Adherence to host extracellular matrix and serum components by Enterococcus faecium isolates of diverse origin

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Abstract

Enterococcus faecium has emerged as an important cause of nosocomial infections over the last two decades. We recently demonstrated collagen type I (CI) as a common adherence target for some E. faecium isolates and a significant correlation was found to exist between acm-mediated CI adherence and clinical origin. Here, we evaluated 60 diverse E. faecium isolates for their adherence to up to 15 immobilized host extracellular matrix and serum components. Adherence phenotypes were most commonly observed to fibronectin (Fn) (20% of the 60 isolates), fibrinogen (17%) and laminin (Ln) (13%), while only one or two of the isolates adhered to collagen type V (CV), transferrin or lactoferrin and none to the other host components tested. Adherence to Fn and Ln was almost exclusively restricted to clinical isolates, especially the endocarditis-enriched nosocomial genogroup clonal complex 17 (CC17). Thus, the ability to adhere to Fn and Ln, in addition to CI, may have contributed to the emergence and adaptation of E. faecium, in particular CC17, as a nosocomial pathogen.

Introduction

Enterococcus faecium, often found as part of the normal mammalian gut flora, has emerged in the last two decades as a significant nosocomial pathogen with high resistance to multiple antibiotics (Murray, 2000; Leavis et al., 2006). Enterococcus faecium can cause a broad range of serious infections, including bacteremia, surgical wound infections, urinary tract infections, meningitis and endocarditis (Murray & Weinstock, 1999). Recent surveillance data revealed a change in the epidemiology of enterococcal infections: before 1997, infections due to E. faecium were infrequent (~10%) of enterococcal clinical isolates), but now, in the nosocomial environment, E. faecium account for ~38% of healthcare-associated enterococcal infections (Hidron et al., 2008; Top et al., 2008). In addition to the acquisition of resistance to antibiotics (e.g. ampicillin, aminoglycosides and vancomycin), hospital-associated E. faecium strains belonging to clonal complex 17 (CC17) genogroup more often carry genes encoding the enterococcal surface protein (esp) (Willems et al., 2001), a putative hyaluronidase (hyl) (Rice et al., 2003), nine of the 14 fms (E. faecium surface protein-encoding) genes reported by Sillanpää et al. (2009) and a functional copy of the collagen adhesin-encoding gene (acm) (Nallapareddy et al., 2003).

Similar to several other gram-positive pathogens, adherence to exposed host extracellular matrix (ECM) is likely the first step in the infection process of E. faecium. ECM is a complex array of secreted molecules, including collagens and a number of other glycoproteins, proteoglycans and glycosaminoglycans (Iozzo, 1997). Earlier reports on ECM protein adherence of E. faecium, including our previous pilot survey of 13 clinical E. faecium isolates, demonstrated the ability of occasional isolates to adhere to one or more ECM proteins, such as fibronectin (Fn), vitronectin and lactoferrin (Lf) (Zareba et al., 1997; Xiao et al., 1998; Styriak et al., 1999; Styriak & Ljungh, 2003). Because relatively few isolates and adherence targets [except collagen type 1 (CI)] were included in these studies, the prevalence of adherence phenotypes to ECM components and their distribution among different populations of E. faecium, such as isolates associated with human infections vs. nonclinical sources,
remains unclear. Recently, Nallapareddy et al. (2003, 2008) assessed a diverse collection of 122 isolates for acme-mediated adherence to CI and found a significant association between clinical (vs. fecal or food) origin and CI adherence.

In this investigation, we examined the ability of 60 E. faecium isolates, derived from human infections, community feces and animals, to adhere to up to 15 ECM and serum components, including various glycoproteins, proteoglycans and glycosaminoglycans. The possible association between adherence and clinical origin was also analyzed.

Materials and methods

Bacterial strains and growth conditions

A total of 60 E. faecium clinical and nonclinical isolates collected between 1990 and 2006 from diverse geographic locations (the United States, Argentina, China, Germany and Spain) were included in this study. These isolates were derived from the blood cultures of endocarditis patients (n = 20, collected between 1992 and 2006), other clinical infections (OCI group; isolated from the blood of nonecarditis patients, bile, catheters, sputum, urine and wounds; n = 19, collected between 1990 and 2006), healthy volunteer human feces from the community (n = 10, collected between 1994 and 2003), and animals (isolated from either feces or rumen contents of cattle, feces of chickens, turkeys or swine; n = 11, collected between 1998 and 2002). Isolates known to be identical by pulsed-field gel electrophoresis to other isolates in this collection were previously excluded (Sillanpää et al., 2009).

