



## Dental Drug Delivery System Used In Periodontitis

James Ding\*

Dental Materials Science, Division of Applied Oral Sciences & Community Dental Care, The University of Hong Kong, Pokfulam, Hong Kong

### Abstract

Periodontitis is a chronic inflammatory disease that affects the supporting structures of teeth, including the periodontal ligament, alveolar bone, and gingiva. It is a leading cause of tooth loss worldwide and is associated with various systemic conditions such as diabetes and cardiovascular diseases. The traditional treatment of periodontitis involves mechanical debridement, including scaling and root planing, in addition to the use of systemic or local antibiotics. However, the challenges of drug bioavailability, targeting, and sustained release have led to the development of innovative drug delivery systems (DDS). These systems are designed to improve drug concentration at the site of infection while minimizing systemic side effects. This article reviews the various dental drug delivery systems employed in the management of periodontitis, focusing on the mechanisms of action, materials used, and their clinical efficacy.

**Keywords:** Periodontitis; Dental drug delivery systems; Periodontal disease management; Local drug delivery; Controlled release systems; Drug targeting; Nanoparticles; Biodegradable polymers; Antimicrobial agents

### Introduction

Periodontitis is a prevalent and serious inflammatory disease that affects the gums and structures supporting the teeth. It is primarily caused by bacterial infection, leading to the breakdown of the soft tissue and bone that hold teeth in place. If left untreated, periodontitis can lead to tooth mobility and eventual tooth loss. According to the World Health Organization (WHO), nearly 10% of the global population suffers from severe periodontitis, and its prevalence increases with age. Periodontitis is not only a significant cause of tooth loss but also has been linked to a variety of systemic health issues, including cardiovascular diseases, diabetes, and respiratory conditions. The conventional management of periodontitis includes mechanical debridement, scaling and root planing (SRP), and the use of antibiotics or antimicrobials. While SRP helps to remove the bacterial plaque and calculus from the root surfaces, it often fails to completely eliminate bacteria from deep periodontal pockets. Systemic antibiotics are also commonly used; however, they may not achieve adequate concentrations at the site of infection, and their long-term use can lead to resistance. Local delivery systems, therefore, have become increasingly important as they allow for higher concentrations of therapeutic agents at the site of infection, improving treatment outcomes while minimizing systemic side effects.

This paper focuses on various dental drug delivery systems (DDS) that have been developed for the treatment of periodontitis, highlighting the different approaches, materials used, and clinical effectiveness.

### Pathophysiology of periodontitis

Periodontitis begins with the accumulation of dental plaque—a biofilm containing bacteria—on the tooth surfaces, particularly at the gumline. The presence of bacteria leads to the activation of the body's immune response, resulting in inflammation. The inflammatory response is initially localized but can become systemic if the infection is not controlled. As the disease progresses, it leads to the breakdown of the periodontal ligament and alveolar bone, which provide support for the teeth. This tissue destruction occurs through the action of various inflammatory mediators, including cytokines, prostaglandins, and matrix metalloproteinases (MMPs).

Periodontitis can be classified into various stages, from gingivitis (the earliest form of periodontal disease, characterized by inflammation

of the gums without bone loss) to severe periodontitis (with significant loss of tooth-supporting structures). The clinical manifestations include bleeding gums, tooth mobility, deep periodontal pockets, and gum recession. The primary goal of treatment is to control the bacterial infection, reduce inflammation, and regenerate the lost tissues.

### Conventional treatment approaches for periodontitis

The treatment of periodontitis typically involves both non-surgical and surgical interventions. Non-surgical treatment is the first line of therapy and includes scaling and root planing (SRP), aimed at removing plaque and calculus from the tooth surfaces. However, SRP alone may not always be sufficient to eliminate the bacteria deep within periodontal pockets, leading to recurrence of infection.

Systemic antibiotics, such as amoxicillin and metronidazole, are often used in conjunction with mechanical debridement to control bacterial growth. However, systemic antibiotic therapy is limited by its inability to concentrate at the site of infection, which may result in suboptimal therapeutic effects. Furthermore, prolonged antibiotic use can lead to resistance, adverse effects, and a disruption of the oral microbiota.

As a result, local drug delivery systems have been explored to enhance the effectiveness of treatment by providing direct and sustained release of antimicrobial agents at the site of infection. These systems aim to achieve higher local drug concentrations, reduce systemic side effects, and offer a controlled release over extended periods [1-5].

