INTRODUCTION

Streptococcus sanguinis is a gram-positive facultative anaerobe and a primary colonizer of the oral cavity (Zhu et al., 2018). Epidemiological studies have shown that S. sanguinis has significantly increased abundance in healthy versus diseased subgingiva as well as higher abundance on sound enamel versus carious lesions (Griffen et al., 2012; Stingu et al., 2008; Zhu et al., 2018). Additionally, in vitro experiments have shown that S. sanguinis can compete against oral pathogens such as Streptococcus mutans and Porphyromonas gingivalis via hydrogen peroxide production (Chen et al., 2012; Kreth et al., 2005; Zhu et al., 2018, 2019). These studies suggest that S. sanguinis contributes to the homeostasis of the oral cavity. It is also known, however, that oral surgery, mastication, or routine hygiene practices...
can produce lesions in the oral mucosa through which S. sanguinis and other oral streptococci can enter the bloodstream (Wilson et al., 2007; Wray et al., 2008). If this happens in a person who has certain preexisting cardiac conditions, these bacteria can colonize the damaged heart valves or other endocardial surfaces, causing a disease called infective endocarditis (IE) (Baddour et al., 2015).

Antibiotics are often used prophylactically to prevent IE (European Society of Cardiology, 2015; Wilson et al., 2007) and therapeutically to treat it (Baddour et al., 2015; Wilson et al., 2007). In the former case, the primary indication is patients with high-risk cardiac conditions (Thornhill et al., 2018) who are preparing to undergo an invasive dental procedure. In the latter case, treatment can last 4 to 6 weeks (Rasmussen et al., 2017), a duration that would be expected to contribute to both the development of antibiotic resistance and to the disruption of the oral microbiome (Shaw et al., 2019). Moreover, efficacy is suboptimal, with valve surgery required in addition to antibiotics in approximately 50% of cases. Even with all available treatments, 1-year readmission and mortality rates of 65% and 33%, respectively, have been reported (Rasmussen et al., 2017).

We previously proposed targeted antimicrobial therapy based on the different essential gene targets present in beneficial versus pathogenic bacterial species in the microbiome (Stone et al., 2015; Stone & Xu, 2017). Here, we propose a second method to selectively target pathogens—in this instance, based on the different genetic requirements of a single bacterial species when present at a disease site versus its usual anatomical location. As applied to IE caused by oral streptococci, we propose to take advantage of potential differences in available nutrients in the oral cavity compared to infected heart valves. If an essential nutrient is present at a lower concentration in the bloodstream than in the oral cavity, genes required for its uptake should be much more important for bacteria on blood-bathed endocardial surfaces than in the oral cavity. A drug that specifically inhibits S. sanguinis in the bloodstream may prove more efficacious than conventional antibiotics for IE prevention or treatment while leaving the oral microbiome intact. In addition, several studies have identified S. sanguinis strains from IE cases that are resistant to penicillin (Pericás et al., 2019), linezolid (Mendes et al., 2013), and vancomycin (Saffar & Rolston, 2006). Thus, finding new drug targets for the selective inhibition of S. sanguinis in the bloodstream could lead to fewer treatment failures.

In the process of identifying the essential genes in S. sanguinis SK36, we constructed a comprehensive library of nonessential genes containing 2046 single-gene mutants (Xu et al., 2011). In this study, we tested the growth of these mutants in human serum (HS) and brain heart infusion (BHI) broth using a sequencing protocol similar to transposon sequencing (Tn-seq) that we refer to as ORF-seq. The network of genes whose corresponding mutants had significantly reduced growth in HS was constructed by predicting the protein–protein interactions in the STRING database (Szklarczyk et al., 2017), to investigate the linkage between the 178 mutants whose corresponding mutants exhibited a significant change in abundance (log2 fold change ≥ 0.585 or ≤ −0.585 and p value ≤ .05) in HS including 138 mutants whose abundance was decreased (SI Spreadsheet 1).

2 | RESULTS

2.1 | Identification of fitness genes of S. sanguinis in HS using ORF-seq

Separate cultures of 2046 S. sanguinis mutants, each of which contains a deletion in a single nonessential gene, were pooled together, harvested, and inoculated into HS, which was incubated at 37°C under microaerobic conditions (6% O2) to mimic the nutrient conditions in the bloodstream. The pooled bacterial cells were collected after 24 hr incubation (T24) and compared with the bacterial cells in the original library (T0). The abundance change of mutants was determined by a procedure with similarities to Tn-seq that we have named ORF-seq (See Materials and Methods for details.) A total of 178 mutants were identified with significant abundance changes (log2 fold change ≥ 0.585 or ≤ −0.585 and p value ≤ .05) in HS including 138 mutants whose abundance was decreased (SI Spreadsheet 1).

