Review Article
Animal Models for Periodontal Disease

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Animal models and cell cultures have contributed new knowledge in biological sciences, including periodontology. Although cultured cells can be used to study physiological processes that occur during the pathogenesis of periodontitis, the complex host response fundamentally responsible for this disease cannot be reproduced in vitro. Among the animal kingdom, rodents, rabbits, pigs, dogs, and nonhuman primates have been used to model human periodontitis, each with advantages and disadvantages. Periodontitis commonly has been induced by placing a bacterial plaque retentive ligature in the gingival sulcus around the molar teeth. In addition, alveolar bone loss has been induced by inoculation or injection of human oral bacteria (e.g., Porphyromonas gingivalis) in different animal models. While animal models have provided a wide range of important data, it is sometimes difficult to determine whether the findings are applicable to humans. In addition, variability in host responses to bacterial infection among individuals contributes significantly to the expression of periodontal diseases. A practical and highly reproducible model that truly mimics the natural pathogenesis of human periodontal disease has yet to be developed.

1. Introduction

1.1. Periodontitis. Periodontitis is a highly prevalent, chronic immunoinflammatory disease of the periodontium that results in progressive loss of gingival tissue, the periodontal ligament, and adjacent supporting alveolar bone [1]. In addition to its significant impact on human health, the annual cost of periodontal therapy is estimated to exceed $14 billion in the USA [2]. Furthermore, periodontitis has been associated with systemic diseases, such as cardiovascular complications [3], rheumatoid arthritis [4], and adverse pregnancy outcomes [5].

Chronic inflammation of the periodontium is initiated by complex subgingival biofilms containing several likely periodontal pathogens. The biofilm generally contains a portion of the gram (−) negative anaerobic commensal microbiota as well as opportunistic pathogens of the oral cavity, including Porphyromonas gingivalis (P. gingivalis) [6]. In response to periodontal pathogens, polymorphonuclear cells (PMNs) release destructive reactive oxygen species (ROS), for example, superoxide, via the respiratory burst [7–9], proteinases, and other factors that can damage host tissues [10–12]. These molecules induce further oxidative damage to gingival tissue, periodontal ligaments, and elicit osteoclastic bone resorption [10, 13–15]. The secreted agents also enhance the production of numerous proinflammatory cytokines that contribute to the disease, including interleukin (IL)-1β, IL-6, and tumor necrosis factor (TNFα), among a broad array of biomolecules that have consistently been reported to be elevated in gingival crevicular fluid (GCF) and tissues of periodontitis patients [16–18], rhesus monkeys [19], and dogs [20]. Levels of these proinflammatory molecules are frequently reduced following periodontal therapy [21, 22].

Because individuals are not equally susceptible to the destructive effects of periodontal infections, periodontitis is not only caused by bacterial infection but also may be associated with host susceptibility [23, 24]. Variability in host responses among individuals contributes significantly to the expression of periodontal diseases [24]. Although human cell cultures were found to be useful models for replicating some aspects of the periodontal disease process at the cellular level, information about the complex host response was not prominent [25]. Thus, research into the host response using animals is critically important in the analysis of periodontal disease and development of improved treatments.
1.2. Animal Models. Animal models have contributed to the generation of new knowledge in biological sciences, including periodontology [19, 20, 26–29]. Periodontal disease can occur naturally or be experimentally induced in animals. Various species have been used to study the pathogenesis of periodontitis and to assess therapeutic modalities against the disease. While animal models have provided a large amount of data, it is sometimes difficult to determine whether the findings are applicable to humans. Thus, a simple and reproducible model that truly mimics human pathogenesis of periodontal disease has yet to be discovered. This paper reviews naturally and experimentally induced animal models used to study different aspects of periodontal diseases.

