyticus (n = 17) were the most prevalent species in ovine milk samples after S. epidermidis. Table 2 shows that only 1 S. chromogenes was classified as BP while 7 (87.5%) S. simulans and 3 (50%) S. caprae were classified as WBP. The details are reported in Table 2.

Human NAS
The distribution of biofilm, autolysins, and MSCRAMMs genes in the 70 human NAS, including S. haemolyticus, S. epidermidis, S. lugdunensis, S. hominis, S. capitis, S. warneri, S. xylosus, S. pasteuri and S. saprophyticus subsp. bovis is shown in Tables 2 and 3.

**Table 1** Distribution of ovine and human NAS isolates according to the specimen origin

| Source        | S. epi | S. chr | S. hae | S. sim | S. cap | S. war | S. sap | S. int | S. mus |
|---------------|--------|--------|--------|--------|--------|--------|--------|--------|--------|
| Milk          | 57     | 29     | 17     | 8      | 6      | 5      | 1      | 1      | 1      |
| Human NAS     |        |        |        |        |        |        |        |        |        |
| Nasal swab    | 6      | 5      | –      | –      | –      | –      | –      | –      | –      |
| Blood         | 4      | 6      | –      | 1      | –      | –      | –      | –      | –      |
| Skin swab     | 2      | 3      | –      | –      | 2      | –      | 1      | –      | –      |
| Pus           | 1      | 3      | –      | –      | –      | –      | –      | –      | 1      |
| Peritoneal fluid | 2   | –      | 2      | –      | –      | –      | –      | –      | –      |
| Seminal fluid | 4      | –      | –      | –      | –      | –      | –      | –      | –      |
| Injury        | –      | 3      | 1      | –      | –      | 1      | –      | –      | –      |
| Ear swab      | 2      | –      | –      | –      | –      | 1      | –      | –      | –      |
| Oral swab     | 1      | 1      | –      | –      | –      | –      | –      | –      | 1      |
| Urine         | 2      | –      | –      | 1      | –      | –      | –      | –      | –      |
| CVC^b         | –      | 2      | –      | –      | –      | –      | –      | –      | –      |
| Ulcer swab    | –      | 1      | –      | –      | –      | –      | –      | –      | –      |
| FVC^b         | –      | 1      | –      | –      | –      | –      | –      | –      | –      |
| Peritoneal swab| 1   | –      | –      | –      | –      | –      | –      | –      | –      |
| Vaginal swab  | –      | –      | 1      | –      | –      | –      | –      | –      | –      |
| Glans swab    | 1      | –      | –      | –      | –      | –      | –      | –      | –      |
| Pleural fluid | –      | –      | –      | 1      | –      | –      | –      | –      | –      |
| Fluid drainage| –      | –      | –      | –      | –      | 1      | –      | –      | –      |
| B.L. fluid^b  | 1      | –      | –      | –      | –      | –      | –      | –      | –      |
| N.P. aspirate^b| –    | –      | –      | –      | 1      | –      | –      | –      | –      |
| Bile          | 1      | –      | –      | –      | –      | –      | –      | –      | –      |
| Biopsy        | –      | 1      | –      | –      | –      | –      | –      | –      | –      |
| Prosthesis    | –      | –      | 1      | –      | –      | –      | –      | –      | –      |
| Total         | 28     | 26     | 4      | 4      | 4      | 3      | 2      | 1      | 1      |

^a Isolate abbreviations: S. hae, S. haemolyticus; S. epi, S. epidermidis; S. chr, S. chromogenes; S. sim, S. simulans; S. lag, S. lugdunensis; S. hom, S. hominis; S. cpr, S. caprae; S. cap, S. capitis; S. war, S. warneri; S. xyl, S. xylosus; S. pas, S. pasteuri; S. int, S. intermedius; S. sap, S. saprophyticus subsp. Bovis; S. mus, S. muscae

^b Specimen abbreviations: C.V.C central venous catheter, F.V.C femoral venous catheter, B.L. fluid bronchoalveolar lavage fluid, N.P. aspirate, nasopharyngeal aspirate

S. haemolyticus
Out of 28 S. haemolyticus isolates examined by MPA, only 1 (3.6%) was classified as WBP. None of the S. haemolyticus isolates harbored icaA/D, bhp and embp. On the contrary, the WBP (from blood) and other 3 non-BP (from nasal swab, blood, and glans swab) isolates possessed the aap gene. No amplification was observed for atlE, aae, sdrF/G, clfA/B, fnbA/B, bhp, cna, fib, epbS, agr locus (Table 3), and toxin genes. However, all isolates harbored the eno gene.