2.2 | Network analysis of S. sanguinis fitness genes in HS

We used STRING, a comprehensive database for the prediction of protein–protein interactions (Szklarczyk et al., 2017), to investigate the linkage between the 178 proteins whose corresponding mutants exhibited a significant change in abundance in HS. The linkage was put into Gephi for visualization and analysis of the network (Bastian et al., 2009). There are 149 proteins that were annotated in STRING and present in the network (Figure 1 and SI Spreadsheet 2). Others are uncharacterized proteins or proteins without information in the STRING database, whose functions cannot be predicted. The main network contains 135 proteins and contains several small, independent proteins groups.

Most of the mutants with altered abundance had reduced rather than increased abundance after 24 hr of incubation in HS (Figure 1). Four of these mutants correspond to genes that we identified previously as encoding virulence factors for IE in a rabbit model: ssaB, purB, lgt, and ccpA (Bai et al., 2019; Das et al., 2009; Paik et al., 2005). This is not surprising because a mutant that cannot grow in serum is unlikely to grow on an infected heart valve with blood as its primary source of nutrition.

The 10 mutants with the largest decrease in abundance were deleted for purA, rplA, clpP, priA, hom, purB, guaA, nadD, guaB, or rpoE (Figure 1 and SI Spreadsheet 2). Four of the mutants (purA, purB, guaA, and guaB) belong to the purine metabolism category in the Clusters of Orthologous Groups (COG) database (Tatusov et al., 2000), indicating that purine metabolism is required for normal growth of S. sanguinis in
HS. Similar findings have been obtained previously with other bacterial species (Le Breton et al., 2013; Samant et al., 2008; Zhang et al., 2017). Details of these four genes will be discussed below.

Functional enrichment analysis was performed using the DAVID database (Huang et al., 2009) for the 138 genes whose mutants exhibited significant decreases in abundance in HS. Three KEGG pathways—ABC transporters, purine metabolism, and arginine and proline metabolism—were found to be significantly enriched (Figure 2a). The biological functions of the proteins in the network were annotated using the COG database and sorted by the number of proteins involved in these functions (Tatusov et al., 2000). The top eight functions are depicted in Figure 2b. The pathway enrichment results were similar to those obtained from the COG annotation.

The size of nodes in the network is positively related to the betweenness centrality of proteins (Figure 2b). There exists at least one shortest path between two nodes in a network. The betweenness centrality describes the number of different shortest paths passing through a node in a network (Freeman, 1977). A protein with a larger betweenness centrality is more central in a network. In our network, GuaA, GuaB, and PurL are the three largest nodes (Figure 2b). Their biological functions are related to the purine metabolism pathway.

Other hub proteins (betweenness centrality ≥ 400) are also apparent, such as Pkn, Gcp, Lgt, and RgpF (Figure 2b). The direct link from these four hub proteins to other proteins in the network is shown in Figure S1. In our ORF-seq results, all four of these hub gene’s mutants had reduced growth in HS (Figure 1). However, the growth reduction exhibited by these four mutants was less pronounced than that of other mutants with less betweenness centrality. For example, ΔssaB and ΔstpA both had larger decreases in abundance in HS than Δlgt, (Figure 1). These data suggest that, although the four hub proteins interacted with other proteins or might modulate other proteins at the posttranslational level (Bugrysheva et al., 2011), proteins directly related to nutrient uptake and synthesis such as PurA, PurB, SsaB, and StpA seemed to be more important (Figure 1).

2.3 | The impact of purine metabolism on S. sanguinis growth in HS

We next repeated the ORF-seq experiment as before, except in BHI. Because the library was constructed in BHI-grown cells, we knew
that all the mutants should be capable of growth, although not at the same rate. We performed the same enrichment analysis described above with genes whose mutants had decreased abundance after 24 hr of incubation in BHI. Interestingly, homologous recombination was the only pathway that was significantly enriched in BHI (SI Spreadsheet 3). By comparing the results of the two ORF-seq studies, we identified 88 mutants whose relative abundance was altered with a log$_2$ value ≥ |0.585| in HS but not in BHI. These 88 mutants are depicted by a bigger node size in Figure 1, in blue in Figure S2 and their locus tags are highlighted in yellow in SI Spreadsheet 1. Although any gene required for growth in HS is of interest, we reasoned that those that were not needed for growth in the rich medium BHI might be more likely to be HS-specific. We noted that within this group, 10 of the 12 mutants with the greatest defect in HS abundance belonged to the purine metabolic pathway. As depicted in the KEGG database, purine metabolism is one of two components of the nucleotide transport and metabolism pathway (Figure 3a) (Kanehisa et al., 2017). The genes on the left side of phosphoribosyl pyrophosphate (PRPP) are responsible for purine salvage and synthesis while the genes on the right side are responsible for pyrimidines. These pathways have been well characterized (Kilstrup et al., 2005; Leonard et al., 2018). Many of the genes in both pathways were found to be essential, as indicated in Figure 3a by the blue coloring. The corresponding mutants could not be constructed and their phenotypes could therefore not be assessed (Xu et al., 2011).

