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Population Genetic Structure of the *Staphylococcus intermedius* Group: Insights into *agr* Diversification and the Emergence of Methicillin-Resistant Strains

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The population genetic structure of the animal pathogen *Staphylococcus intermedius* is poorly understood. We carried out a multilocus sequence phylogenetic analysis of isolates from broad host and geographic origins to investigate inter- and intraspecies diversity. We found that isolates phenotypically identified as *S. intermedius* are differentiated into three closely related species, *S. intermedius*, *Staphylococcus pseudintermedius*, and *Staphylococcus delphini*. *S. pseudintermedius*, not *S. intermedius*, is the common cause of canine pyoderma and occasionally causes zoonotic infections of humans. Over 60 extant STs were identified among the *S. pseudintermedius* isolates examined, including several that were distributed on different continents. As the *agr* quorum-sensing system of staphylococci is thought to have evolved along lines of speciation within the genus, we examined the allelic variation of *agrD*, which encodes the autoinducing peptide (AIP). Four AIP variants were encoded by *S. pseudintermedius* isolates, and identical AIP variants were shared among the three species, suggesting that a common quorum-sensing capacity has been conserved in spite of species differentiation in largely distinct ecological niches. A lack of clonal association of *agr* alleles suggests that assortive recombination may have contributed to the distribution of *agr* diversity. Finally, we discovered that the recent emergence of methicillin-resistant strains was due to multiple acquisitions of the *mecA* gene by different *S. pseudintermedius* clones found on different continents. Taken together, these data have resolved the population genetic structure of the *S. intermedius* group, resulting in new insights into its ancient and recent evolution.

*Staphylococcus intermedius* is a member of the normal flora of dogs and is also a major opportunistic pathogen responsible for the common canine skin condition pyoderma (19). *S. intermedius* has also been found in association with other animal species (1, 3, 12, 16) and can occasionally cause severe infections of humans (26, 29, 37).

Despite its prevalence among many members of the animal kingdom, an understanding of the population genetics of *S. intermedius* is lacking. Until very recently, studies of the diversity of *S. intermedius* populations have been limited to phenotypic and molecular typing approaches (1, 3, 5, 12, 16–18, 40). A relatively high degree of phenotypic diversity exists within the species, leading to the identification of different biotypes associated with specific animal hosts (16, 28), and ribotyping studies have shown that strains are typically associated with only a single host species (1, 5). Canine strains were represented by a small number of distinct ribotypes, in contrast to strains from other animals, which generally had greater diversity (17). The level of phenotypic and genotypic diversity observed among *S. intermedius* isolates has led some investigators to speculate that an *S. intermedius* group (SIG) consisting of several species or subspecies may exist (5, 28). A closely related species, *Staphylococcus pseudintermedius*, was recently described (7) and has been isolated from several animal species, including healthy dogs in a veterinary clinic in Japan (32), and from a human infection (38). Very recently, a study of Japanese isolates based on DNA sequences of *sodA* and *hsp60* genes identified three different species, *S. intermedius*, *S. pseudintermedius*, and *Staphylococcus delphini*, among isolates phenotypically identified as *S. intermedius* and suggested a reclassification of the species (33). However, the distribution of these different species among isolates from outside Japan was not examined.

Worryingly, methicillin-resistant strains of *S. intermedius* (MRSI) and *S. pseudintermedius* (MRSP) have recently emerged in veterinary clinics around the world (15, 22, 32, 44), but their evolutionary histories have not been examined. An understanding of the population genetics of *S. intermedius* is required to provide a much-needed framework for studies of the evolution, pathogenesis, and emerging methicillin resistance of *S. intermedius*. In order to investigate the population genetic structure of *S. intermedius*, we developed a multilocus
sequencing approach that included five gene loci with a range of predicted nucleotide diversities to facilitate inter- and intraspecies differentiation. Broad new insights into the diversity of _S. intermedius_ populations were obtained.

