Virulence Genes, Antibiotic Resistance and Capsule Locus Polymorphisms in *Enterococcus faecalis* isolated from Canals of Root-Filled Teeth with Periapical Lesions

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**ABSTRACT**

Frequent isolation of *Enterococcus faecalis* from root canal treated teeth with apical periodontitis, has proposed the role of this organism in endodontic treatment failures. Different factors have been suggested in the pathogenicity of this organism. In this study, 22 *E. faecalis* isolates from canals of root-filled teeth were identified, and phenotypic and genotypic characteristics were investigated. No resistance to vancomycin and gentamicin was noted, and most isolates (91%) were susceptible to ampicillin. Biofilm formation was detected in 73% of the isolates and may be considered as the most important virulence factor involved in the pathogenesis of these isolates.

**Keywords:** Enterococcus faecalis; Apical periodontitis; Virulence factors; Antibiotic resistance; Biofilm

Although *Enterococcus faecalis* is not commonly considered a microorganism of the normal oral microbiota, it is frequently isolated from obturated root canals of teeth associated with chronic periodontitis, which indicates its pathogenic role in chronic endodontic treatment failure. The gastrointestinal and genitourinary tracts, but not the oral cavity, are recognized as typical habitats of this organism, but the origin of enterococcal infections in root canals is unclear. Some evidence points to exogenous sources; however, the role of normal host microflora cannot be ruled out [1].

Different virulence factors, including the enterococcal surface protein (Esp), collagen-binding protein (Ace), endocarditis- and biofilm-associated pili (Ebp), cytolysin (Cyl), gelatinase (GelE), bile salt hydrolase, capsule production, and biofilm formation [2] contribute to the pathogenicity of this organism.

Ace mediates *E. faecalis* adherence to human collagen type IV, mouse laminin and human dentin. The ebp locus, which encodes pili, is well known for its role in endocarditis, but its
Characterization of *E. faecalis* dental isolates

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Role in endodontic diseases is unclear. *E. faecalis* pathogenicity is also reinforced by biofilm formation, which leads to chronic infections, increased antimicrobial resistance, and protection from the host immune response, and endodontic treatment failure [2].

As the knowledge about the characteristics of *E. faecalis* strains isolated from root canals may guide new strategies for treatment of related infections, this study was designed to investigate phenotypic and genotypic characteristics of such isolates. Over a period of 12 months (from May 2016 to May 2017), 70 samples were collected from canals of root-filled teeth (associated with apical periodontitis) of seventy patients seen at the Dentistry School of the Kerman University of Medical Sciences or in private endodontic clinics. The study was approved by the ethics committee of the Bam University of Medical Sciences (Opinion number: MUBAM.REC.1395.3) and was conducted in accordance with the Declaration of Helsinki (2008). Written informed consent was obtained from all patients. All teeth had been root filled (at least one year ago) and apical periodontitis was confirmed based on radiographic evidence. Failure of root-canal treatment was confirmed by an endodontist via clinical and radiographic examinations. Patients who underwent antibiotic therapy within the last three months prior to the study or had systemic diseases were excluded.

The sampling procedure was carried out as follows: After removal of the coronal restorations, resin posts and carious lesions and access cavity preparation, the affected tooth was individually separated from the oral cavity using a rubber dam and disinfected using 5.25% sodium hypochlorite solution inactivated with 5% sodium thiosulfate, in order to avoid interference with the bacterial sampling. A single root canal of each patient was sampled, in order to limit the microbiological assessment to a single ecological environment. In multi-rooted teeth, we sampled the root with the periapical lesion, and if all roots had periapical lesions, the widest canal was selected. The root pulp was removed using Gates-Glidden drills and endodontic files without the use of chemical solvents. Irrigation with sterile saline solution was carried out in order to remove any remaining materials and to moisten the canal prior to sample collection [3]. The sample was then obtained by introducing a sterile paper point into the full length of the canal (as determined with a preoperative radiograph), and kept in place for 60 s. Next, the introduced paper points were transferred to microtubes containing liquid *Streptococcus faecalis* (SF) medium (BBL, Sparks, NV, United States), which were incubated for 48 h at 37°C. After bacterial growth (characterized by color change of the medium), the bacterial pellet was collected, and identification was initially performed using conventional methods [4]. *E. faecalis* identification was confirmed by a polymerase chain reaction (PCR) assay targeting the *ddl* gene [4]. The disk diffusion method was used to test susceptibility to antibiotics. Multidrug-resistance (MDR) was defined as resistance to three or more different classes of antibiotics [5, 6]. Gelatinase production was assessed using the method described by Marra et al. with minor modifications [7]. Biofilm formation was assessed by the microtiter plate assay proposed by Upadhyaya et al. [8], with *Staphylococcus epidermidis* (strain ATCC 35984 / RP62A) being used as the positive control. All the assays were performed in triplicate.