S. epidermidis
Among the 26 S. epidermidis isolates, 9 (34.6%) were classified as WBP; 4 of them (2 from catheter and 2 from blood) harbored both icaA and icaD whereas the remaining 5 isolates were icaA/D negative. The other 5 icaA/D positive isolates, classified as non-BP, were also positive for the other biofilm genes analyzed. Out of 17 non-BP
and icaA/D negative isolates, 3 (2 from pus and 1 from oral swab) harbored aap and embp while 1 (from skin swab) harbored all bhp, aap, and embp genes. Overall, 16 (61.5%), 25 (96.1%) and 23 (88.5%) isolates carried the bhp, aap, and embp genes, respectively (Table 2). Data on the prevalence of autolysins and MSCRAMM genes by PCR are shown in Table 3. Concerning autolysin genes, all isolates were PCR positive for aae and almost all (25/26 = 96.1%) were positive for altE. Regarding adhesion factors, all S. epidermidis isolates were positive for sdrG, eno, and epbS, while 21/26 (80.8%) for sdrF. In four of them (2 from pus, 1 from biopsy and 1 from skin swab), a PCR product smaller than the expected size was observed. Sequence analysis of these amplicons showed the absence of an 84 bp fragment. In contrast, no amplification was observed for clfA/B, fnbA/B, bhp, cna, and fib. Determination of the agr type was performed in all S. epidermidis isolates: 17 (65.4%) belonged to agr-1se whilst 9 (34.6%) to agr-2se (Table 4). Among the 17 agr-1se isolates, only 3 (1 from femoral venous catheter, 1 from nasal swab and 1 from pus) carried simultaneously icaA/D, bhp, aap, embp, atlE, aae, sdrG, eno, and epbS, associated with biofilm formation. Of these, 1 was WBP and two non-BP. Among the 9 agr-2se isolates, only 1 (from blood) possessed these genes and it was a WBP isolate. Regarding the pyrogenic toxin genes, amplification was not observed in the S. epidermidis isolates or the remaining staphylococci.

Minor human NAS
Out of the 4 S. lugdunensis isolates examined, 3 (2 from peritoneal fluid and 1 from vaginal swab) were considered as WBP. However, they harbored only sdrG and eno.

Table 2 Phenotypic characterisation of biofilm production by MPA and genotypic detection by PCR of ica, bhp, aap and embp genes from 125 ovine and 70 human NAS isolates

| Ovine Isolates (n) | Biofilm production (MPA) | Biofilm genes | Proteinaceous factors |
|-------------------|--------------------------|---------------|----------------------|
|                   | WBP<sup>a</sup> | B.P<sup>b</sup> | icaA | icaD | icaA/D | bhp | aap | embp |
|                   | n | % | n | % | n | % | n | % | n | % | n | % |
| S. epi (57)       | 17 | 30 | 11 | 19 | 1 | 1.7 | 1 | 1.7 | 1 | 1.7 | 30 | 52.6 | 20 | 16 | 54 | 94.7 |
| S. chr (29)       | 0 | 0 | 1 | 3.4 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 4 | 13.8 | 0 | 0 | 8 | 27.5 |
| S. hae (17)       | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. sim (8)        | 7 | 87.5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. cpr (6)        | 3 | 50 | 0 | 0 | 1 | 16.6 | 0 | 0 | 0 | 0 | 0 | 1 | 16.6 | 0 | 0 | 0 | 0 |
| S. war (5)        | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 2 | 40 | 0 | 0 | 0 | 0 |
| S. sap (1)        | 1 | 100 | 0 | 0 | 1 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. int (1)        | 1 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. mus (1)        | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Total (125)       | 29 | 23.2 | 12 | 9.6 | 3 | 2.4 | 1 | 0.8 | 1 | 0.8 | 37 | 29.6 | 22 | 17.6 | 63 | 50.4 |

| Human Isolates (n) | Biofilm production (MPA) | Biofilm genes | Proteinaceous factors |
|---------------------|--------------------------|---------------|----------------------|
|                     | WBP<sup>a</sup> | B.P<sup>b</sup> | icaA | icaD | icaA/D | bhp | aap | embp |
|                     | n | % | n | % | n | % | n | % | n | % | n | % |
| S. hae (28)         | 1 | 3.6 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. epi (26)         | 9 | 34.6 | 0 | 0 | 10 | 38.5 | 10 | 38.5 | 10 | 38.5 | 16 | 61.5 | 25 | 96.1 | 23 | 88.5 |
| S. lug (4)          | 3 | 75 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. hom (4)          | 4 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. cap (3)          | 3 | 100 | 0 | 0 | 3 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. war (2)          | 2 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 50 |
| S. xyl (1)          | 1 | 100 | 0 | 0 | 1 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. pas (1)          | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| S. sap (1)          | 1 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Total (70)          | 24 | 34.3 | 0 | 0 | 14 | 20 | 10 | 14.3 | 10 | 14.3 | 16 | 22.8 | 34 | 48.6 | 24 | 34.3 |

Isolate abbreviations: S. hae, S. haemolyticus; S. epi, S. epidermidis; S. chr, S. chromogenes; S. sim, S. simulans; S. lug, S. lugdunensis; S. hom, S. hominis; S. cpr, S. caprae; S. cap, S. capitis; S. war, S. warrenii; S. xyl, S. xylosus; S. pas, S. pasteuri; S.int, S. intermedius; S. sap, S. saprophyticus subsp. bovis; S. mus, S. muscae