**MATERIALS AND METHODS**

**Bacterial strains.** In total, 105 isolates were examined, including 99 from an array of diseased and healthy animal species, such as dogs, humans, horses, camels, and pigeons, which were previously identified as _S. intermedius_ in different centers in the United States, Canada, Japan, United Kingdom (including Scotland and England), France, Belgium, Czech Republic, Germany, and Sweden (see Table S1 in the supplemental material). Identification was carried out with standard phenotypic tests that varied slightly depending on the laboratory, but included production of coagulase and anaerobic acid from mannitol, sucrose, and _D_-trehalose and lack of hyaluronidase production, lack of growth on _P_ agar containing acriflavine, and the production of _β_-galactosidase when grown in _o_ -ni trophenyl-β-D-galactopyranoside broth according to the original description of the species by Hajek (16). In addition, the type strain of _S. delphini_, ATCC 49171 (39), four strains of the recently described _S. pseudointermedius_, including the type strain, LMG 22237 (7), and the type strain of _Staphylococcus schleiferi_ subspe. subspe., ATCC 43808 (11), were included in the study.

**Bacterial growth conditions and genomic DNA isolation.** Staphylococcal strains were grown on tryptic soy agar (Oxoid) at 37°C overnight or in tryptic soy broth (Oxoid) at 37°C overnight with shaking at 200 rpm. Genomic-DNA extraction was carried out with a bacterial genomic-DNA purification kit (Edge Biosystems) according to the manufacturer's instructions. Prior to incubation at 37°C for 10 min, 125 _μg/ml_ lysostaphin (Ambo) was included.

**Gene loci selected for DNA sequence analysis.** Five loci, the 16S _rRNA_, _cpn60_ (330 bp), _tuf_, _pta_, and _agrD_ genes, were selected for DNA sequence analysis. 16S _rRNA_, _cpn60_, and _tuf_ have been used previously in single-locus approaches to differentiating staphylococcal species (7, 13, 24, 25), and _cpn60_, _tuf_, and _pta_ have been incorporated into multilocus sequence-typing schemes for differentiating strains of several different bacterial species (2, 10, 43). The accessory gene _agrD_ has been used previously in single-locus approaches to differentiate _S. intermedius_ strains into _S. intermedius_ subsp. _intermedius_, _S. intermedius_ subsp. _hominis_, _S. schleiferi_, and _S. delphini_. In addition, the type strain of _Staphylococcus schleiferi_ was characterized by a double peak for a single site in the sequencing trace for eight isolates. A mixed or contaminated template was characterized by a single peak for a single site in the sequencing trace for eight isolates. A mixed or contaminated template was indicated by the existence of at least three single-locus variants.

**PCR amplification of gene fragments.** Oligonucleotide primers (Sigma-Genosys or Invitrogen) were designed for the 16S _rRNA_, _cpn60_, and _agrD_ genes based on gene sequences available for _S. intermedius_ in GenBank (accession numbers Z26897, AF019773, AY965912, and AF346723), and _S. intermedius_ _pta_ nucleotide sequence (unpublished data). Oligonucleotide sequences specific for the _tuf_ and _meCA_ genes were designed in previous studies (21, 31). The oligonucleotide sequences and predicted PCR products are given in Table 1. Gene fragments were amplified with the following: 16S _rRNA_ (570 bp), 5'-CCCTTTGCGAGACGACAAAGTAAGC-3' (forward) and 5'-GACCGGCGGATGTGATCC-3' (reverse); _tuf_ (500 bp), 5'-CAATACTGCGAATCACTCACG-3' (forward) and 5'-GCTTCAACCTAAATCAATAC-3' (reverse); _cpn60_ (560 bp), 5'-GGCGACTGTATACTCAGCCGAA-3' (forward) and 5'-AACTTACACCGCAGCTTTAC-3' (reverse); _agrD_ (430 bp), 5'-GGCGGCTTTTATACTAGTTC-3' (forward) and 5'-CGTATGAGCGTTGAGCGCTC-3' (reverse); _meCA_ (310 bp), 5'-GGGGATGTTGCAAATTCGAATTAAGC-3' (forward) and 5'-GGGGACGTGGTCTGGCTGAC-3' (reverse); _pta_ (570 bp), 5'-GGCAGGTATGTTGTTGGAGGAGG-3' (forward) and 5'-AAGAGTCCGACGGAATATGC-3' (reverse) and 5'-GCTATGAAATCTGGTGAAGAC-3' (reverse). PCR products were purified by incubation with exonuclease I and Antarctic phosphatase (NEB) for 1 min at 72°C, with a final extension for 7 min at 72°C (36). PCR products were purified by incubation with exconuclease I and Antarctic phosphatase (NEB) at 37°C for 15 min, followed by inactivation at 80°C for 15 min.