The deletion of pur genes whose products catalyze de novo purine synthesis from PRPP/L-glutamine to AMP/GMP reduced the growth of S. sanguinis in HS, suggesting that de novo synthesis of AMP and GMP was important in HS (Figure 3a). In contrast, most of the genes required solely for purine salvage had little impact on growth, including punA, deoD, apt, and xpt (Figure 3a). This could either be because salvage is only important for serum growth when de novo synthesis is blocked by mutation or because the gene products, which are nucleotide phosphorylases (PunA, DeoD) or phosphoribosyltransferases (Apt, Xpt), are at least partially redundant. The mutation of hpt, which also encodes a phosphoribosyltransferase (hypoxanthine → IMP), did reduce abundance in HS significantly (log$_2$ fold change = −2.15), suggesting that purine salvage contributed to the growth of S. sanguinis in HS (Figure 3a). Nevertheless, the deletion of hpt had less impact than deletion of genes such as purH, purC, and purE, indicating that de novo synthesis was more important than purine salvage from inosine for growth in HS (Figure 3a and SI Spreadsheet 1). Finally, it should also be noted that the four mutants (purA, purB, guaA, and guaB) that displayed the largest reductions in abundance in HS out of all the purine mutants tested are each blocked in the conversion of inosine to either AMP or GMP, which are reactions shared by the de novo and salvage pathways. The combined results suggest that salvage plays a role in purine replenishment in HS, but that de novo synthesis makes a greater contribution.

Given that salvage requires less energy than synthesis, we predicted that its limited contribution to growth in HS likely results from purine nucleoside concentrations being too low to support normal growth of S. sanguinis. To test this hypothesis, we consulted the Human

![Figure 2](https://wileyonlinelibrary.com)
Metabolome Database (HMDB) (Wishart et al., 2018), which showed that the concentration of L-glutamine in blood was more than 500 μM, whereas the concentrations of adenosine, guanosine, and inosine were less than 1 μM for most of the results reported (Figure 3b), in agreement with our expectations. We then added 0.15 mg/ml (560 μM) inosine to HS and the growth of S. sanguinis wild type (WT), the ΔpurC mutant, and the ΔpurH mutant was measured after 24 hr of incubation in individually inoculated tubes. The addition of inosine did not significantly impact the colony-forming unit (CFU) numbers for WT (Figure 3c), but did increase the CFUs of ΔpurC and ΔpurH, which was also consistent with our hypothesis (Figure 3c). In other words, only when the pathway from L-glutamine/PRPP to AMP/GMP was blocked, would the production of AMP/GMP from inosine salvage noticeably improve growth. When cells were grown in BHI, the precursors for de novo synthesis and nucleosides for salvage were sufficient to support normal growth of the pur mutants (ΔpurA, ΔpurB, ΔpurC, ΔpurD, ΔpurE, ΔpurF, ΔpurH, ΔpurK, ΔpurL, ΔpurM, and ΔpurN) (Figure S2 and SI Spreadsheet 1). Interestingly, however, the guaA and guaB mutants grew poorly in BHI (although better than in serum), suggesting that guanosine levels in BHI are not sufficient to support normal growth in the absence of conversion of inosine to GMP (SI Spreadsheet 1).

Compared to purine metabolism, genes involved in pyrimidine metabolism had less impact on S. sanguinis growth in HS (Figure 3a). The genes responsible for biosynthesis from PRPP/L-aspartate/L-glutamine to UMP had little to no effect on the abundance of S. sanguinis in HS (Figure 3a). One possibility was that there was enough

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**FIGURE 3** The effect of nucleotide metabolism on the growth of S. sanguinis in HS. (a) The KEGG nucleotide transport and metabolism pathway is shown. The abundance changes of the respective mutants in HS in ORF-seq are indicated by color according to the scale shown. (b) Metabolite concentrations in blood in the form of mean ± standard deviation were downloaded from the Human Metabolome Database (HMDB) and are depicted here. (C) The growth of ΔpurC and ΔpurH was measured after 24 hr of incubation with or without supplementation with 0.15 mg/ml inosine. The CFUs of each strain in HS+ inosine are expressed relative to that of the same strain in HS. Means and standard deviations from triplicate experiments are shown. **p value ≤ .01, Student’s t test [Colour figure can be viewed at wileyonlinelibrary.com]
UMP from another pathway to compensate for loss of de novo biosynthesis. The HMDB indicated concentrations of uridine greater than 3 μM for all three reported results, whereas most of the reported concentrations of adenosine, guanosine, and inosine were lower than 1 μM, which may explain why de novo biosynthesis of UMP was less important than AMP/GMP (Figure 3b).

### 2.4 | Transporters important for the growth of S. sanguinis in HS

In gram-positive bacteria, transport proteins localize to the sole cell membrane. Therefore, many of the component proteins are surface-accessible making them appealing drug targets. Using the KEGG pathway annotation tool (Kanehisa et al., 2017), we identified four types of ABC transporters whose mutants had reduced abundance in HS. These included six putative metal transport genes (ecfA2, ecfT, stpA, ssaA, ssaB, and czcD), four peptide and nickel transport genes (oppF, oppD, oppB, and amiF), three phosphate and amino acid transport genes (pstB1, livF, and SSA_2099), and one sugar transporter (SSA_1007) (SI Spreadsheet 1). Deletion of the metal transport genes caused the greatest reduction in S. sanguinis growth in HS (SI Spreadsheet 1). We examined the most promising mutants by individual inoculation into tubes containing HS or saliva, followed by dilution plating and colony counts.