All of the E. faecium isolates were previously identified to the species level by biochemical tests and confirmed by high-stringency colony hybridization using intragenic efu506 and aac(6')-li probes (Singh et al., 1998; Duh et al., 2001). For routine growth, organisms were grown using brain–heart infusion (BHI; Difco Laboratories, Detroit, MI) broth/agar and incubated overnight at 37 °C.

ECM components

The 15 host components used for adherence studies in this study can be classified into five groups. (1) Classic ECM proteins: CI (rat tail, Sigma, St. Louis, MO), collagen type V (CV) (human placenta, Sigma), Fn (human plasma, Enzyme Research Laboratories, South Bend, IN), vitronectin (human placenta, Sigma) and laminin (Ln, natural mouse Ln from EHS cells, Invitrogen, Carlsbad, CA). (2) Plasma proteins: fibrinogen (Fg, human plasma, Enzyme Research Laboratories), Lf (human milk, Sigma), transferrin (Tf, human, Sigma) and plasminogen (human plasma, Enzyme Research Laboratories). (3) Small leucine-rich proteoglycans: decorin (bovine articular cartilage, Sigma) and biglycan (bovine articular cartilage, Sigma). (4) Glycosaminoglycans: heparan sulfate (bovine kidney, Sigma), heparin (porcine intestinal mucosa, Sigma) and hyaluronic acid (human umbilical cord, Sigma). (5) Other: mucin (porcine mucosa, Sigma).

Adherence of radiolabeled cells of E. faecium isolates to immobilized ECM proteins

Enterococcus faecium cells were inoculated at an initial cell density of OD600 nm = 0.01 to 5 mL BHI broth containing 10 μCi mL−1 35S label, and were grown at 37 °C for 8, 10 or 12 h. Adherence of bacteria to immobilized host proteins was tested by an assay described previously with minor modifications (Xiao et al., 1998; Nallapareddy & Murray, 2008). Briefly, 1 μg of CI, CV and Ln in phosphate-buffered saline (PBS), pH 7.4, and Fn and Fg in 50 mM carbonate–bicarbonate buffer, pH 9.6, was immobilized in the wells. Because we have not previously assessed the coating efficiencies of Lf, Tf, decorin, plasminogen, heparin, hyaluronic acid and mucin using this assay, we initially immobilized 1, 5 and 10 μg of each protein in PBS or in the above carbonate–bicarbonate buffer to the wells to test adherence of eight E. faecium isolates. Peak adherence was observed by some of the isolates to Lf and Tf at 5 μg and without further increase at 10 μg, while the remaining isolates were nonadherent to any of the tested proteins at all three concentrations. No adherence to bovine serum albumin (BSA) was observed at these concentrations. Furthermore, coating with host proteins dissolved in the carbonate–bicarbonate buffer did not alter the binding of these test strains. Hence, 5 μg in PBS was used in subsequent experiments for coating these proteins. Adherence was determined from three independent experiments using triplicate wells. CI (1 μg) was included as a positive control (Nallapareddy et al., 2003, 2008) and BSA (5 μg, Sigma) as a negative control. The following formula was used for calculating the percentage of adhered bacteria: (radioactivity of adhered cells/total radioactivity of added cells) × 100.

Isolates were classified as adhering to ECM proteins if ≥5% of total labeled cells bound to wells (Nallapareddy & Murray, 2008).

Sequencing of the purK allele

The allelic profile for purK was determined by sequencing for 25 of the 60 isolates, which had not previously been assessed for the presence of the purK1 allele (a marker for CC17 lineage; Willems et al., 2001; Nallapareddy et al., 2008) using primer sequences available at http://efaecium.mlst.net/misc/info.asp
**Statistical analysis**

The two-tailed Fisher’s exact test was used to determine the statistical significance of the differences observed in adherence to various ECM molecules for different *E. faecium* isolates. *P*-values < 0.05 were considered to be statistically significant.