Genomic DNA was extracted using an appropriate extraction kit (Gene All, Seoul, Korea), according to the manufacturer’s instruction. PCR assays were carried out using specific primer pairs for the *esp*, *getE*, *cylA*, *ebp*, *ace*, *cib* and *cps* operons, as previously described (Table 1) [4, 9-11]. The *cps* operon encodes the polysaccharide capsule and has 11 open reading frames (from *cpsA* to *cpsK*) and 3 polymorphisms: CPS 1, comprising *cpsA* and *cpsB*; CPS 2, comprising all 11 genes in the operon; and CPS 5, comprising all genes, except *cpsF*. Only CPS 2 and 5 encode the capsular polysaccharide [9].

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E. faecalis was isolated from 22 out of 70 specimens and all isolates were susceptible to vancomycin, teicoplanin, and linezolid. MDR was detected in 4/22 (18%) isolates. Antimicrobial susceptibility results are shown in Table 2. Gelatinase production was detected in 7/22 (32%) isolates. A total of 18 (~80%) isolates were biofilm producers, among which 16 (89%) produced strong biofilm. Optical density (OD) ≥0.24 was considered indicative of strong biofilm formation [4]. The most prevalent virulence genes were cbh and ace each with 72.7%. No esp-positive isolates were found. The frequency of virulence-associated genes is shown in Table 2. Amplification of the cps operon (Fig.1) revealed that CPS 1 was the most common polymorphism, expressed by 63% of the isolates, while only two isolates (9%) expressed CPS2.

High prevalence of E. faecalis in treated root canals of teeth associated with periodontitis emphasizes the role of this microorganism in endodontic treatment failures. In this study, 22 E. faecalis isolates from root canals associated with apical periodontitis were analyzed in terms of antimicrobial resistance, biofilm formation, gelatinase production, expression of virulence-associated genes, and capsule locus polymorphism.

Inside root canals, E. faecalis is protected against the action of antibiotics and the immune system. Beta-lactam antibiotics, such as ampicillin, are used for treating dental infections. It has been reported that ampicillin resistance is not usual in E. faecalis isolated from patients with periodontitis [12]. This is corroborated by our results showing an ampicillin-susceptibility rate of 91%.

Up to 40% of E. faecalis strains isolated from canals of root-filled teeth associated with apical periodontitis were found to be capsule producers in previous reports [9, 13]. The
A high proportion of the isolates were shown to produce biofilm, and multiple virulence-associated genes were detected in individual isolates. A study by Sun et al. found that all *E. faecalis* strains isolated from patients with chronic periodontitis were able to produce biofilm *in vitro* [15]. Regarding virulence genes, the role of the *esp* gene in the *E. faecalis* adaptation to the root canal environment has been reported [13]. In the present study, none of the isolates expressed this gene, which is in contrast with studies reporting an expression rate of 30% and 61% in strains recovered from treated root canals of teeth with or without apical periodontitis [1, 16]. Cytolysin can cause damage to the dentin and preapical tissues via erythrocyte lysis and destruction of host cells. Virulence determinants such as cytolysin are rare in endodontic isolates and previous reports suggest that cytolysin may have little or no relevance to the pathogenicity of *E. faecalis* in endodontic infections [13]. Herein, 63.6% of the isolates expressed the *cylA* gene, but none of them was hemolytic. Our rate was higher than the 38% rate reported by Sun et al. [15], among which 17% were hemolytic isolates.
This lack of phenotypic/genotypic concordance may suggest the loss of the responsible gene in this operon [10]. Another putative virulence determinant is the gelE gene, which encodes gelatinase. Gelatinase hydrolyzes gelatin and other peptides, leading to direct and indirect damages to the host tissues and is possibly involved in the biofilm formation on dentin [17, 18]. Gelatinase contributes to periodontal inflammation by degradation of specific host proteins and disruption of the periodontal organic matrix [13]. The present study revealed that 13 (31.8%) dental isolates produced gelatinase, which is in accordance with previous studies [1, 15, 16]. Such observations demonstrate that gelatinase can affect the E. faecalis virulence and, consequently, the pathogenesis of apical and marginal periodontitis [15]. This lack of genotypic/phenotypic concordance has been reported in other studies [17]. Expression of the gelE gene is regulated by a quorum-sensing system encoded by the fsr gene cluster. A deletion in the upstream sequence of the fsr gene cluster determines the gelatinase-negative phenotype of some E. faecalis isolates [19].

The fact that most isolates were biofilm producers suggests biofilm formation may be considered as the most important virulence factor involved in the pathogenesis of enterococcal infections and biofilm eradication should be targeted in therapy strategies.

REFERENCES

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