### Dental drug delivery systems (DDS)

Dental drug delivery systems (DDS) are designed to improve the management of periodontal diseases by providing direct and

**\*Corresponding author:** James Ding, Dental Materials Science, Division of Applied Oral Sciences & Community Dental Care, The University of Hong Kong, Pokfulam, Hong Kong E-mail: jamesding21465@gmail.com

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controlled drug release to the site of infection. The key goals of DDS in periodontitis treatment include:

- **Increased local drug concentration:** DDS allows higher drug concentrations at the infection site compared to systemic delivery, improving treatment efficacy.
- **Sustained release:** DDS can offer a prolonged release of therapeutic agents, ensuring continuous antimicrobial activity over a longer duration.
- **Minimized systemic side effects:** By localizing the drug delivery to the periodontal tissue, DDS minimizes the risk of adverse effects that may arise from systemic drug exposure.
- **Targeted delivery:** DDS can be engineered to specifically target the infected periodontal sites, increasing the specificity and reducing side effects.

Various DDS technologies have been developed to address these goals, including biodegradable polymers, nanoparticles, hydrogels, and liposomes. These DDS platforms are designed to deliver various therapeutic agents, such as antibiotics, antimicrobials, enzymes, and growth factors, directly to the site of infection.

## Types of dental drug delivery systems

### Biodegradable Polymers

Biodegradable polymers are one of the most commonly used materials in DDS. They offer the advantage of releasing the drug over an extended period as they break down within the body. Some of the commonly used biodegradable polymers include poly(lactic acid) (PLA), poly(lactic-co-glycolic acid) (PLGA), and chitosan. These polymers are biocompatible, biodegradable, and have been extensively studied for local drug delivery.

PLGA-based systems are widely used in periodontitis therapy due to their ability to control the release of drugs. These systems can be incorporated into various forms, such as microspheres, films, or scaffolds, and are effective in delivering antibiotics like tetracycline or minocycline directly to the periodontal pocket. Once in place, the polymer degrades over time, releasing the drug in a controlled manner.

### Nanoparticles and nanomaterials

Nanoparticles have gained significant attention in recent years due to their unique properties, such as small size, high surface area, and ability to cross biological barriers. Nanoparticles can be engineered to deliver a variety of drugs, including antibiotics, anti-inflammatory agents, and growth factors. These particles are often designed to target specific cells or tissues and offer sustained release of the drug.

Nanoparticles can be fabricated from a variety of materials, including lipids, polymers, and ceramics. In the treatment of periodontitis, nanoparticles can be used to encapsulate drugs like chlorhexidine, doxycycline, or minocycline, ensuring high local concentrations at the infection site. Additionally, nanoparticles can be surface-modified to enhance their stability and improve their interactions with the periodontal tissue.

### Hydrogels

Hydrogels are water-based polymers that can absorb large amounts of water and maintain their shape and consistency. Hydrogels can be used to create a matrix for drug delivery, as they are highly flexible and can be applied directly to the periodontal tissue. Hydrogels are typically

loaded with antimicrobial agents and can provide both a barrier to protect the tissue and a sustained release of drugs.

One of the advantages of hydrogels is their ability to form in situ, meaning they can be applied as a liquid and will gel at body temperature. This makes them easy to apply to periodontal pockets and ensures good retention of the drug. Hydrogels are also highly biocompatible and can be engineered for controlled release, making them an excellent choice for periodontal therapy.

### Liposomes

Liposomes are spherical vesicles made up of lipid bilayers that can encapsulate both hydrophilic and hydrophobic drugs. They can be used to deliver a wide range of therapeutic agents, including antibiotics, anti-inflammatory drugs, and growth factors. Liposomes are especially advantageous in drug delivery due to their ability to fuse with cell membranes, enhancing drug uptake.

In periodontitis treatment, liposomal formulations have been developed to deliver drugs like tetracycline and clindamycin directly to the periodontal tissue. Liposomes can protect the drug from degradation and provide controlled release, ensuring a sustained therapeutic effect.

## Clinical Efficacy of Dental Drug Delivery Systems

Numerous studies have investigated the clinical efficacy of DDS in the management of periodontitis. The results indicate that local drug delivery systems can significantly improve clinical outcomes when compared to conventional treatments. For example, studies have shown that the use of locally delivered antibiotics such as doxycycline or minocycline in combination with SRP leads to a greater reduction in probing depth and improvement in clinical attachment levels than SRP alone.