**DNA sequencing.** Sequencing reactions were carried out using 5 _μl_ purified PCR-amplified DNA (approximately 50 to 90 ng) plus 1 _μl_ sequencing primer (3.2 _pmol/μl_) with the BigDye Terminator v3.0 Ready Reaction cycle sequencing kit (Applied Biosystems) according to the manufacturer's instructions. Each sequencing reaction included 25 cycles with denaturation for 3 min at 95°C, annealing for 20 s at 50°C, and extension for 4 min at 60°C. The nucleotide sequence was determined with a 3730 DNA analyzer (Applied Biosystems). For the 16S _rRNA_, _cpn60_, _tuf_, and _pta_ genes, forward and reverse sequencing reactions were carried out with independently amplified PCR products to rule out the possibility of Taq polymerase-generated errors. For _agrD_, forward and reverse sequencing reactions were carried out from a single PCR product generated with Vent DNA polymerase (NEB), which contains proofreading activity and has a much lower predicted error rate than _Taq_ polymerase.

**DNA sequence and molecular evolutionary analyses.** DNA sequences were assembled using BioEdit Sequence Alignment Editor software (http://www.mbio.ncsu.edu/BioEdit/bioedit.html) and the Staden package (35). Nucleotide and amino acid sequences were aligned using AlignX in Vector NTI Advance10 (Invitrogen). Phylogenetic analyses were carried out with MEGA v.3.1 software (23). The neighbor-joining method was applied to construct phylogenetic trees using the Kimura two-parameter model, while the degree of statistical support for DNA sites on the minimum evolution tree was evaluated using 1,000 neighbor-joining tree bootstrap recoveries. Percent recoveries in 1,000 resample bootstrap trees were used for estimating recombination, RDP software, which includes the programs GENECONV, BOOTSCAN, MAXIMUM κ², CHIMAERA, and SISTER SCANNING, was used (27). The index of association standardized (I_σ²) between the different gene loci was calculated using the LIAN program (version 3.1; Department of Biotechnology and Bioinformatics, University of Applied Sciences, Weihenstephan, Germany) (http://adenein.biz.fh-weihenstephan.de/lian3.1). The value of the I_σ² would be expected to be zero for linkage equilibrium when recombination events occur frequently. If the I_σ² value differs significantly from zero (σ < 0.05), recombination should be rare.

**eBURST analysis.** Predicted lines of evolutionary descent and clonal complexes in our collection of isolates were identified using the eBURST algorithm (http://eburst.mlst.net). Sequence types (STs) were included in the same group if they shared four of the five gene loci with at least one other ST within the group. Subgroups were defined by the existence of at least three single-locus variants.

**Nucleotide sequence accession numbers.** The DNA sequences generated in this study were deposited in GenBank, accession no. EU157185 to EU157715.

**RESULTS**

**Multilocus sequence analysis of _S. intermedius_.** Overall, 192,482 bp representing 521 sequences from 105 isolates, including 99 isolates phenotypically identified as _S. intermedius_, 4 isolates of _S. pseudointermedius_, and the type strains of _S. delphini_ and _S. schleiferi_ subspe. subspe., were generated. 16S _rRNA_ gene sequences contained 1.37%-variable sites and 7 alleles. _tuf_ sequences contained 2.4% and 9 alleles, _cpn60_ contained 12.76% and 28 alleles, _pta_ contained 10.98% and 18 alleles, and _agrD_ contained 13.04% polymorphic nucleotide sites and 9 alleles (Table 1). 16S _rRNA_ gene sequence analysis revealed the existence of five ambiguous nucleotide sites characterized by a double peak for a single site in the sequencing trace for eight isolates. A mixed or contaminated template was ruled out by repurification of the strain to a single colony before genomic-DNA isolation, repeat PCR, and sequencing. These data are consistent with the existence of intragenomic 16S _rRNA_ gene polymorphisms, which have been previously observed (6). Phylogenetic reconstructions were carried out using concatenated sequences, which included both possible 16S _rRNA_ alleles to determine if they influenced tree topology, but no significant difference in topology was observed.
Phylogenetic analysis reveals that *S. pseudintermedius* and not *S. intermedius* is the common canine pyoderma pathogen. Neighbor-joining trees were constructed with MEGA3.1 using the Kimura two-parameter model combined with 1,000 resampling tests. The sequences were not included in the concatenated sequence analysis (Fig. 1). Phylogenetic analysis based on the concatenated sequence data indicated the existence of three major phylotypes with strong confidence (bootstrap values, 100% for each node) (Fig. 1). For comparison, phylogenetic-tree reconstruction was also carried out using the Minimum Evolution and unweighted-pair group method using average linkages, and each resulted in a topology that was cognate with the neighbor-joining trees (data not shown).