1.3. Nonhuman Primates. Nonhuman primates have oral structures and teeth similar to those of humans and have naturally occurring dental plaque, calculus, oral microbial pathogens (e.g., *P. gingivalis*), and periodontal disease. In particular, rhesus monkeys (*Macaca mulatta*), cynomolgus monkeys (*Macaca fascicularis*), and baboons (*Papio anubis*) are susceptible to naturally occurring periodontal disease [30]. To accelerate periodontitis, however, plaque-accumulating devices, such as orthodontic elastic ligatures or sutures, are commonly placed apical to the interproximal region around selected molars to promote plaque formation [31]. Ligatures are changed at 1-2-week intervals until periodontal pocket formation is confirmed by probing [32–36]. The use of nonhuman primates was later modified to include inoculation with human pathogens. Cynomolgus monkeys with no previously detectible human pathogen *P. gingivalis* were treated with the organism. About 5 months later, infection by *P. gingivalis* was confirmed and plaque formation leading to bone loss was observed [26].

Although periodontitis in primates most closely resembles the human disease, the expense of and special husbandry requirements for these animals limit their use in periodontal studies. In addition, they are prone to infectious diseases such as tuberculosis [30], which makes them a less practical model for periodontal diseases.

1.4. Miniature Pigs. Miniature pigs have oral and maxillofacial structures similar to those of humans in terms of anatomy, physiology, and disease development [37]. The Minnesota miniature pig (minipig) was developed about 60 years ago [38] and has been used extensively in biomedical research [39]. After the age of 6 months, minipigs usually develop gingivitis, manifested by inflamed gingival tissue, accumulated plaque and calculus, and bleeding when probed [37]. There is infiltration of inflammatory cells in the gingival tissue that results in progression to severe periodontal inflammation at 16 months of age with identical histopathology to that seen in humans. Periodontitis in minipigs is promoted in about 4–8 weeks using ligatures, and in association with bacterial inoculations of *P. gingivalis*, *S. mutans*, and *A. actinomycetemcomitans* [37]. Minipigs can be suitable for periodontal as well as orofacial investigations. However, minipigs are relatively expensive, with husbandry issues and few studies to support their use.

1.5. Dogs. Dogs provide an appropriate model to study naturally occurring gingivitis and periodontitis [20]. In dogs, the subgingival plaque involves predominantly anaerobic gram (−) negative cocci and rods, *P. gingivalis* and *F. nucleatum*, similar to human bacteria [40, 41]. The severity of the disease increases with age and frequently results in loss of tooth. Susceptibility or resistance to periodontal disease in different breeds is mainly dictated by genetic variations [42] rather than the diet [43]. In addition, dogs are used for surgical manipulations, including wound healing and regeneration in periodontal pockets [44].

As a limitation of the natural periodontal diseases, the extent and localization of periodontal lesions are not always synchronized in dogs [45, 46]. In dogs, the complete width of marginal gingiva is also affected rather than only the tissue lateral to the gingival pocket wall. In addition, animal care regulations, including daily companionship, exercise, space, and maintenance, make use of dogs less desirable in periodontal studies.

1.6. Rodent Models. Rodents provide some unique characteristics to evaluate microbial and host responses to complement primate and human periodontal studies. Rodents have only one incisor and 3 molars in each quadrant. Studies using rodents have elicited disease via placement of ligatures in the gingival sulcus around the molar teeth by increasing biofilm accumulation, as well as disrupting the gingival epithelium, enhancing osteoclastogenesis and bone loss [29]. In alternative models, these animals are orally infected with select human pathogens, attempting to document the virulence of these species in rodents [9, 47]. These approaches have also enabled the use of genetically manipulated strains to focus on individual components of the host response and to thereby describe their role in the disease process [48, 49]. More recently, different investigators have used gingival tissue inoculated with chemicals [28, 50], microorganisms [51], or their products [52, 53] to elicit periodontal disease.

1.6.1. Rats. Rats are often used in models of experimental periodontitis because periodontal anatomy in the molar region shares some similarities with that of humans. Furthermore, rats are easy to handle and can be obtained with different genomes and microbial status. There is clear evidence from the literature demonstrating horizontal bone loss in rats infected with *Aggregatibacter (Actinobacillus) actinomycetemcomitans* [54–59] or *P. gingivalis* [56–60]. Periodontitis has been induced in rats by placing a bacterial plaque retentive silk or cotton ligature in the gingival sulcus around the molar teeth [61]. In addition, alveolar bone loss has been induced by the injection of *P. gingivalis* [62].