<sup>a</sup> WBP weak biofilm-producing isolate

<sup>b</sup> BP biofilm-producing isolate

Out of the 4 S. lugdunensis isolates examined, 3 (2 from peritoneal fluid and 1 from vaginal swab) were considered as WBP. However, they harbored only sdrG and eno.
Table 3  Results of testing 125 ovine and 70 human NAS isolates for autolysins and MSCRAMMS genes by PCR

| Isolate Abbreviations | OVINE Isolates (n) | Autolysins | MSCRAMMS |
|-----------------------|-------------------|-------------|-----------|
|                       |                   | atlE | aae | sdrF | sdrG | clfA | clfB | fnbA | fnbB | bbp | cna | flb | eno | epbS |
|                       |                   | n   | %   | n   | %   | n   | %   | n   | %   | n   | %   | n   | %   | n   | %   | n   | %   | n   | %   | n   | %   |
| S. epi (57)           |                   | 48  | 84.2| 46  | 80.7| 45  | 78.9| 50  | 87.7| 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 56  | 98  | 52  | 91  |
| S. chr (29)           |                   | 1   | 3.4 | 0   | 0   | 9   | 31  | 13  | 22.8| 0   | 0   | 2   | 7   | 7   | 1   | 3.4 | 0   | 0   | 7   | 7   | 16  | 55  | 2   | 7   |
| S. hom (17)           |                   | 1   | 5.8 | 1   | 5.8 | 1   | 5.8 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 16  | 88  | 0   | 0   |
| S. sim (8)            |                   | 0   | 0   | 1   | 12.5| 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   |
| S. cap (6)            |                   | 0   | 0   | 0   | 0   | 1   | 16.6| 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   |
| S. war (5)            |                   | 0   | 0   | 1   | 20  | 3   | 60  | 2   | 40  | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 4   | 80  | 0   | 0   |
| S. int (1)            |                   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   |
| S. mus (1)            |                   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   |
| Total (125)           |                   | 50  | 40  | 49  | 39.2| 59  | 47.2| 66  | 52.8| 0   | 0   | 2   | 1.6 | 2   | 1.6 | 1   | 0.8 | 1   | 0.8 | 2   | 1.6 | 2   | 1.6 | 100 | 80  | 54  | 43.2|
| HUMAN Isolates (n)    |                   |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
|                       |                   |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| S. epi (26)           |                   | 25  | 96.1| 26  | 100 | 21  | 80.8| 26  | 100 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 26  | 100 | 26  | 100 |
| S. lug (4)            |                   | 0   | 0   | 0   | 0   | 0   | 0   | 4   | 100 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   |
| S. cap (3)            |                   | 0   | 0   | 0   | 0   | 0   | 0   | 3   | 100 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 3   |
| S. war (2)            |                   | 1   | 50  | 1   | 50  | 1   | 50  | 2   | 100 | 1   | 50  | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 2   |
| S. xyl (1)            |                   | 0   | 0   | 1   | 100 | 1   | 100 | 1   | 100 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 1   | 100 | 0   | 0   |
| S. pas (1)            |                   | 0   | 0   | 1   | 100 | 0   | 0   | 1   | 100 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 1   | 100 | 0   | 0   |
| S. sap (1)            |                   | 0   | 0   | 1   | 100 | 0   | 0   | 1   | 100 | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 1   | 100 | 0   | 0   |
| Total (70)            |                   | 26  | 37.1| 30  | 42.8| 23  | 32.8| 42  | 60  | 1   | 50  | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 0   | 70  | 100 | 26  | 100 |

Isolate abbreviations: S. hae, S. haemolyticus; S. epi, S. epidermidis; S. chr, S. chromogenes; S. sim, S. simulans; S. lug, S. lugdunensis; S. hom, S. hominis; S. capr, S. caprae; S. cap, S. capitis; S. war, S. warneri; S. xyl, S. xylolus; S. pas, S. pasteuri; S. int, S. intermedius; S. sap, S. saprophyticus subsp. bovis; S. mus, S. muscae
All 4 S. hominis isolates were phenotypically WBP but were PCR-positivity only for aap, sdrG, and eno. All 3 WBP S. capitis isolates were PCR-positive for icaA, sdrG, and eno. Tables 3 and 4 report the PCR results for S. warneri, S. xylosus, S. pasteuri, and S. saprophyticus subsp. bovis. Of note, one WPB S. warneri isolate (from fluid drainage) was negative for icaA/D and bhp genes but was positive for the aap, embp, atlE, aae, sdrG/F, clfA, eno and agr genes. However, we were not able to type the agr locus.

**Discussion**

We established a collaboration between animal and human health care professionals aimed at understanding if non-aureus staphylococci (NAS) responsible for human diseases share genetic similarities with those circulating in sheep, in consideration of the high number of dairy sheep farmed in the island and of the prominent role of these bacteria as mastitis agents.