The use of biodegradable polymer-based systems, such as those containing tetracycline, has also been shown to reduce periodontal pocket depth and bacterial load effectively. Similarly, nanoparticle-based systems and hydrogels have demonstrated promise in providing sustained antimicrobial effects at the site of infection.

One of the key advantages of these systems is their ability to maintain high drug concentrations in the periodontal pocket for extended periods, thus reducing the frequency of application and the risk of bacterial resistance. Furthermore, the localized delivery of drugs minimizes the systemic side effects associated with oral antibiotics.

## Discussion

The development of DDS for periodontitis has opened new possibilities for more effective and targeted treatment. By addressing the challenges of poor drug bioavailability, rapid drug clearance, and bacterial resistance, DDS technologies offer significant improvements over traditional therapies. Biodegradable polymers, nanoparticles, hydrogels, and liposomes each offer unique advantages in terms of drug release profiles, biocompatibility, and ease of application.

Despite the promising results from clinical studies, there are still several challenges that need to be addressed. These include the optimization of drug release rates, the development of systems that can target specific bacteria or tissues, and the improvement of patient compliance. Additionally, the long-term safety and effectiveness of DDS need to be thoroughly evaluated in larger clinical trials.

Overall, dental drug delivery systems represent a promising

approach for the management of periodontitis, providing a more effective, localized, and controlled treatment modality. As research continues to advance, it is likely that DDS will play an increasingly important role in periodontal therapy, offering improved outcomes for patients and reducing the burden of this common and debilitating disease [6-10].

## Conclusion

Periodontitis remains a major cause of tooth loss and is associated with a wide range of systemic health conditions. While conventional treatments such as scaling and root planing and systemic antibiotics are effective to some extent, they are often insufficient in completely eliminating the infection and promoting tissue regeneration. Local drug delivery systems have emerged as an innovative solution to improve treatment outcomes by offering sustained, targeted drug release directly at the site of infection. Biodegradable polymers, nanoparticles, hydrogels, and liposomes are the primary DDS technologies that have shown promise in the management of periodontitis. These systems improve drug concentration at the infection site, reduce systemic side effects, and offer prolonged release of therapeutic agents. Clinical studies have demonstrated that DDS can significantly improve clinical outcomes, including reductions in periodontal pocket depth and bacterial load.

## Acknowledgment

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## Conflict of Interest

None

## References

1. Kesse-Guyot E, Péneau S, Jeandel C, Hercberg S, Galan P (2011) Thirteen-year prospective study between fish consumption, long-chain n-3 fatty acids intake and cognitive function. *J Nutr Health Aging* 15: 115-120.
2. Appelon K, Woodside JV, Yarnell JWG, Arveiler D, Haas G (2007) Depressed mood and dietary fish intake: Direct relationship or indirect relationship as a result of diet and lifestyle. *J Affect Disord* 104: 217-223.
3. Hakkarainen R, Partonen T, Haukka J, Virtamo J (2005) Is low dietary intake of omega 3 fatty acids associated with depression?. *Am J Psychiatry* 161: 567-569.
4. Schiepers OJG, De Groot RHM (2010) Fish consumption, not fatty acid status, is related to quality of life in a healthy population. *Prostaglandins Leukot Essent Fatty Acids* 83: 31-35.
5. Lansdowne ATG, Provost SC (1998) Vitamin D3 enhances mood in healthy subjects during winter. *Psychopharmacology* 135: 319-323.
6. Therasse P, Eisenhauer EA, Verweij J (2006) RECIST revisited: a review of validation studies on tumour assessment. *Eur J Cancer* 42: 1031-1034.
7. Tuma RS (2006) Sometimes size doesn't matter: reevaluating RECIST and tumor response rate endpoints. *J Natl Cancer Inst*. 98: 1272-1274.
8. Gore ME, Escudier B (2006) Emerging efficacy endpoints for targeted therapies in advanced renal cell carcinoma. *Oncology* 20: 19-24.
9. Hoos A, Parmiani G, Hege K (2007) A clinical development paradigm for cancer vaccines and related biologics. *J Immunother* 30: 1-15.
10. Escudier B, Eisen T, Stadler WM (2007) Sorafenib in advanced clear-cell renal-cell carcinoma. *N Engl J Med* 356: 125-134.