The third major phylotype was represented by isolates from horse, camel, and pigeon hosts and included the *S. delphini* type strain, ATCC 49171, isolated from a dolphin (39), suggesting that *S. delphini* may be commonly misidentified as *S. intermedius*. Taken together, these data indicate the existence of at least three closely related but distinct species among isolates that are identified as *S. intermedius* and not *S. intermedius* is the common canine pyoderma pathogen.

*S. pseudintermedius* has a largely clonal population structure. As mentioned above, trees constructed with sequences from each housekeeping locus appeared to be generally in good agreement, indicating that recombination has not played a major role in the evolution of these genes and consistent with a clonal population structure (see Fig. S1 in the supplemental material). To further investigate the population structure, we calculated the degrees of linkage disequilibrium within the whole SIG and within the *S. pseudintermedius* and *S. delphini* species (Table 2). The value of the *I*^*s* for the SIG as a whole was 0.1510 (*P* < 0.001), and for the *S. pseudintermedius* and *S. delphini* species it was 0.0877 (*P* < 0.001) and 0.3699 (*P* < 0.001), respectively, indicating that there is linkage disequilibrium within each group tested (Table 2). In addition, the RDP2 program was used to search for evidence of recombination among the selected gene loci (27). For each locus, none of the recombination detection methods, GENECONV, BOOTSCAN,
significant from zero, recombination should be rare.

MAXIMUM $\chi^2$, CHIMAERA, and SISTER SCANNING, detected any evidence for recombinant sequences among the isolates examined (data not shown). Taken together, these data indicate a largely clonal population structure for the SIG as a whole and for the $S.\ pseudintermedius$ and $S.\ delphini$ species.

$S.\ pseudintermedius$ clonal diversity and intercontinental distribution. We identified the number of alleles at each of the five gene loci and defined unique complements of alleles as STs (see Table S1 in the supplemental material). $S.\ pseudintermedius$ is characterized by the existence of 61 different STs among 89 isolates examined, indicating considerable clonal diversity within the species (see Table S1 in the supplemental material). The limited number of $S.\ pseudintermedius$ isolates that were from human and noncanine animal sources commonly shared identical or closely related genotypes with $S.\ pseudintermedius$ canine commensal strains, consistent with zoonotic transfer from dogs (Fig. 1). In addition, identical or closely related $S.\ pseudintermedius$ STs were found on different continents, indicating broad geographic dissemination of successful clones (Fig. 1).

Identification and distribution of $agr$ types among the SIG of closely related species. DNA sequencing of the $agrD$ locus revealed the presence of four predicted AIP variants among the strains examined, including a novel fourth variant that has not been described previously (designated type IV) which is specific to 27% of all strains (Table 3). The AIP variants may correspond to distinct $agr$ interference groups (8, 30, 36, 42). All staphylococcal species examined to date have been shown to encode AIP peptides that are unique to each species. In $S.\ pseudintermedius$, four AIP variants (I to IV) were encoded among the isolates; AIP variants I, II, and IV were identified among the $S.\ delphini$ group of isolates; and the $S.\ intermedius$ type strain and related pigeon isolates all encoded AIP type I (see Table S1 in the supplemental material).

A lack of clonal association of $agrD$ alleles was observed, and several isolates shared identical genotypes by 16S rRNA gene, $tuf$, $cpn60$, and $pta$ but encoded different AIP variants (Fig. 1). Further, the $agr$ gene tree topology was markedly different from those of the other gene trees. These data and the lack of recombination breakpoints identified within $agrD$ sequences using the RDP2 suite of programs suggest that assortive (whole-gene) recombination may have contributed to the distribution of $agr$ alleles. There was no identifiable association between $agr$ type and host, clinical, or geographic origin.