Rice Rats. The swamp rice rat, or rice rat (*Oryzomys palustris*), is a native American species found wildly in the southern US [63]. These animals are highly susceptible to periodontal disease, beginning as early as 2 weeks of age [64]. The gingival tissues become swollen, with pocket formation, accumulation of debris, and ulceration at about 3 months
Prior to infection, mice were given antibiotics for 5–9 weeks. Rice rats have been used to evaluate the dietary effects in mice with genetic defects in their phagocytes, although their ability to control their indigenous bacteria is compromised by genetic defects in their phagocytes, although the presence of antibiotics prevents the development of the disease [75].

Unlike the chronic process in human periodontitis, mice can be utilized to understand the host-parasite interaction [74]. Young mice are one or two of at least 150 microbial types present in any dental plaque biofilm. However mice can be utilized to assess the virulence of periodontal pathogens, specific pathogen-free female BALB/c mice (10 weeks old) were orally infected with strains of A. actinomycetemcomitans and/or P. gingivalis [70–72]. Prior to infection, mice were given antibiotics (sulfamethoxazole and trimethoprim) in their water for about 10 days to suppress the normal oral microflora. Mice were treated by oral gavage five times at 2-day intervals with one type or an admixture of bacteria resuspended in carboxymethylcellulose to establish the infection. Alveolar bone loss was detected after 10 weeks. It was speculated that P. gingivalis initiated experimental periodontitis, at least in part, by modifying the endogenous subgingival biofilm to acquire enhanced virulence [73]. Mice naturally develop periodontitis starting at about 9 months of age with further increases as a function of age, similar to human periodontitis. This model, however, may not reproduce all aspects of human periodontitis initiation and progression; the bacteria used are one or two of at least 150 microbial types present in any dental plaque biofilm. However mice can be utilized to understand the host-parasite interaction [74]. Young mice also can develop periodontitis caused by their own flora, if their ability to control their indigenous bacteria is compromised by genetic defects in their phagocytes, although the presence of antibiotics prevents the development of the disease [75].

Chemically Induced Mouse Model. An alternative method for inducing inflammation of oral tissues is by using trinitrobenzene sulfonic acid (TNBS) or dextran sulphate sodium (DSS) [28, 50]. These chemicals are often utilized to induce acute (1 cycle) and chronic inflammation (3–5 cycles) in the gut to evaluate progress of inflammatory bowel disease (IBD) [76–79]. TNBS delivered rectally and DSS provided orally elicit gastrointestinal inflammation, linked with the natural microbiota of the murine gut [80–82]. DSS acts to undermine the epithelial barrier and is an immune cell activator, resulting in innate immune damage to the tissues. TNBS appears to function as a hapten to modify autologous proteins and induce a T-cell-mediated response, resulting in autoimmune-like inflammatory responses [83]. In addition, these compounds upregulate ROS to create a reproducible model of IBD [76–83]. Oral delivery of DSS or TNBS for an extended period of 18 weeks resulted in chronic oral mucosal inflammation and alveolar bone loss [26, 50]. Mice treated biweekly with DSS in their diet developed systemic disease manifestations, including diarrhea and colitis and dysregulated hepatic concentrations of antioxidants in a time-dependent manner that correlated with a significant increase in alveolar bone resorption. Mice treated orally with TNBS 2 times/week developed no systemic clinical symptoms [28, 50]. Oral administration of TNBS resulted in a localized action on periodontal tissues with alveolar bone loss observed in both maxilla and mandibles with progression in a time-dependent manner. In contrast, TNBS injection into gingival tissues caused a localized but severe and acute infiltration of inflammatory cells, granuloma formation, and rapid and extensive alveolar bone loss. Implementation of these inflammatory bone resorption models will enable determination of ROS contributions to inflammatory disease lesions in the oral cavity [28, 50]. Mice have 3 molars and 1 rootless incisor in each quadrant (Figure 1) and provide minute amount of gingival tissue. Therefore, relatively large numbers of animals per group are needed.
injected for 3 days into the gums of lower incisors with Abscess model, outbred ICR mice (3–6 weeks old) were and are continually erupting. To induce a gum pocket Rodent incisors have no roots Murine Incisor Abscess Model. The murine back abscess model mimic chronic halitosis caused by microbial infection. and has a limited use in studying gum pocket abscess to gum. This model needs repeated injections of the bacteria staining showed granuloma formation within the inflamed gum. This model needs repeated injections of the bacteria and has a limited use in studying gum pocket abscess to mimic chronic halitosis caused by microbial infection.