The czcD gene has been studied in S. pneumoniae, where it is responsible for transporting Zn out of cells (Martin & Giedroc, 2016). The ortholog in S. sanguinis (SSA_2321) is a closely related to the pneumococcal protein (SPD_1638), and the two are reciprocal best hits by BlastP analysis. The czcD deletion mutant is sensitive to Zn toxicity (Martin & Giedroc, 2016). Metabolomics data from HMDB showed that the concentration of Zn in blood is higher than that in saliva (Figure 4a), which might explain why ΔczcD had poor growth in HS. In individual inoculation assays, growth of ΔczcD was normal in saliva but growth was much lower in HS after 6 hr of incubation, which was consistent with this hypothesis (Figure 4b). However, the final CFUs of ΔczcD cultures were similar to WT after 24 hr, suggesting that the deletion of czcD reduced the growth rate rather than achievable density (Figure 4b).

The Mn transporter mutant ΔssaA was similar to ΔssaB, with both showing defective growth in HS as assessed by ORF-Seq (SI Spreadsheet 1). Although the ΔssaC mutant displayed a decreased but insignificant abundance change by ORF-seq (SI Spreadsheet 1), the reduction in growth in HS was significant when the ssA mutant was assessed in an individual inoculation assay (Figure 4d). Metabolomics data from HMDB showed a seemingly higher concentration of Mn in saliva than in blood or serum, although the difference was not significant (Figure 4c). Individual inoculation growth studies performed in HS and saliva revealed that the three ssa gene deletion mutants demonstrated reduced growth in both HS and saliva (Figure 4d).

ECF genes belong to the energy-coupling factor (ECF) family of ABC transporters (Rempel et al., 2019). Deletion of ecfA2, ecfT, or stpA attenuated the growth of S. sanguinis in both HS and saliva by individual inoculation (Figure 4e). However, the growth defects of these three mutants were more severe in HS than in saliva (Figure 4e); the average decrease of these three mutants in HS was ~2.4 and ~6 times larger than that in saliva at 6 and 24 hr, respectively. These three mutants will be examined further as described below.

### 2.5 | Mutants with defective growth in HS but normal growth in saliva

In addition to the transport mutants described above, we selected a small number of mutants whose growth was reduced in HS but not in BHI and compared their growth in HS versus saliva by individual inoculation. Two mutants, ΔSSA_0048 and ΔSSA_2241, were identified that were recovered from HS at significantly lower numbers than WT at 24 hr, but demonstrated normal growth in saliva (Figure 4f). Both of the corresponding genes are annotated as encoding hypothetical proteins (Xu et al., 2007). Further studies will be required to determine their functions.

### 2.6 | Characterization of ECF transporters and contribution to endocarditis virulence

Given the importance of the ECF transporters to serum growth and their lesser effect on growth in saliva, we decided to examine them further. We began by attempting to determine the function of these genes, which were annotated as cobalt (Co) transporters (Xu et al., 2007). We first deleted the ecfA2, ecfT, and stpA genes, which belong to the same operon, to create a triple mutant (ΔECF), then, complemented that mutant by placing the three genes at an ectopic expression site used previously for this purpose (Sentry Turner et al., 2009). We then employed inductively coupled plasma-optical emission spectroscopy (ICP-OES) to measure the levels of Co and other divalent cations in SK36, the ΔECF mutant, and the complemented mutant. As shown in Figure 5a, there was no significant difference in Co levels in the mutant compared to SK36 or the complemented mutant. This was also true for Fe and Zn. Surprisingly, Mn levels were significantly lower in the ΔECF mutant and significantly higher in the complemented mutant (Figure 5a). We were interested in further investigating this finding, since we had previously linked the virulence reduction of an ssaB mutant to reduced Mn uptake (Crump et al., 2014). We therefore combined the ΔssaACB mutation with the ΔECF mutation and tested this combined mutant alongside SK36, the two ECF strains, and the ΔssaACB mutant by ICP-OES. As shown in Figure 5a, Mn levels in the ΔECF mutant while lower than WT, were significantly higher than those in the ΔssaACB mutant, and deletion of the ECF genes from the ΔssaACB mutant did not result in a significant reduction in Mn levels compared to the ΔssaACB mutation alone. We also tested the same strains for their ability to grow in rabbit serum in 12% of O₂, the same level found in arterial blood (Atkuri et al., 2007), since this would allow a direct comparison with data collected previously for the ssaB (Crump et al., 2014) and ΔssaACB (Murgas et al., 2020) mutants. As shown in Figure 5b, all of the
strains grew indistinguishably in the overnight BHI precultures used to inoculate the rabbit serum ("0 h" samples); however, the ΔECF and the ΔssaACB mutants showed no apparent growth, whereas the complemented ΔECF mutant grew to WT levels. Interestingly, the combined ΔECF ΔssaACB mutant appeared to experience net death during the course of the 24-hr incubation in rabbit serum, exhibiting CFU values that were significantly lower than at the time of inoculation and also significantly lower than every other strain at 24 hr.