MRSP strains have evolved by multiple mecA gene acquisitions by different clones. The presence of the mecA gene in 16 of 105 isolates was detected by PCR (see Table S1 in the supplemental material). Sequencing of the mecA gene in representative isolates revealed a high degree of homology with mecA of Staphylococcus aureus origin (95 to 100%; data not shown). These strains had previously been identified as MRSI in the centers where they were isolated and had oxacillin MIC levels of $>256 \mu g/ml$ for all strains except one (strain 13), which had an MIC of 0.75 $\mu g/ml$. All strains previously identified as MRSI belonged to the $S.\ pseudintermedius$ phyotype and are therefore reclassified as MRSP. MRSP genotypes are distributed widely across the diversity identified within the $S.\ pseudintermedius$ phylogenetic tree, including one isolate, 3279, that is genetically indistinguishable from methicillin-sensitive isolates (ST29), suggesting horizontal transfer of the mecA gene (Fig. 1). The existence of multiple MRSP clones is supported by eBURST analysis, which shows that MRSP isolates belong to five distinct STs (ST29, ST68, ST69, ST70, and ST71) that differ at two to four of the five loci examined, strongly suggesting that they do not share a very recent ancestor (Fig. 2). Taken together, these data suggest that the mecA gene has been acquired multiple times by different $S.\ pseudintermedius$ strains. Of note, 9 out of 10 MRSP isolates from five different centers in Sweden and Germany belong to ST71, indicating that a common clone may predominate in North and Central Europe (see Table S1 in the supplemental material).
was no sharing of STs among European and U.S. MRSP isolates (see Table S1 in the supplemental material).

DISCUSSION

Hajek and colleagues first described the novel coagulase-positive species *S. intermedius*, which was isolated from pigeons, dogs, minks, and horses in 1976 (16). The type strain, NCTC 11048 (also known as ATCC 29663, CCM 5739, or LMG 13351), isolated from a pigeon, has typically been used to represent *S. intermedius* in species differentiation studies (7, 11, 13, 24, 25). Very recently, Sasaki and colleagues carried out nucleotide sequence analysis of the *sodA* and *hsp60* (*cpn60*) genes and identified *S. intermedius*, *S. pseudintermedius*, and *S. delphini* among isolates from Japan phenotypically identified as *S. intermedius* (33). Here, we show that isolates obtained from 10 different countries on three continents have a population structure that is consistent with that identified among Japanese isolates by Sasaki et al. (33). Our data indicate that the pigeon isolate represents a distinct taxon that is not representative of the majority of isolates commonly identified as *S. intermedius*. All canine isolates examined (*n* = 75) belong to the phylotype that includes four isolates of the recently described *S. pseudintermedius* species, including the type strain (7). Importantly, two *S. pseudintermedius* strains (ST5) were genetically indistinguishable from canine isolates phenotypically identified as *S. intermedius*. These data indicate that the newly described *S. pseudintermedius* species and not *S. intermedius* is the common cause of canine pyoderma. *S. pseudintermedius* is characterized by over 60 different STs identified among the 89 isolates examined, revealing considerable clonal diversity within the species. Identical and closely related STs were identified in several countries on different continents, indicating global dissemination of the most successful clones. The clonal diversity and broad geographic distribution of *S. pseudintermedius* suggests that it has coevolved with its canine host for a long time in evolutionary terms and possibly since the evolution of dogs between 40 and 50 million years ago (20, 41). STs of *S. pseudintermedius* isolates infecting humans are identical or closely related to commensal isolates of dogs, suggesting that human infections are due to zoonotic transmission from a canine host. Indeed, transmission of *S. pseudintermedius* has been shown to occur frequently between dogs affected by deep pyoderma and their owners (14).

The third major phylotype found among the isolates previously identified as *S. intermedius* was associated with several different host species, including horse, camel, and pigeon, and was phylogenetically aligned with the *S. delphini* type strain isolated from a dolphin (39). A previous DNA-DNA hybridization analysis confirmed that the *S. delphini* type strain rep-
resented a distinct species compared to the recently described \textit{S. pseudintermedius} and the \textit{S. intermedius} type strain (7). Taken together, these data suggest that \textit{S. delphini} is commonly misidentified as \textit{S. intermedius} and may be more clinically important than was previously thought. In fact, very few studies have reported the identification of \textit{S. delphini} strains since the original description of the species, but one study in Norway isolated \textit{S. delphini} from a case of bovine mastitis, extending the broad host range observed in the current study (4). Of note, Sasaki et al. indicated that more than one species may exist among isolates genetically allied with \textit{S. delphini} (33).