A total of 195 NAS isolates, 125 from ovine mastitis and 70 from human clinical specimens, were analyzed for biofilm production and presence of autolysins, pyrogenic toxins, and MSCRAMM genes. We also typed agr alleles by PCR because the quorum sensing system regulates many virulence determinants involved in staphylococcal infections, including autolysins, adhesins, and toxins [19]. In sheep, the primary NAS detected were S. epidermidis followed by S. chromogenes and S. haemolyticus. At the same time, in humans we found primarily S. haemolyticus and S. epidermidis, followed by S. lugdunensis, S. hominis, S. capitis, S. warneri, S. xylosus, S. pasteuri, S. caprae, S. intermedius, S. saprophyticus subsp. bovis, S. muscae, S. muscae.
the most frequently isolated species from ovine mastitis and human clinical specimens [1, 26, 27].

Overall, 65 NAS were able to form biofilm in vitro; however, the percentage of biofilm producers in sheep isolates was slightly lower than in human isolates. Moreover, we found a correlation between biofilm production and ica operon presence only in S. epidermidis isolates, 4 human and 1 ovine. Some authors proposed to use this correlation as a pathogenesis marker to distinguish invasive from commensal isolates [28, 29]. However, we and others [11, 30, 31] demonstrated that PCR positivity for icaA/icaD genes can also be found in non-biofilm producers. Since the correlation between biofilm production and positivity for ica, bhp, aap, and embp genes is not clearly defined, we suggest considering all isolates that possess such genes as potentially invasive. In this work, only one S. epidermidis with these characteristics was isolated from ovine mastitis while the other 4 derived from catheters and blood. Noteworthy, a high positivity for the genes encoding the bifunctional adhesins/autolysins AtLE and Aae was found in both animal and human S. epidermidis isolates [3, 5]. In addition to bacteriolytic activity, AtLE and Aae act as adhesins by binding noncovalently to vitronectin and by causing the release of extracellular DNA (eDNA), a critical adherence/aggregation factor in biofilm formation [32]. The presence of atLE and aae in S. epidermidis was accompanied by a high prevalence of embp, sdrG, sdrF, eno and epbS, all genes that mediate adherence to substrates containing fibronectin, fibrinogen, collagen, laminin and elastin, respectively [4, 33–35]. The ability of S. epidermidis to bind these substrates might represent a relevant mechanism by which it can adhere to and colonize different host sites. The eno gene was the only gene found in all NAS analyzed, except for S. simulans. In human NAS, the prevalence is 100%. Therefore, the ability of NAS to bind laminin, a major component of basal membrane of the vasculature, might play a possible role in to tissue invasion and blood dissemination.

The agr locus is a regulatory system that responds to host and environmental stimuli and controls the production of many virulence factors [24]. In S. epidermidis, three distinct agr groups have been recognized [25]. Li et al. [36] have linked the genetic polymorphism of the agr locus to pathogenicity; group-1se was associated with pathogenicity, while healthy people mainly carried group-2se. In our human S. epidermidis isolates, agr-1se was predominant (n = 17), followed by agr-2se (n = 9). It is interesting to notice that almost all isolates possessing ica genes belonged to agr-1se. This may suggest a correlation of these virulence genes with a specific agr locus. However, other 8 icaA−/D− isolates were present in the group-1se. The feature shared by all 17 isolates belonging to this group was the PCR positivity for the attE, aae, sdrG, eno and epbS genes. On the other hand, among the 9 isolates grouped in the agr-2se, 1 (from blood) was icaA+/D−, while the remaining ones were ica-negative. The common denominator of these 9 isolates was the PCR positivity for aap, aae, sdrG, embp, eno, and epbS. These findings suggest that the relationship between agr groups and S. epidermidis pathogenicity will require further investigation. As observed in our previous study [30], agr-3se (n = 33) was predominant among ovine S. epidermidis isolates followed by agr-1se (n = 21). These results may suggest a possible transmission of S. epidermidis isolates from the milkers to the ewes.

Unlike S. chromogenes, S. haemolyticus, S. warneri and S. pasteuri from ovine mastitis and S. pasteuri from human specimens, the other NAS were classified as WBP by the microplate adhesion technique but did not harbor icaA/D. According to Fredheim et al. [37], S. haemolyticus mainly produces a PIA-independent biofilm. However, we detected only the aap (2/17) and embp (1/17) genes in the present study by PCR. Only 4 human S. haemolyticus isolates possessed the aap gene coding a protein that mediates biofilm formation in strains lacking the ica genes [16]. Our data suggest that ica and bhp genes do not contribute significantly to S. haemolyticus biofilms’ protein components.

In S. aureus and in many other bacteria, toxins are critical contributors to aggressive virulence, even though S. epidermidis is not generally accepted as an enterotoxin producer [38, 39]. Based on our findings, the primary enterotoxin genes (sea, seb, sec, sed and see) and the tsst-1 gene were absent in all ovine and human NAS analyzed. On the contrary, Pedroso et al. [16] and Da Cunha et al. [40] detected high percentages of sea and sec genes in coagulase-negative staphylococci from hospitals of Brazil; also, Giormezis et al. [39] found a higher number of isolates positive for tsst among NAS from hospitals in Greece.