In a previous study that included SK36 and an ssaB mutant, growth in rabbit serum in 12% of O₂ predicted IE virulence in a rabbit model (Crump et al., 2014), with SK36 recovered from infected hearts in numbers ~10⁵-fold greater than the ssaB mutant. Given that the ΔECF mutant exhibited a defect in serum growth similar to that of the ΔssaACB mutant, we were interested in testing the ΔECF mutant in our rabbit endocarditis model as well (Crump et al., 2014). The ΔECF mutant, the complemented ΔECF mutant, and JFP36, a derivative of SK36 that possesses a gene for erythromycin resistance (Emr) inserted into the same ectopic locus as the ECF complemented mutant, were co-inoculated into each rabbit. We then used plating on selective antibiotics to determine CFU numbers in the inoculum and the infected vegetations, as we have done previously (Crump et al., 2014). Figure 6 shows that, as with the serum growth study shown in Figure 5b, the ΔECF mutant was recovered at levels ~10⁵-fold lower than the Emr WT strain. The complemented mutant was recovered at levels 1.6-fold higher than—though not significantly different from—the WT strain. Thus, the virulence defect of the ΔECF mutant is comparable to that of the ssaB mutant.

**Figure 4** Comparison of HS and saliva for metal content and cultivation of S. sanguinis strains. (a and c) Metal concentrations in blood and saliva downloaded from HMDB in the form of mean ± standard deviation. Data were analyzed by unpaired t test, *** p value ≤ .001. (b, d, and e) Growth of strains inoculated singly into saliva or HS and incubated for 6 or 24 hr. The CFUs of mutants were expressed relative to those of WT grown under the same condition. (f) Growth of strains inoculated singly into saliva or HS and incubated for 24 hr. CFUs are expressed relative to WT grown under the same condition. Means and standard deviations from triplicate experiments are shown for all growth studies. * p value ≤ .05, ** p value ≤ .01, *** p value ≤ .001, Student’s t test [Colour figure can be viewed at wileyonlinelibrary.com]
3 | DISCUSSION

Antibiotic resistance is a risk of long-term antibiotic use. To address this issue, we have proposed the idea of developing new drugs to selectively inhibit the growth of bacteria in the bloodstream but not in other locations of the body. Such drugs could potentially be more effective for treatment, but their greatest benefit would likely be for IE prevention. Advisory groups differ in opinion regarding the merit of antibiotic prophylaxis prior to dental procedures, with those opposed suggesting that the cumulative risk from daily bacteremias likely exceeds the risk from infrequent dental procedures (Cahill et al., 2017). An antibiotic that targets bacteria in the bloodstream but not at other body sites might find acceptance by both sides for daily use in patients who have cardiac conditions that put them at
Despite its obvious successes, our ORF-seq approach has some limitations. One is that, as with Tn-seq, all of the mutants are combined into pools. This may mask the effect that some mutations would have on the ability of the corresponding mutant to grow because a growth-limiting metabolite could be provided by the other mutants. Such fitness genes cannot be identified using this method. Another potential issue could be polar effects on the expression of downstream genes, which might lead to misleading results. Both of these problems could be partially addressed by a systems biology approach, in which conclusions come from multiple lines of evidence or from different mutants. For example, our conclusion concerning the importance of purine metabolism for bacterial growth in HS was verified through findings derived from multiple mutants.

We discovered two potential selective drug targets for the inhibition of S. sanguinis growth in HS that we predict would preserve growth in saliva. To our knowledge, this is the first report of selective gene targets suppressing bacterial growth in HS but not in saliva. In our previous study, we identified three essential functions of S. sanguinis: cell envelope development, energy production, and genetic information processing (Kong & Zhu, 2019; Xu et al., 2011). Based on differences in the essential genes of different bacterial species, we have developed a selective approach that targets key pathogens in a microbiome while sparing commensal bacteria (Stone et al., 2015).

We also characterized a mutant lacking the common components of an ECF transport system. As far as we are aware, this is only the second such system to have been identified as important for virulence, with the first being an apparent heme transporter in S. pyogenes (Chatterjee et al., 2020). This is also the first ECF system shown to be important for IE virulence in any bacterium. Based on our ICP-OES analysis, the ECF proteins that we characterized play no role in Co transport in S. sanguinis. This is not surprising. The ecfA2, ecfT, and stpA genes appear to encode the common components of a Group II transport system. In such systems, the common components drive the transport of multiple substrates by interacting with distinct substrate-binding proteins encoded by noncontiguous genes (Rempel et al., 2019). Moreover, SK36 is annotated as containing a complete Co transport system of the Group I variety, where the substrate-binding protein is encoded adjacent to the other three components. This system is located within a larger locus devoted to synthesis of cobalamin, or vitamin B12 (Xu et al., 2007). Considering that cobalamin requires Co as a cofactor, this annotation is inherently plausible. In addition, misannotation of EcFA and EcFT proteins as Co transporters is a common error (Rempel et al., 2019) and the recent reannotation of these genes in GenBank no longer suggests Co transport.