The accessory gene regulator (\textit{agr}) is conserved throughout the staphylococci but has diverged along lines that appear to parallel speciation within the genus (8, 42). This divergence has given rise to a novel type of intraspecies and interspecies cross-inhibition that likely represents an essential aspect of staphylococcal biology and may be a predominant feature of the evolutionary forces that have driven it. In order to further investigate the diversity of \textit{SIG} populations, we examined the diversity of the \textit{agrD} locus, which encodes the AIP (8, 20, 36, 42). Sung et al. identified three \textit{agrD} alleles corresponding to \textit{agr} specificity groups encoding different AIP variants among 20 strains of \textit{S. intermedius} isolated from dogs in a single veterinary hospital in the United Kingdom (36) and demonstrated biological activity for two of them. Here, we identified the same three predicted AIPs, in addition to a novel fourth AIP, which was encoded by approximately 27\% of strains (Table 3).

Of note, all of the predicted AIP variants of the \textit{SIG} contain a scissile in place of the conserved cysteine found in the AIPs of the other staphylococcal species analyzed to date, resulting in a lactone rather than a thiolactone ring (20). \textit{S. aureus} populations are also divided into four distinct groups based on \textit{agr} allelic variation. Interference in virulence gene expression caused by different \textit{S. aureus} \textit{agr} groups has been suggested to be a mechanism for isolating bacterial populations and a fundamental basis for subdividing the species (42). All previously examined staphylococcal species encode AIPs that are unique to each species (8). However, we found that \textit{agr} alleles were shared among the three closely related species of the \textit{SIG}, indicating that a common quorum-sensing capacity has been maintained despite species differentiation in largely distinct ecological niches.

A previous study of \textit{agr} evolution in \textit{S. aureus} reported that genotypes were not associated with more than one \textit{agr} type, indicating that \textit{agr} radiation preceded clonal diversification and that recombination has played a very limited role in the distribution of \textit{agr} diversity (42). The study concluded that the species was phylogenetically structured according to \textit{agr} group (42). More recently, Robinson et al. provided evidence that the \textit{S. aureus} species could be divided into two subgroups that both contain multiple clonal complexes and \textit{agr} groups (30). The authors proposed that recombination events had resulted in the sharing of \textit{agr} groups between the two subgroups but concluded that recombination of \textit{agr} has not occurred very frequently within \textit{S. aureus} populations, as \textit{agr} and clone tree topologies within the two subgroups were in agreement (30). In contrast, the association of different \textit{agr} alleles with strains of \textit{S. pseudintermedius} of identical genotype identified in the current study suggests that assortive recombination has frequently contributed to the distribution of \textit{agr} alleles among \textit{S. pseudintermedius} populations. The markedly different topology of the phylogenetic tree constructed with \textit{agrD} sequences compared to trees based on the other four gene loci is consistent with this theory (Fig. 1). Overall, we have found that \textit{S. pseudintermedius} has a largely clonal population structure, but recombination appears to have played an important role in the distribution of \textit{agr} alleles within the \textit{S. pseudintermedius} species and the \textit{SIG} as a whole. The sharing of \textit{agr} alleles between different species of staphylococci has not been previously observed. This discovery indicates that \textit{agr} differentiation has not occurred strictly along lines that parallel speciation. The lack of an association between \textit{agr} type and \textit{SIG} species, host, disease, and clinical or geographic origin identified in the current study leads to the question of what selective pressure is driving \textit{agr} diversification. The basis for \textit{agr} diversity and the importance of its biological activity in the \textit{SIG} remain to be elucidated.

Recently, an increasing number of episodes of \textit{S. intermedius} infections that were refractory to treatment with methicillin have been reported (15, 22, 32, 44). Here, we found that the methicillin-resistant \textit{SIG} strains examined are all classified as \textit{S. pseudintermedius} and have evolved by multiple acquisitions of the \textit{mecA} gene by different \textit{S. pseudintermedius} clones. Of note, a common clone was identified among isolates from five different centers in Sweden and Germany, indicating the existence of a widespread successful clone in Northern and Central Europe. MRSP clones were not shared between Europe and North America, indicating geographic restriction and probably reflecting the very recent emergence of methicillin-resistant strains. The identification of the common methicillin-resistant clones in the current study means that surveillance can be carried out to monitor MRSP clonal dissemination and strategies for the control of MRSP infections can be targeted to the most widespread clones.

Taken together, these data have resolved the population genetic structure of the \textit{SIG}, resulting in broad new insights into the ancient and recent evolution of this important group of animal pathogens.

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