Conclusion

In conclusion, we detected intercellular adhesion genes (icaAB) and other genes related to biofilm formation only in S. epidermidis, although we found icaA in ovine S. caprae and S. saprophyticus, and in human S. capitis and S. xylosus. The remaining isolates carried few virulence determinants. The ability to form biofilm observed in NAS isolates, especially S. epidermidis, might constitute a significant virulence factor facilitating colonization, infection, diffusion, and resistance.
Methods

Isolate collection

Ovine isolates: A total of 125 NAS were isolated from sheep milk samples that routinely arrive at the Istituto Zooprofilattico Sperimentale della Sardegna. Milk samples, collected from farms with mastitis problems in different provinces of Sardinia (Italy), were analyzed over 9 months (April-December 2017) (Table 1). The geographic distribution of these isolates is reported in Supplementary Fig. S1. Isolates were identified as: S. epidermidis (n = 57), S. chromogenes (n = 29), S. haemolyticus (n = 17), S. simulans (n = 8), S. caprae (n = 6), S. warneri (n = 5), S. saprophyticus (n = 4), S. intermedius (n = 1), and S. muscae (n = 1), by means of PCR-RFLP [27].

Human isolates: During the same period, 70 NAS isolates were collected from different clinical specimens at the microbiology laboratories of three major Sardinia hospitals. Isolates were anonymized without patient identifiers. Isolates and their origin are summarized in Table 1; 90% of the human NAS were recovered from hospitalized patients in intensive care unit, hematology, and orthopedics. The 70 isolates were identified by PCR-RFLP as S. haemolyticus (n = 28), S. epidermidis (n = 26), S. lugdunensis (n = 4), S. hominis (n = 4), S. capitis (n = 3), S. warneri (n = 2), S. xylosus (n = 1), S. pasteurii (n = 1), and S. saprophyticus subspp. bovis (n = 1) [27].

Statements of owner consent or patient consent were not required in this case since personal or sensitive data never accompanied samples. All Isolates were anonymized regarding the originating animal, flock, or patient, and were processed for phenotypic and molecular analyses without any original information linked to them.

Phenotypic evaluation of biofilm production by the microtiter plate assay (MPA)

All 195 isolates were tested using the MPA technique, described by Vasileiou et al. [8] with some modifications. Briefly, a colony of each isolate was inoculated into a tube containing 1 mL Tryptone Soy Broth (TSB, Oxoid, Basingstoke, UK) for 16 h at 37°C. Overnight culture was diluted 1:40 with TSB containing 0.25% glucose, and 200 μL per well were seeded in a sterile 96-well flat-bottomed microplate (Thermo Fisher, Rodano, IT) at 37°C for 24 h. After three washes in PBS pH 7.4, the microplate was dried at 45°C for 20 min, and wells were then stained with 1% crystal violet for 15 min at room temperature. After three washes with distilled water and subsequent drying at 45°C for 20 min, 200 μL of 33% acetic acid were added to each well. Biofilm growth was measured at 630 nm in a microplate spectrophotometer (Multiskan GO, Thermo Fisher). Unincoculated wells containing TBS with glucose served as blanks. In each microplate, S. epidermidis ATCC 35984 and S. epidermidis ATCC 12228 were included as the positive and negative controls, respectively. Each isolate and both controls were tested in triplicate, and the assay was repeated two times at different dates. Isolates were classified into three categories based upon the median OD of isolates and positive and negative controls: biofilm-producing (OD isolate ≥ OD of the positive control), weak biofilm-producing (OD negative control < OD isolate < OD positive control) and non biofilm-producing (OD isolate ≤ negative control).

Detection of biofilm, autolysins, MSCRAMMs and pyrogenic toxins genes

Genomic DNA was extracted from all 195 NAS isolates and Reference Strains (RS) according to Onni et al. [41]. Single-tube PCRs were performed for detecting genes related to biofilm production (icaA/D, bhp, aap, embp) [30, 31, 42, 43], autolysins (atlE and aae) [44, 45], MSCRAMMs (encoding clumping factor-clfA/B, fibronectin-binding protein-fnbA/B, encoding bone sialoprotein-binding protein-bbp, collagen-binding protein-cna, fibrinogen-binding protein-fib, laminin-binding protein-enol, elastin-binding protein-ebpS, and serine-aspartate repeat protein-repeat protein-sdrF/G) [19, 30, 46–49] and pyrogenic toxins (sea, seb, sec, sed, see and tsst-1) [50–52]. Primer sets are reported in Supplementary Table S1. PCR tests were carried out in a GeneAmp9700 DNA thermal cycler (Applied Biosystems, now Thermo Fisher Scientific, Waltham, MA, USA). The following RS were used as positive controls: S. epidermidis ATCC 35984 (icaA/D, bhp, aap, embp, atlE, aae, sdrF, sdrG), S. aureus ATCC 25923 (clfA/B, bhp, epbs), S. aureus ATCC 35591 (fnbA/B, cna, fib, enol), S. aureus ATCC 13565 (sea), S. aureus ATCC 14458 (seb), S. aureus ATCC19095 (sec), S. aureus ATCC 23235 (sed), S. aureus ATCC 27664 (see) and S. aureus ATCC 33586 (tsst-1).

