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 trehalose and lack of hyaluronidase production, lack of growth on P agar containing acriflavin, and the production of β-galactosidase when grown in ρ-nitrophenyl-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 isolate, LM 22229 (7); and the type strain of Staphylococcus schleiferi subsp. schleiferi, 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 (bps60), 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 regulator (agr) quorum-sensing system found in the majority of staphylococcal species examined (8) is thought to have diversified along lines of speciation, giving rise to a number of subspecies groups that have the capacity for intra- and interspecies inhibition of virulence (30, 42). We examined the allelic variation of agrD, which encodes the autinducing peptide (AIP), to further investigate both inter- and intraspecies differentiation.

**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 16S tuf sequences contained 1.37% variable sites and 7 alleles, and cpn60 sequences contained 2.4% and 9 alleles, whereas 16S rRNA alleles to determine if they influenced tree topologies were purified by incubation with exonuclease I and Antarctic phosphatase (NEB) kit (Applied Biosystems) according to the manufacturer’s instructions. Each reaction included 25 cycles with denaturation for 30 s at 95°C, annealing for 20 s at 50°C, and extension for 4 min at 60°C. The nucleotide 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), 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.unc.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 tree topology was assessed using 1,000 bootstrap replications. Predictions of evolutionary events, RDP software, which includes the programs GENECONV, BOOTSCAN, MAXIMUM κ², CHIMAERA, and SISTER SCANNING, was used (27). The index of association standardized (Iₐ), which increases with recombination events occur frequently. If the Iₐ value differs significantly from zero (p < 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://ebrust.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. EU157195 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 subsp. schleiferi, 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 resample trees by the bootstrap test using sequences for each individual locus (see Fig. S1 in the supplemental material). Trees constructed with sequences from each housekeeping locus appeared to be generally in good agreement, indicating a robust phylogenetic signal (see Fig. S1 in the supplemental material). In contrast, the topology of the tree generated with the *agrD* nucleotide sequence was very different from those of the other gene sequences, suggesting 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).

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, suggesting 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,
The small number of isolates of the *S. intermedius* phenotype precluded carrying out this analysis. ND, not done.

**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 staphyloccoccal species examined to date have been shown to encode AIP peptides that are unique to each species. In contrast, the three closely related species examined in the current study had AIP peptide variants in common, suggesting the existence of a conserved *agr* quorum-sensing system. All four AIP variants (I to IV) were encoded among the *S. pseudintermedius* 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 MRSA in the centers where they were isolated and had oxacillin MIC levels of >256 µg/ml for all strains except one (strain 13), which had an MIC of 0.75 µg/ml. All strains previously identified as MRSA belonged to the *S. pseudintermedius* phenotype 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). There

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**TABLE 2.** Analysis of linkage disequilibrium by calculation of $I_d^S$ for the SIG and for the *S. pseudintermedius* and *S. delphini* phylotypes

<table>
<thead>
<tr>
<th>Group selection</th>
<th>$I_d^S$</th>
<th>$P$ value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>SIG</td>
<td>0.151</td>
<td>0.0000</td>
</tr>
<tr>
<td><em>S. pseudintermedius</em></td>
<td>0.0877</td>
<td>0.0000</td>
</tr>
<tr>
<td><em>S. delphini</em></td>
<td>0.3699</td>
<td>0.0000</td>
</tr>
<tr>
<td><em>S. intermedius</em></td>
<td>ND</td>
<td>ND</td>
</tr>
</tbody>
</table>

$^*$ The value of the $I_d^S$ would be expected to be zero for linkage equilibrium when recombination events occur frequently. If the $I_d^S$ ($P < 0.05$) value differs significantly from zero, recombination should be rare.

**TABLE 3.** Amino acid sequences of the predicted *agrD*-encoded AIPs identified among the strains examined

<table>
<thead>
<tr>
<th>AIP type (AIP)</th>
<th>Amino acid sequence</th>
<th>No. of strains (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>RIPTSTGFF</td>
<td>16 (15)</td>
</tr>
<tr>
<td>II</td>
<td>RIPSTGFF</td>
<td>31 (30)</td>
</tr>
<tr>
<td>III</td>
<td>KIFTSTGFF</td>
<td>29 (28)</td>
</tr>
<tr>
<td>IV*</td>
<td>KYPTSTGFF</td>
<td>28 (27)</td>
</tr>
</tbody>
</table>

* Novel AIP variant identified in the current study.