These proteins may play a role in Mn transport. It seems unlikely that reduced Mn transport is responsible for the poor serum growth or virulence of the ΔECF mutant, however, given that this mutant transported far more Mn than the ΔssaACB mutant (Figure 5a) and that serum growth was not restored to the ΔECF mutant by addition of Mn to 100 µM (data not shown). For comparison, we found previously that 5 µM Mn was sufficient to fully restore the serum growth of the ΔssaACB mutant (Murgas et al., 2020). Finally, the ECF components that we characterized presumably interact with all other Group I substrate-binding protein components present in the cell. It is not clear...
how many such components there are or the substrates they transport, although the RAST-based annotation of the SK36 genome present on the PATRIC website (https://patricbrc.org/) identifies eight, most of which are thought to transport vitamins or cofactors. It is thus not surprising that the ΔECF mutant displays a severe growth defect.

To date, very few S. sanguinis genes have been confirmed as virulence factors using an animal model (Bai et al., 2019; Crump et al., 2014; Ge et al., 2008; Martini et al., 2020). We have likely identified many additional promising candidates. By ORF-seq, the ΔssaB and ΔpurB mutants exhibited a log₂ fold change in HS of −2.49 and −3.99, respectively. If these values are compared to the abundance of these mutants relative to WT in an IE animal model (log₂ fold change −7.31, respectively), the defect is clearly much greater in the animals. Similarly, the log₂ fold changes in HS for the individual ECF mutants ranged from −1.58 to −1.85, whereas the log₂ fold change for the triple mutant in the IE model was −16.67 (Figure 6). In contrast, in our previous study (Crump et al., 2014) growth of ΔssaB in rabbit serum matched the results of the IE model (log₂ fold change = −14.14 for IE vs. −19.19 for serum), just as it did for the ΔECF mutant in this study (log₂ fold change = −15.95 for serum vs. −16.67 for IE; Figure 5b vs. Figure 6). The ECF result was not due to a difference between the triple ΔECF mutant versus the individual mutants or human versus rabbit serum, since we observed similar results when each of the three individual ECF mutants was grown in rabbit serum (not shown). The explanation is most likely the degree to which the inocula were diluted for the two experiments. For ORF-seq, the mutant pool was diluted 100-fold into HS, whereas each strain was diluted 10⁴-fold into rabbit serum. Thus, a WT strain would be expected to attain an abundance ~100-fold that of a nongrowing strain by ORF-seq, whereas a ~10⁶-fold difference would be expected in the rabbit serum studies. The latter experiment is a better model of in vivo disease, where we have observed colonization of cardiac tissues by ~10⁷ CFU and recovery the following day of ~10⁴ CFU per animal (Crump et al., 2014). We can therefore conclude that the ORF-seq study provides a highly conservative view of the virulence defects that we would likely see in the IE model or in human disease. Thus, many of the mutants displaying lesser defects by ORF-seq may be worth further examination with our standard rabbit serum growth assay, followed by testing in the in vivo model.

4 | MATERIALS AND METHODS

4.1 | Bacterial strains

Strains used in the ORF-seq analysis were from our comprehensive mutant pool, which was generated from strain SK36 (Kilian et al., 1989) in our previous study (Ge & Xu, 2012). Additional strains used in this study are described below and listed in Table S1. The ΔECF mutant was constructed by replacing the three ECF genes (SSA_2367, SSA_2366, and SSA_2365, in that order) with the ada9 cassette encoding ScΔ (Senty Turner et al., 2009) by overlap extension PCR (Ho et al., 1989). The ada9 cassette, including its promoter, was inserted immediately downstream from the stop codon of the upstream gene, SSA_2368. We note that we relied on the 2020 GenBank annotation for this deletion, in which the start codon of SSA_2367 is separated by 1 bp from the stop codon of SSA_2368, rather than the two ORFs overlapping as in the earlier annotation. The ada9 cassette extended exactly to the stop codon of SSA_2365, which was 210 bp from the downstream ORF, SSA_2364. Based on expression data from multiple RNA-seq studies, this ORF appears to be in a separate operon from the ECF genes (data not shown). For complementation of this mutant, the same three genes were placed behind the aphA-3 gene with its own promoter used for construction of the nonessential gene library (Ge & Xu, 2012). These four genes were then inserted into the SSA_0169 gene. Although this disrupts the gene, we have shown previously that this has no effect on growth, IE virulence, or a number of other phenotypes (Senty Turner et al., 2009). To make it possible to use different antibiotics for selection of the ΔECF mutant and the complemented mutant, we performed an exact replacement of the ada9 cassette in the complemented mutant with the tetM cassette including its promoter, encoding tetracycline resistance (Senty Turner et al., 2009). The ΔssaACB mutant (JFP169) was constructed by an exact replacement of the ssaACB genes with the promoterless version of the aphA-3 gene, also used for construction of the nonessential gene library (Ge & Xu, 2012). This strain relies on the strong, native ssaACB promoter for expression of the aphA-3 gene. This strain has been characterized previously (Murgas et al., 2020) (also Puccio, et al, 2020). For construction of the ΔECF-ΔssaACB mutant, the ada99 cassette with its flanking DNA in the ΔECF mutant was amplified by PCR and introduced into the ΔssaACB mutant by transformation. All mutants were confirmed to possess the expected sequences by Sanger sequencing (Retrogen, Inc.).