Typing of agr alleles

A 200 bp conserved region of the agr operon was amplified as described previously [19]. For isolate typing, we targeted agr-1a to agr-3s sequences [53]. As controls, we used S. epidermidis ATCC 35984 (agr-1α), S. epidermidis isolate 1037 (agr-2s) and S. epidermidis isolate 43027 (agr-3s). Supplementary Table S1 reports primer sets and related references.
Abbreviations
CRA : Congo Red Agar; MSCRAMMs: Microbial surface components recognizing adhesive matrix molecules; NAS: Non-aureus staphylococci; QS: Quorum-sensing system.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s12917-022-03257-w.

Additional file 1: Supplementary Table S1. Oligonucleotide primers used for the detection of genes related to biofilm production (icaA/B), hla, fnbA, fnbB, ebp, cap, msr, sar, seb, sec, sed, see and tst-1) and for agr typing.

Additional file 2: Supplementary Figure S1. Map of Sardinia showing the location of all 125 non-aureus staphylococci isolates in every municipality. No copyright permission was required.

Acknowledgements
Not applicable.

Authors’ contributions
EA: carried out isolation and identification of ovine NAS, performed the experiments; analyzed the data. CML: carried out isolation and identification of ovine NAS, performed the experiments; SA, SS, and MS: carried out isolation and identification of human NAS; MFA: interpreted the data; drafted and revised the manuscript; ST: Conceived the study, analyzed, and interpreted the data; performed the experiments; SA, SS, and MS: carried out isolation and identification of ovine NAS; JMC: interpreted the data; drafted and revised the manuscript; CT: supervised the project. All authors read, edited, and approved the final version of the manuscript.

Funding
This work was supported by Istituto Zooprofilattico Sperimentale della Sardegna, Italy. The funding body have no role in the design of the study, writing the manuscript and the collection, analysis and interpretation of data.

Availability of data and materials
All data supporting these research findings are included within the manuscript. The databases are available from the corresponding author upon request.

Declarations
Ethics approval and consent to participate
Full name of all participating hospitals: Ospedale “San Francesco”, Ospedale “A. Segni” and Azienda Ospedaliera Universitaria.

Human material:
Following diagnostic routine procedures carried out at each hospital, isolates obtained from biological human material, were stored for future analyses. All samples were fully anonymised. We understand that each hospital obtained approval and patient consent. For our research study, special permissions were granted by each hospital to use these isolates for our analyses. In accordance with national regulations and the institutional rules for Good Scientific Practice, our study protocol did not require separate approval by an ethical committee.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no conflicts of interest associated with this study.

Author details
1 Istituto Zooprofilattico Sperimentale della Sardegna “G. Pegreffi”; Via Vienna 2, 07100 Sassari, Italy. 2 Ospedale “San Francesco”, 08100 Nuoro, Italy. 3 Azienda Ospedaliera Universitaria, 07100 Sassari, Italy. 4 Ospedale “A. Segni”, 07014 Ozieri, Italy. 5 Dipartimento di Medicina Veterinaria, Università degli Studi di Milano, 26900 Lodi, Italy.