Murine Back Abscess Model. The murine back abscess model has been used to investigate the interactions of both oral microbial species and host responses to various oral pathogens as monomicrobial infections leading to soft tissue destruction (e.g., P. ginvialis [85–87] and Treponema denticolata [88]). Mixed infections (P. ginvialis and F. nucleatum [89]; P. ginvialis and A. actinomycetemcomitans [90]) have been shown to result in formation of larger abscess compared to a mono infection [89]. Coinfection with S. constellatus and F. nucleatum caused death in mice, while monoinfection with these organisms was not lethal [91]. In addition, the mouse subcutaneous chamber model has been used to study host bacteria interactions and to determine virulence variations among P. ginvialis strains leading to tissue damage and invasion [92]. Although, the lesions are not located in the oral cavity, this model has some value for examining bacterially induced infections/coinfections that result in soft tissue destruction.

1.7. Other Animals

1.7.1. Horses. Common naturally occurring oral diseases in horses include buccal abrasions, calculus, gingival recession, and periodontal pockets. According a recent equine survey, the prevalence of periodontal pockets and gingival recession is highest in older horses and mostly associated with other dental disorders and tooth loss [93]. Because of their size and husbandry considerations, horses are not a practical model for basic science studies of periodontitis or for testing of potential therapies.

1.7.2. Rabbits. Characterization of the oral microorganisms in rabbits showed numerous pathogenic bacteria, including F. nucleatum, P. heparginolytica, Prevotella spp., P. micros, S. milleri group, A. israelii, and A. haemolyticum, which is somehow consistent with the flora related to periodontal disease in humans [94]. Rabbits have been used for creation of surgically induced periodontal defect and to study periodontal regeneration, but they have been found less suitable for regeneration of periodontal ligament [94, 95].

1.7.3. Ferrets. Ferrets (Mustela putorius) naturally develop calculus and periodontal disease similar to humans [96, 97]. Unlike rodents, calculus formation in ferrets does not depend on the diet and can be scored in live ferrets [96]. Ferrets are a suitable model to study calculus; however, they can easily escape from standard cages and they need special maintenance.

1.7.4. Hamsters. Hamsters have a dental formula similar to that of rats, and they develop experimental periodontitis using ligatures around the molar teeth [98–100]. In addition, hamsters have buccal pouches lined with stratified squamous epithelium that are useful for studying oral carcinoma [101]. The disease development is very similar to rats.

2. Conclusion

Each animal model for periodontal disease has advantages and disadvantages (summarized in Table 1). Several show similarities to human disease. While nonhuman primates are most similar to the human condition, their cost and husbandry issues preclude widespread use for both basic science and therapeutic studies of periodontal disease. Rodents are less expensive and easier to handle; however, they do not reproduce all aspects of human periodontitis.
progression. For example, ligatures and/or seeding with exogenous (human) pathogens comprising only one or two from hundreds of microbes that constitute dental plaque biofilm are often needed to induce disease. In addition, rodents have their unique dental anatomical differences. Nonetheless, rats and mice are useful for understanding some aspects of the host-microbe interaction and therapies.

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References