4.2 | Growth conditions

Unless otherwise stated, strains were grown under microaerobic conditions (6% O₂, 7.2% CO₂, 7.2% H₂, and 79.6% N₂) at 37°C using an Anoxomat® (Spiral Biotech, Norwood, MA) with an Anoxomat® system (Spiral Biotech, Norwood, MA). HS was prepared from frozen serum obtained commercially from anonymous donors (normal pool, MP Biomedicals™, cat No. MP092930149) that was thawed at room temperature and heat inactivated by incubation for 30 min at 56°C. Heat inactivation was performed to eliminate possible differences in complement levels among different serum lots. For the ORF-seq experiments, the mutant library was grown in BHI (Difco Inc., Detroit, MI) or HS for 24 hr. For in vitro growth studies involving individual mutants, strains were grown individually in BHI overnight, and then, diluted 10⁴-fold into BHI, HS, or human saliva obtained commercially from anonymous donors (pooled human donors, LeeBio, Catalog Number: 991-05-P) for 24 hr unless otherwise indicated. Growth in rabbit serum was examined as described previously (Baker et al., 2018). Briefly, cells were grown overnight at 37°C in BHI with 6% of O₂, then, diluted 10⁶-fold into pooled rabbit serum (Gibco) that had been preincubated in 12% of O₂, 12% O₂, 4.3% H₂, 4.3% CO₂, and 80% N₂. Incubation was continued in this atmosphere for 24 hr, then, 1 ml was removed, sonicated, diluted in...
PBS, and plated onto BHI agar for enumeration. Apart from the minute amounts in the inocula, BHI was not added to HS, human saliva, or rabbit serum to support growth. When needed, antibiotics were added at the following concentrations: kanamycin (Km), 500 μg/ml; spectinomycin (Sc), 200 μg/ml; tetracycline (Tc), 5 μg/ml; erythromycin (Em), 10 μg/ml.

### 4.3 ORF-seq

The tag that was linked to the kanamycin resistance gene (aphA-3) used to create each mutant (Xu et al., 2011) served as the basis for determining the abundance of each mutant in the pooled library via ORF-seq. Our method is a modification of that described by Turner et al. (2015), which employed a poly(C) tail. In ORF-seq, the aphA-3 gene is present in every mutant and is thus analogous to the transposon present in every mutant in the Tn-seq procedure. A 5' primer complementary to the 3' end of aphA-3 was used in conjunction with a 3' primer containing a poly-G tail to amplify DNA sequences located adjacent to the aphA-3 gene to identify each mutant. Each mutant in the library was separately cultured, combined, and washed with PBS. The library mixture was then diluted 100-fold into HS or BHI and incubated microaerobically for 24 hr at 37°C. Cells were harvested by centrifugation at 1,700 g for 15 min at 4°C and washed with PBS. Genomic DNA was isolated using a QIAamp DNA Mini Kit (Qiagen, USA). Subsequently, genomic DNA was sheared by sonication using a Covaris S2 Ultrasonicator (Qsonica, USA). PolyC was added to 3' ends of the fragmented library by Terminal Deoxynucleotidyl Transferase (Promega, USA). The DNA fragments with polyC were purified by AMPure XP beads (Beckman, USA) for use as template for the next step of PCR. PCR was performed using primers olj 376 and K10_Truseq (Table S1). The PCR program was as follows: 98°C for 45 s, then, 20 cycles of 98°C for 10 s, 60°C for 30 s, and 72°C for 30 s. This was followed by 72°C for 5 min and a hold at 4°C. The PCR products were purified by AMPure XP beads, following by the second round of PCR using the primers Truseq_HT01 for the BHI set and Truseq_HT02 for the HS set (Table S1). PE1npKan was used in both sets as the 3’ primer (Table S1). The PCR program was as follows: 98°C for 45 s, 10–15 cycles of 98°C for 10 s, 60°C for 30 s, and 72°C for 30 s, followed by 72°C for 5 min and a hold at 4°C. The PCR products were purified by AMPure XP Beads and sent to the VCU DNA Core Facility for NGS sequencing. Each experiment was performed in triplicate.