Received: 3 November 2020 Accepted: 20 April 2022

References
1. Marogna G, Rollesu S, Lolli S, Tola S, Leori G. Clinical findings in sheep farms affected by recurrent bacterial mastitis. Small Rumin Res. 2010;88:119–25. https://doi.org/10.1016/j.smallrumres.2010.11.019.
2. Onni T, Vidilli A, Bandino E, Marogna G, Schianchi S, Tola S. Identification of coagulase-negative staphylococci isolated from caprine milk samples by PCR-RFLP of groEL gene. Small Rumin Res. 2012;104:185–90. https://doi.org/10.1016/j.smallrumres.2011.10.004.
3. Martins KB, Faccioli PY, Bonesso MP, Fernandes S, Oliveira AA, Dantas A, et al. Characteristics of resistance and virulence factors in different species of coagulase-negative staphylococci isolated from milk of healthy sheep and animals with subclinical mastitis. J Dairy Sci. 2017;100:2184–95. https://doi.org/10.3168/jds.2016-11583.
4. Vanderhaeghen W, Pieters P, Leroy F, Van Coillie E, Haesebrouck F, De Vliegher S. Identification, typing, ecology and epidemiology of coagulase negative staphylococci associated with ruminants. Vet J. 2015;203:44–51. https://doi.org/10.1016/jivet.2014.11.001.
5. Von Eiff C, Peters G, Heilmann C. Pathogenesis of infections due to coagulase-negative staphylococci. Lancet Infect Dis. 2002;2:677–85. https://doi.org/10.1016/S1473-3099(02)00438-3.
6. Mack D, Davies AP, Harris LG, Rohde H, Horstktte MA, Knobloch JK-M. Microbial interactions in Staphylococcus epidermidis biofilms. Anal Bioanal Chem. 2007;387:399–408. https://doi.org/10.1007/s00216-006-0745-2.
7. Stepanovic S, Yukovic D, Dakic I, Savic B, Svbacic-Vlahovic M. A modified microtiter-plate test for quantification of staphylococcal biofilm formation. J Microbiol Methods. 2000;40:175–9. https://doi.org/10.1016/S0167-7012(00)00123-6.
8. Vasilieou NG, Chatzopoulos DC, Gougooula D, Sarrou S, Katsafadou AI, Spyrou V, et al. Slime-producing staphylococci as causal agents of sub-clinical mastitis in sheep. Vet Microbiol. 2016;224:95–9. https://doi.org/10.1016/j.vetmic.2018.08.022.
9. Arcioli CR, Campoccia D, Gamberini S, Cervellati M, Donati E, Montanaro L. Detection of slime production by means of an optimised Congo red agar plate test based on a colourimetric scale in Staphylococcus epidermidis clinical isolates genotyped for ica locus. Biomaterials. 2002;23:4233–40. https://doi.org/10.1016/S0142-9612(02)00171-0.
10. Kogan G, Gavish A, Chagin N, Chokr A, Jabbour S. Biofilms of clinical strains of Staphylococcus do not contain polysaccharide intercellular adherant. FEMS Microbiol Lett. 2006;255:11–6. https://doi.org/10.1111/j.1574-6968.2005.00043.x.
11. Rohde H, Burandt EC, Siemssen N, Frommelt L, Burdelski C, Wurster S, et al. Polysaccharide intercellular adhesion or protein factors in biofilm accumulation of Staphylococcus epidermidis and Staphylococcus aureus isolated from prosthetic hip and knee joint infections. Biomaterials. 2007;28:1711–20. https://doi.org/10.1016/j.biomaterials.2006.11.046.
12. Fey PD, Olson ME. Current concepts in biofilm formation of Staphylococcus epidermidis. Future Microbiol. 2010;5:917–33. https://doi.org/10.2217/ fbm.10.56.
13. Heilmann C, Hussain M, Peters G, Götz F. Evidence for autolysin-mediated primary attachment of Staphylococcus epidermidis to a polystyrene surface. Mol Microbiol. 1997;24:1013–24. https://doi.org/10.1046/j.1365-2958.1997.4101774.x.
14. Heilmann C, Thrumm G, Chhatwal GS, Hartleib J, Uekötter A, Peters G. Identification and characterization of a novel autolysin (Aae) with adhesive properties from Staphylococcus epidermidis. Microbiology. 2003;149:2769–78. https://doi.org/10.1099/mic.0.26527-0.
15. Otto M. Molecular basis of Staphylococcus epidermidis infections. Semin Immunopathol. 2012;34:201–14. https://doi.org/10.1007/s00281-011-0296-2.
16. Pei L, Flock JI. Lack of fib, the gene for a fibrinogen-binding protein from Staphylococcus epidermidis, reduces its adherence to fibrinogen coated surfaces. Microb Pathog. 2001;31:185–93. https://doi.org/10.1006/impat.2001.0462.
33. Christner M, Franke GC, Schommer NN, Wondt U, Wegert K, Pehle P, et al. The giant extracellular matrix-binding protein of Staphylococcus epidermidis mediates biofilm accumulation and attachment to fibronectin. Mol Microbiol. 2010;75:187–207. https://doi.org/10.1111/j.1365-319X.2009.06981.x.

34. Caniero CRW, Postol E, Nizomor R, Reis LFL, Brentani RR. Identification of enolase as a laminin-binding protein on the surface of Staphylococcus aureus. Microb Infect. 2004;6:604–8. https://doi.org/10.1016/j.micinf.2004.02.003.

35. Downer R, Roche F, Park PW, Mecham RP, Foster TJ. The elastin-binding protein of Staphylococcus aureus (EbpS) is expressed at the cell surface as an integral membrane protein and not as a cell wall-associated protein. J Biol Chem. 2002;277:243–50. https://doi.org/10.1074/jbc.M107621200.

36. Li M, Guan M, Jiang XF, Yuan FY, Xu M, Zhang WZ, et al. Genetic polymorphism of the accessory gene regulator (agr) locus in Staphylococcus epidermidis and its association with pathogenicity. J Med Microbiol. 2004;53:545–9. https://doi.org/10.1099/jmm.0.035406-0.

37. Feddehem EGA, Klingenberg C, Rohde H, Frankenberger S, Gaustad P, Flægstad T, et al. Biofilm formation by Staphylococcus haemolyticus. J Clin Microbiol. 2009;47:1172–80. https://doi.org/10.1128/JCM.01091-08.