**Results and discussion**

*Enterococcus faecium* adherence to CI, Fn and Fg at three different growth phases

Previous studies on *E. faecalis* and *Streptococcus bovis* (Nallapareddy & Murray, 2008; Sillanpää *et al*., 2008b) revealed that the ability of bacterial strains to adhere to immobilized CI, Fn and Fg is often affected by their growth stage. To examine whether different growth phases influence the adherence phenotypes of *E. faecium* isolates, we selected three endocarditis strains: TX0016 (ST16; also known as strain DO and for which the genome sequence is available), TX0074 (ST337; non-CC17 strain) and TX0082 (ST17; founder sequence type of CC17), and compared their ability to adhere to CI, Fn and Fg after growth in BHI for 8 h (approximately late exponential phase), 10 h (approximately the time of entry into the stationary phase), and 12 h (2 h after entry into the stationary phase). Consistent with our earlier findings, there was no appreciable binding (< 5% of cells adhered) to CI, Fn, Fg or BSA by TX0016 (Xiao *et al*., 1998; Nallapareddy *et al*., 2003), TX0074 showed a high level of adherence to CI, and TX0082 to CI and Fg (Nallapareddy *et al*., 2006) as well as Fn (Fig. 1). TX0074 and TX0082 showed higher adherence levels to tested proteins after growing for 10 and 12 h compared with 8 h, while binding to BSA by both TX0074 and TX0082 was less at 10 h than at 12 h. Thus, for subsequent analyses, cells harvested at 10 h were used.

**Adherence of 60 *E. faecium* isolates to ECM components**

We initially examined the adherence of eight *E. faecium* endocarditis isolates to all 15 ECM molecules. Because no binding was detected to vitronectin, biglycan and heparan sulfate in this screening assay, we limited our subsequent assays to 11 ECM molecules: CV, Fn, Ln, Fg, Tf, decorin, plasminogen, heparin, hyaluronic acid and mucin. Results of adherence from our first assay with all 60 isolates, using triplicate wells for each isolate, showed no binding by any of the isolates to plasminogen, heparin, hyaluronic acid, decorin and mucin. Results of adherence from our first assay with all 60 isolates, using triplicate wells for each isolate, showed no binding by any of the isolates to plasminogen, heparin, hyaluronic acid, decorin and mucin. Because these proteins, along with vitronectin, biglycan and heparan sulfate above, appeared not to be significant adherence targets for *E. faecium* isolates, we limited our two replicative assays to Fn, Fg, Ln, CV, Tf and Lf, as at least one isolate exhibited adherence to these proteins in our first assay; hence, three independent assays with nine wells in total were performed for Fn, Fg, Ln, CV, Tf and Lf. Despite the lack of adherence to the nine proteins above in this study, we cannot rule out the possibility that adherence to some of these proteins could be elicited under...
other culture conditions, for example in the presence of host-derived material, or in vivo in the host.

With cells grown in BHI for 10 h, 16 of the 60 isolates (27%) were considered adherent (≥5% of bacteria adhered) to one or more ECM molecules, while adherence to BSA was minimal (≤1.4%) for all 60 isolates (Fig. 2). Most of the 16 isolates exhibited adherence to one (six isolates) or two (six isolates) of the six ECM proteins, while only one isolate (TX0082, previously shown to express acm, scm and ebpfm; Nallapareddy et al., 2006; Sillanpää et al., 2008a) was adherent to all six. Adherence to Fn was most commonly observed, with 12 of the 60 isolates (20%) showing >5% adherence and six of these showing adherence levels >10% (Fig. 2). Adherence to Fg and Ln was slightly less common, with 10 of the 60 isolates (17%) exhibiting adherence to Fg and eight (13%) to Ln. Two isolates were considered adherent to Lf; one of these was the only isolate showing adherence to CV and Tf. It is notable that these adherence phenotypes were clustered in some of the isolates, for example all eight isolates exhibiting adherence to Ln, with one exception, were also adherent to Fn and seven of the 10 isolates that adhered to Fg also adhered to Fn. While yet to be investigated, this clustering of positive adherence within certain isolates could be a result of a generalized regulation mechanism, leading to expression of functional adhesins in a subset of the isolates during in vitro growth.