### 4.4 Metal analysis

Cellular metal content was determined by ICP-OES exactly as described previously (Baker et al., 2018). In brief, cells grown in BHI were harvested, washed in cold, Chelex-treated (Bio-Rad) PBS, and resuspended in the same buffer. An aliquot was removed for lysis and protein determination using a bicinchoninic acid (BCA) protein assay kit (Pierce). The remainder of the sample was centrifuged, resuspended in Chelex-treated deionized water, to which was then added concentrated (67% to 70%) trace metal-grade nitric acid. The mixture was then placed into a modified polytetrafluoroethylene (TFM) digestion vessel in a Multiwave Go microwave digestion system (Anton Paar) for digestion at 20 × 10⁵ Pa. Following dilution in Chelex-treated water, samples were analyzed on an Agilent 5110 ICP-VDV OES with a Pb internal standard (MSPB-10PPM; Inorganic Ventures) against a standard curve made with a CMS-5 metal standard (Inorganic Ventures, Inc.). Each sample was analyzed for Co, Mn, Fe, and Zn. Results were expressed relative to protein amounts determined above.

### 4.5 Virulence studies

Specific-pathogen-free, male New Zealand White rabbits (2–4 kg; Charles River Labs Inc., Wilmington MA) were used to assess virulence as previously described (Crump et al., 2014). Rabbits were sedated and anesthetized before insertion of a catheter into the right carotid artery. The catheter was trimmed and sutured in place to induce minor damage to the aortic valve. The incision site was closed with staples. The following day, all strains were inoculated into BHI for overnight growth in 6% of O₂. The next morning, the cultures were diluted 10-fold into fresh, prewarmed BHI and incubated for 3 hr before harvesting by centrifugation and washing twice in PBS. The washed cells were diluted and combined in PBS to yield an inoculum containing approximately 10⁷ CFU/ml of each strain. Bacterial enumeration of the remaining inoculum was performed by removing 1 ml of the prepared inoculum, sonicating, and plating a dilution series onto replicate BHI agar plates, each of which contained one of the three antibiotics employed in the experiment, using an Eddy Jet 2 spiral plater (Advanced Instruments, Inc., Norwood, MA). The peripheral ear vein of each rabbit was injected with 0.5 ml of the combined inoculum. Rabbits were sacrificed via intravenous injection of Euthasol approximately 20 hr after inoculation. Catheter placement was assessed upon necropsy and cardiac vegetations were collected, and then, placed into PBS and homogenized. The homogenates were sonicated, serially diluted, and plated onto replicate BHI agar plates, each containing one of three antibiotics as per the inoculum, and incubated overnight anaerobically. Resulting colonies were counted to determine CFU/rabbit for each strain, with numbers normalized to account for small differences in inoculum numbers for each strain. All animal experiments were approved by the Institutional Animal Care and Use Committee of the Virginia Commonwealth University, and were performed in compliance with local, state, and federal regulations.

### 4.6 Data analysis

Reads obtained from ORF-seq were aligned against the S. sanguinis SK36 genome using Rockhopper (Tjaden, 2015). Analyses were run using
default parameter settings. Significance was determined by a $p$ value adjusted for a false discovery rate of 1%. Three replicates were examined for data analysis. The set of genes whose mutants had significant abundance changes was input into the STRING database for the construction of a protein–protein interaction network (Szklarczyk et al., 2017). The results of the STRING analysis were imported to the Gephi software program for the visualization and calculation of the betweenness centrality (Bastian et al., 2009). Genes in the STRING network were classified by COG annotation (Tatusov et al., 2000). Genes identified as affecting abundance in HS or BHI were input into the DAVID database for the KEGG pathways enrichment analyses. The significantly enriched KEGG pathways ($p$ value $\leq .05$) were identified. The purine and pyrimidine metabolism pathways were analyzed by mapping related $S.\ sanguinis$ genes onto the KEGG nucleotide transport and metabolism pathway (Kanehisa et al., 2017). The function of transporters in $S.\ sanguinis$ was also classified based on the information obtained from the KEGG database (Kanehisa et al., 2017). The HMDB was used to explore the concentration of compounds in blood and saliva (Wishart et al., 2018). The data of metabolite concentrations in the form of mean ± standard deviation was viewed. Note that samples are labeled as "blood" in this database even if they are actually plasma or serum (Wishart et al., 2018). The growth of individual $S.\ sanguinis$ mutants in HS and saliva was investigated by dilution plating of bacterial cultures. Three replicates were performed to measure means and standard deviations. Significance was assessed using Student’s $t$ test. The growth of strains in rabbit serum and their recovery in the rabbit endocarditis model were assessed by ANOVA and Tukey posttest applied to log-transformed data.

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CONFLICT OF INTERESTS

The authors declare no competing interests.

AUTHOR CONTRIBUTIONS

XG and PX conceived and designed the ORF-seq study. XG carried out ORF-seq assays. BZ, LB, HN, and HG performed HS growth studies. SG designed and performed the metal analysis and rabbit serum growth studies. SG, TP, and TK designed and performed the virulence study. Data analysis was performed by BZ, HG, and SG. BZ, TK, HN, and PX wrote this manuscript with assistance from SG and TP. All authors reviewed and discussed the manuscript.

DATA AVAILABILITY STATEMENT

All data generated or analyzed during this study are included in this published article (and its supplementary files).
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**SUPPORTING INFORMATION**

Additional Supporting Information may be found online in the Supporting Information section.

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