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**FIG. 1.** Phylogenetic tree constructed by the neighbor-joining method with 1,000 bootstrap replicates of concatenated 16S rRNA gene, *tuf*, *cpn60*, and *pta* sequences. (A) Radiating tree indicating the three major phylotypes, with species type strains indicated by triangles. (B) Branch style tree indicating the host and geographic origins of all isolates. Methicillin-resistant, *mecA*-positive isolates (dots); human isolates (squares); previously identified *S. pseudintermedius* isolates (diamonds); and species-type isolates (triangles) are indicated, along with the *agr* type in roman numerals in parentheses. Bootstrap values over 50% are indicated.
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 \textit{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 \textit{S. intermedius} in species differentiation studies (7, 11, 13, 24, 25). Very recently, Sasaki and colleagues carried out nucleotide sequence analysis of the \textit{sodA} and \textit{hsp60} (\textit{cpn60}) genes and identified \textit{S. intermedius}, \textit{S. pseudintermedius}, and \textit{S. delphini} among isolates from Japan phenotypically identified as \textit{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 \textit{S. intermedius}. The clonal diversity and broad geographic distribution of \textit{S. pseudintermedius} suggest 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 \textit{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 \textit{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 \textit{S. intermedius} was associated with several different host species, including horse, camel, and pigeon, and was phylogenetically aligned with the \textit{S. delphini} type strain isolated from a dolphin (39). A previous DNA-DNA hybridization analysis confirmed that the \textit{S. delphini} type strain rep-

FIG. 2. Schematic diagram of the clonal relatedness of \textit{S. pseudintermedius} STs predicted by eBURST analysis. Each ST is represented by a black dot, the size of which is proportional to the number of isolates of that ST. Blue and yellow dots denote the predicted group founder and subgroup founders, respectively. Single-locus variants are linked by lines. STs that include MRSP strains are indicated by red circles. STs that are not linked by branches to other STs do not share at least four identical alleles with any other ST.
resented a distinct species compared to the recently described *S. pseudintermedius* and the *S. intermedius* type strain (7). Taken together, these data suggest that *S. delphini* is commonly misidentified as *S. intermedius* and may be more clinically important than was previously thought. In fact, very few studies have reported the identification of *S. delphini* strains since the original description of the species, but one study in Norway isolated *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 *S. delphini* (33).

The accessory gene regulator (*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 interstrain 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 SIG populations, we examined the diversity of the *agrD* locus, which encodes the AIP (8, 20, 36, 42). Sung et al. identified three *agrD* alleles corresponding to *agr* specificity groups encoding different AIP variants among 20 strains of *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 SIG contain a scerine 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). *S. aureus* populations are also divided into four distinct groups based on *agr* allelic variation. Interference in virulence gene expression caused by different *S. aureus* *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 *agr* alleles were shared among the three closely related species of the SIG, indicating that a common quorum-sensing capacity has been maintained despite species differentiation in largely distinct ecological niches.

A previous study of *agr* evolution in *S. aureus* reported that genotypes were not associated with more than one *agr* type, indicating that *agr* radiation preceded clonal diversification and that recombination has played a very limited role in the distribution of *agr* diversity (42). The study concluded that the species was phylogenetically structured according to *agr* group (42). More recently, Robinson et al. provided evidence that the *S. aureus* species could be divided into two subgroups that both contain multiple clonal complexes and *agr* groups (30). The authors proposed that recombination events had resulted in the sharing of *agr* groups between the two subgroups but concluded that recombination of *agr* has not occurred very frequently within *S. aureus* populations, as *agr* and clone tree topologies within the two subgroups were in agreement (30).

In contrast, the association of different *agr* alleles with strains of *S. pseudintermedius* of identical genotype identified in the current study suggests that assortive recombination has frequently contributed to the distribution of *agr* alleles among *S. pseudintermedius* populations. The markedly different topology of the phylogenetic tree constructed with *agrD* sequences compared to trees based on the other four gene loci is consistent with this theory (Fig. 1). Overall, we have found that *S. pseudintermedius* has a largely clonal population structure, but recombination appears to have played an important role in the distribution of *agr* alleles within the *S. pseudintermedius* species and the SIG as a whole. The sharing of *agr* alleles between different species of staphylococci has not been previously observed. This discovery indicates that *agr* differentiation has not occurred strictly along lines that parallel speciation. The lack of an association between *agr* type and SIG species, host, disease, and clinical or geographic origin identified in the current study leads to the question of what selective pressure is driving *agr* diversification. The basis for *agr* diversity and the importance of its biological activity in the SIG remain to be elucidated.

Recently, an increasing number of episodes of *S. intermedius* infections that were refractory to treatment with methicillin have been reported (15, 22, 32, 44). Here, we found that the methicillin-resistant SIG strains examined are all classified as *S. pseudintermedius* and have evolved by multiple acquisitions of the *medA* gene by different *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 SIG, resulting in broad new insights into the ancient and recent evolution of this important group of animal pathogens.

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