**Comparison of ECM protein adherence among clinical and nonclinical isolates**

Ten of the 20 (50%) endocarditis-derived *E. faecium* isolates and three of the 19 (16%) other clinical isolates showed adherence to one or more ECM molecules, and the percentages of adherent cells among these strains ranged from 5% to 23%. In contrast, only one of 10 stool isolates from community volunteers (10%) adhered to at least one ECM molecule, while two of 11 animal-derived isolates (18%) adhered at a lower level, as shown in Fig. 2. For Fn adherence, a statistically significant difference was noted between the percentages of adherent strains in clinical groups (endocarditis plus OCIs) and nonclinical groups (stool isolates from community volunteers and animal sources) (∗∗P = 0.04); this was driven by the higher percent of endocarditis-derived *E. faecium* isolates showing adherence to Fn compared with nonclinical isolates (stool isolates from community volunteers and animal sources) (∗∗∗P = 0.003) (Table 1). Similarly, there was a statistically significant difference between the percentages of adherent strains in clinical groups vs. nonclinical groups for adherence to Ln (∗∗∗P = 0.04); again, more endocarditis-derived isolates adhered to Ln vs. nonclinical isolates (∗∗∗P = 0.003) (Table 1). The percentage of Fg-adherent strains in clinical groups was found to be not significantly different (18%) from...
nonclinical groups (14%). One of the 20 endocarditis isolates adhered to CV and Tf and two to Lf, while none of the OCI isolates, stool isolates from healthy volunteers or animal sources showed adherence to these three ECM molecules. Taken together, these results indicate that in vitro-grown clinical E. faecium isolates, especially those derived from endocarditis, commonly exhibit adherence to the major protein components of the ECM, Fn, Ln and Fg, as well as CI (Nallapareddy et al., 2003, 2008), while adherence to the other ECM components tested is uncommon.

### Adherence to ECM proteins in the E. faecium genogroup CC17

Our previous analysis of 35 of the 60 isolates studied here for their ancestral relationships identified 18 of them as CC17 and 17 as non-CC17 (Nallapareddy et al., 2008). Because the purK allele has been defined as an epidemic marker of the CC17 lineage by multilocus sequence typing (MLST) studies (Willems et al., 2001), we sequenced the purK alleles of the 25 remaining isolates. Three of the five endocarditis and eight of the 10 OCI isolates, but none of the four normal stool and six animal isolates were found to carry the CC17-specific purK1 allele. Thus, the overall combined results of MLST and purK allele analyses implicated 29 of the 39 clinical isolates and none of the 21 nonclinical isolates as being related to CC17.

For each of the six ECM molecules, a higher percentage of CC17 isolates exhibited adherence (ranging from 3% to 35%) compared with non-CC17 isolates (ranging from 0% to 13%). Statistically significant enrichment among CC17 isolates was detected for adherence to Fn ($P = 0.009$) and to Ln ($P = 0.023$) (Table 2). The percentage of Fg-adherent strains in the CC17 group was found to be not significantly different (21%) from the non-CC17 group (13%). The increased adherence to Fn and Ln among the endocarditis isolates of this study, of which most (17 of 20) were CC17-related, could be a ‘cause,’ i.e. it might contribute to their ability to cause endocarditis, or an ‘effect,’ i.e. it may have been elicited during the presence of the organism in the vegetation for a prolonged period.

### Lack of correlation of the observed adherence phenotypes with the presence of fms genes encoding putative and characterized microbial surface components recognizing adhesive matrix molecules (MSCRAMMs) and pilus family proteins

We recently evaluated the distribution of 14 fms genes encoding putative MSCRAMMs and/or pilus proteins (Hendrickx et al., 2007; Sillanpää et al., 2008a) among 433 isolates, including the 60 isolates of this study (Sillanpää...