38. Otto M. Staphylococcus epidermidis - the "accidental" pathogen. Rev Med Microbiol. 2009;7:555–67. https://doi.org/10.1038/rmmicro2182.

39. Giormeizis N, Kolonitsiou F, Foka A, Drougka E, Liakopoulos A, Makri A, et al. Coagulase-negative staphylococcal bloodstream and prosthetic-device-associated infections: the role of biofilm formation and distribution of adhesin and toxin genes. J Med Microbiol. 2016;65:1500–10. https://doi.org/10.1099/jmm.0.05259-0.

40. Da Cunha MDRS, Calosilari RAQ, Araujo JP. Detection of enterotoxin and toxic shock syndrome toxin 1 genes in Staphylococcus, with emphasis on coagulase-negative staphylococcal. Microbiol Immunol. 2007;51:381–90. https://doi.org/10.1111/j.1348-0421.2007.tb03925.x.

41. Onn T, Sanna G, Larsen J, Tola S. Antimicrobial susceptibilities and population structure of Staphylococcus epidermidis associated with ovine mastitis. Vet Microbiol. 2011;148:45–50. https://doi.org/10.1016/j.vetmic.2010.07.024.

42. Vandencaestelee SJ, Peetersmans WE, Merckx RR, Rijnders BJA, Van Eldere J. Reliability of the Ica, aap and atg genes in the discrimination between invasive, colonizing and contaminating Staphylococcus epidermidis isolates in the diagnosis of catheter-related infections. Clin Microbiol Infect. 2009;15:114–9. https://doi.org/10.1016/j.cmi.2009.05.044.

43. Gu J, Li H, Li M, Vuong C, Otto M, Wen Y, et al. Bacterial insertion sequence IS256 as a potential molecular marker to discriminate invasive strains from commensal strains of Staphylococcus epidermidis. J Hosp Infect. 2005;61:342–8. https://doi.org/10.1016/j.jhin.2005.04.017.

44. Maretra T, Hermansen L, Holck AL, Sidhu MS, Rudi K, Langvird S. Biofilm formation and the presence of the intercellular adhesion locus ica among staphylococci from food and food processing environments. Appl Environ Microbiol. 2003;69:5648–55. https://doi.org/10.1128/AEM.69.9.5648-5655.2003.

45. Lou Q, Zhu T, Hu J, Ben H, Yang J, Yu F, et al. Role of the SaeRS two-component regulatory system in Staphylococcus epidermidis autoaggregation and biofilm formation. BMC Microbiol. 2011;11:146. https://doi.org/10.1186/1471-2180-11-146.

46. Stephan R, Annemüller C, Hassan AA, Lämmler C. Characterization of enterotoxigenic Staphylococcus aureus strains isolated from bovine mastitis in north-East Switzerland. Vet Microbiol. 2001;78:373–82. https://doi.org/10.1016/S0378-1135(00)00134-2.

47. Peacock SJ, Moore CE, Justice A, Kanzanou M, Story L, Mackie K, et al. Virulent combinations of adhesin and toxin genes in natural populations of Staphylococcus aureus. Infect Immun. 2002;70:4987–96. https://doi.org/10.1128/IAI.70.9.4987-4996.2002.

48. Vancraeynest D, Hermansen K, Hassebroeck F. Genotypic and phenotypic screening of high and low virulence Staphylococcus aureus isolates from rabbits for biofilm formation and MSCRAMMs. Vet Microbiol. 2004;103:241–7. https://doi.org/10.1016/j.vetmic.2004.09.002.

49. Tristan A, Ying L, Bes M, Etienne J, Vandenesch F, Lina G. Use of multiplex PCR to identify Staphylococcus aureus adhesins involving in human hematogenous infection. J Clin Microbiol. 2003;41:4465–7.

50. Monday SR, Bohach GA. Use of multiplex PCR to detect classical and newly described pyrogenic toxin genes in staphylococcal isolates. J Clin Microbiol. 1999;37:3411–4. https://doi.org/10.1128/jcm.37.10.3411-3414.1999.
51. Johnson WM, Tyler SD, Ewan EP, Pollard DR, Rozee KR. Detection of genes for enterotoxins, exfoliative toxins, and toxic shock syndrome toxin 1 in Staphylococcus aureus by the polymerase chain reaction. J Clin Microbiol. 1991;29:426–30. https://doi.org/10.1128/jcm.29.3.426-430.1991.

52. Løvseth A, Loncarevic S, Berdal KG. Modified multiplex PCR method for detection of pyrogenic exotoxin genes in staphylococcal isolates. J Clin Microbiol. 2004;42:3869–72. https://doi.org/10.1128/JCM.42.8.3869-3872.2004.

53. Lina G, Boutite F, Tristan A, Bes M, Etienne J, Vandenesch F. Bacterial competition for human nasal cavity colonization: role of staphylococcal agr alleles. Appl Environ Microbiol. 2003;69:18–23. https://doi.org/10.1128/AEM.69.1.18-23.2003.

Publisher's Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.