### Table 1. Adherence of 60 temporally and geographically diverse Enterococcus faecium isolates to six ECM proteins

<table>
<thead>
<tr>
<th>ECM molecule</th>
<th>Clinical isolates</th>
<th>Nonclinical isolates</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Endo (n = 20)</td>
<td>OCI (n = 19)</td>
</tr>
<tr>
<td></td>
<td>Total (n = 39)</td>
<td></td>
</tr>
<tr>
<td>Fibronectin</td>
<td>9 (45%)</td>
<td>2 (11%)</td>
</tr>
<tr>
<td></td>
<td>11 (28%)</td>
<td></td>
</tr>
<tr>
<td>Fibrinogen</td>
<td>6 (30%)</td>
<td>1 (5%)</td>
</tr>
<tr>
<td></td>
<td>7 (18%)</td>
<td></td>
</tr>
<tr>
<td>Laminin</td>
<td>7 (35%)</td>
<td>1 (5%)</td>
</tr>
<tr>
<td></td>
<td>8 (21%)</td>
<td></td>
</tr>
<tr>
<td>Collagen type V</td>
<td>1 (5%)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>1 (3%)</td>
<td></td>
</tr>
<tr>
<td>Transferrin</td>
<td>1 (5%)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>1 (3%)</td>
<td></td>
</tr>
<tr>
<td>Lactoferrin</td>
<td>2 (10%)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>2 (5%)</td>
<td></td>
</tr>
</tbody>
</table>

*17 of 20 isolates had the CC17-associated purK1 allele.

12 of 19 had purK1.

None had purK1.

$^b P < 0.004$, endocarditis isolates versus nonclinical isolates (stool isolates from community and animal sources).

$^c P = 0.04$, clinical isolates (endocarditis and OCI) versus nonclinical isolates (stool isolates from community and animal sources).

Endo, isolates from E. faecium endocarditis patients; OCI, other clinical isolates from various tissues/infections (see Materials and methods).

### Table 2. Comparison of ECM protein adherence between CC17 and non-CC17 isolates

<table>
<thead>
<tr>
<th>ECM molecule</th>
<th>CC17 (n = 29)</th>
<th>Non-CC17 (n = 31)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fibronectin</td>
<td>10 (35%)</td>
<td>2 (6%)</td>
</tr>
<tr>
<td>Fibrinogen</td>
<td>6 (21%)</td>
<td>4 (13%)</td>
</tr>
<tr>
<td>Laminin</td>
<td>7 (22%)</td>
<td>1 (3%)</td>
</tr>
<tr>
<td>Collagen type V</td>
<td>1 (3%)</td>
<td>0</td>
</tr>
<tr>
<td>Transferrin</td>
<td>1 (3%)</td>
<td>0</td>
</tr>
<tr>
<td>Lactoferrin</td>
<td>2 (7%)</td>
<td>0</td>
</tr>
</tbody>
</table>

$^* P = 0.009$ against the non-CC17 isolates of our study.

$^1 P = 0.023$ against the non-CC17 isolates of our study.
et al., 2009). Here, we tested for a possible association between gene presence and adherence phenotype and found no apparent differences in gene distribution profiles of adherent vs. nonadherent isolates (to individual ECM and serum proteins). However, it is plausible that these genes are expressed at varying levels among different strains, resulting in adherence of some of the isolates but not others, as we recently found with acm (Nallapareddy et al., 2003, 2008), or that they are present as pseudogenes in many of the isolates, as shown for acm and three of the 14 genes, fms15, fms16 and fms19 (Nallapareddy et al., 2003, 2008; Hendrickx et al., 2007; Sillanpää et al., 2008a), or that some of them are functionally redundant, i.e. two or more of these genes code for adhesins that bind to the same ligand.

In summary, we have reported here a systematic evaluation of the ability of a large collection of E. faecium isolates from clinical and nonclinical sources to adhere to various host ECM and serum components and detected adherence to six ECM proteins: Fn, Fg, Ln, CV, Tf and Lf. Our observation of the more frequent adherence to Fn and Ln by clinical isolates, particularly endocarditis isolates, as compared with fecal isolates from community volunteers and animal isolates, and the significant association of these two adherence phenotypes with CC17-related isolates, suggests that these interactions may have contributed to the emergence of E. faecium in nosocomial infections. Our ongoing studies are aimed at determining the specific adhesins involved in E. faecium pathogenesis and their host ligands.

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References


Sillanpää J, Nallapareddy SR, Prakash VP, Qin X, Hook M, Weinstock GM & Murray BE (2008a) Identification and phenotypic characterization of a second collagen adhesin, Scm, and genome-based identification and analysis of 13 other predicted MSCRAMMs, including four distinct pilus loci, in Enterococcus faecium. Microbiology 154: 3199–3211.


