

Journal of Pharmaceutical Research International

33(49B): 166-172, 2021; Article no.JPRI.76444 ISSN: 2456-9119 (Past name: British Journal of Pharmaceutical Research, Past ISSN: 2231-2919, NLM ID: 101631759)

Localised Aggressive Periodontitis – A Review

Milind Wasnik^{1≡*}, Suryakant Kumar¹[∞], Arun Sajjanar^{1#}, Niharika Gahlod^{1≡}, Sneha Khekade^{1≡}, Miranda George^{2≡}, Durga Bhattad^{1†} and Pratima Kolekar^{1†}

¹Department of Pedodontics and Preventive Dentistry, Swargiya Dadasaheb Kalmegh Smruti Dental College and Hospital, Wanadongri, Nagpur, Maharashtra, India. ²Department of Pedodontics and Preventive Dentistry, Pushpagiri College of Dental Sciences, Thiruvalla, India.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/JPRI/2021/v33i49B33353 <u>Editor(s)</u>: (1) Ana Cláudia Coelho, University of Trás-os-Montes and Alto Douro, Portugal. <u>Reviewers</u>: (1) Abdelrahman Magdi Ahmed Mohamed, Zhengzhou University, Egypt. (2) Mohamed Ahmed Mohamed Helal, Al-Azhar University, Egypt. Complete Peer review History: <u>https://www.sdiarticle4.com/review-history/76444</u>

Review Article

Received 04 September 2021 Accepted 08 November 2021 Published 12 November 2021

ABSTRACT

The term "periodontal diseases" includes any inherited or acquired disorders of the tissues that are supporting the teeth i. e Gingiva, Cementum, PDL, and Alveolar bone. The periodontal disease can be either localized or generalized. Localized aggressive periodontitis (LAgP) patients have interproximal attachment loss on at least two permanent first molars and incisors, with attachment loss on no more than two teeth other than first molars and incisors. In children and adolescents LAgP occurs without clinical evidence of systemic disease and it is characterized by the severe loss of alveolar bone around permanent teeth [10]. Most commonly the disease is localized to the permanent first molars and incisors.

Aim: The aim of this review article is explain in details about aggressive periodontitis including different management aspect of the same.

Methods: This paper presents a review of the aggressive periodontitis in children. An electronic search was conducted using Pub Med_®/MEDLINE, and Google search using the terms:

- [#]Professor and HOD;
- [†]PG Student;

[■]Senior Lecturer;

[@]Reader;

^{*}Corresponding author: E-mail: milind.wasnik@sdk-dentalcollege.edu.in;

Periodontium, Localized Aggressive Periodontitis, Children and periodontal health, periodontal health in adolescents, gingival disease in children, periodontal disease in children, gingivitis, periodontitis, gingival disease and its prevalence, periodontal disease and its prevalence.

Keywords: Periodontium; Localized aggressive periodontitis; children.

1. INTRODUCTION

The word periodontium comes from the Greek terms, 'peri-' which means "around," and odons', which means "tooth." Periodontium includes the tissues that surround and support the teeth which included Gingiva, Cementum, Periodontal ligaments, and Alveolar bone [1,2]. The periodontium of the primary dentition differs from that of the permanent dentition in numerous aspects [3]. In case of primary dentition the gingiva appears to be more reddish, vascular, and flabby and it lacks stippling [1,4]. Periodontal ligaments in children are wider and they have less dense fibers [1,3,4]. The alveolar bone in case of primary dentition has less trabecula and calcification, more marrow spaces, and they have greater blood supply and lymphatic drainage [1,3,4]. At the molecular level periodontium of the primary dentition resorbed more easily as it contains more amount of sialoprotein and osteoprotein, which ultimately facilitates the binding of odontoclast [1,5-7].

The periodontal disease can be either localized generalized. Localized or aggressive periodontitis (LAgP) patients have interproximal attachment loss on at least two permanent first molars and incisors, with attachment loss on no more than two teeth other than first molars and incisors. The onset of these diseases in young individuals is often circumpubertal. Some investigators have found that the localized form of aggressive periodontitis appears to be selflimiting, [8] while others recommend that it is not [9].

In children and adolescents LAgP occurs without clinical evidence of systemic disease and it is characterized by the severe loss of alveolar bone around permanent teeth [10]. Most commonly the disease is localized to the permanent first molars and incisors. However, bone loss around the primary teeth can be an early finding in the disease as found by some retrospective data obtained from LAgP patients [11].

The prevalence of LAgP in geographically diverse adolescent populations range from 0.1 to 15% [11-20]. Most reports suggest a low prevalence of 0.2%, but it is distinctly greater in

African American populations of 2.5%.Generally LAgP form a very little supragingival dental plaque or calculus [10,21]. which is in contrast with other investigators who found plaque and calculus at levels similar to other periodontal diseases [22,23]. The most common bacteria of apparent etiologic importance include highly virulent strains of *A.actinomycetemcomitans* in combination with Bacteroides-like species [24-27]. In some populations, *Eubacterium sp.* have been associated with the presence of LAgP [28,29].

In LAgP patients a diversity of functional defects have been reported in neutrophils [30-32]. These include anomalies of chemotaxis, [33-35] [36.37] bactericidal phagocytosis. activity, [38]superoxide production, [39-43] FcallIB expression, [44] (CD16) leukotriene R4 generation, [45,46] and Ca2+-channel and second messenger activation [47-51]. The defect in chemo taxis is thought to be an intrinsic defect by some investigators [52-55] and an induced defect by others [56]. The influence of these functional defects on the susceptibility of individuals to LAgP is unknown, but it is possible that they play a role in the clinical course of disease in some patients. Indeed, in some cases exhibiting phagocyte abnormalities, neutrophil defects may still be present after treatment [57]. Molecular markers of LAgP can include an abnormally low number of chemo attractant receptors [58-61] and an abnormally low amount of another cell surface glycoprotein designated GP-110 [62,63]. Adherence receptors on neutrophils and monocytes, such as LFA-1 and Mac-1, are normal in LAgP patients [62,63].

Treatment methods for aggressive periodontitis are often similar to those used in chronic periodontitis. These include:

- 1. Oral hygiene instructions.
- 2. Reinforcement and evaluation of the patient's plaque control.
- 3. Supragingival and sub gingival scaling and root planning.
- 4. Control of other local factors.
- 5. Occlusal therapy, if necessary.
- 6. Periodontal surgery, if necessary.
- 7. Periodontal maintenance

Aggressive periodontitis may, however, require additional treatments beyond those of chronic periodontitis. A general medical evaluation may rule out underlying systemic diseases. Adjunctive antimicrobial therapy, as well as microbial identification and antibiotic testing, should be considered [5]. The long-term outcome may depend on patient compliance. When the primary teeth are affected, the eruption of the permanent teeth and their attachment Levels should be monitored. Finally, evaluation and counseling of family should be done.

2. METHODS

This paper presents a review of the aggressive periodontitis in children. An electronic search was conducted using Pub Med /MEDLINE and Google search using the terms: Periodontium, Localized Aggressive Periodontitis, Children and periodontal periodontal health. health in adolescents, gingival disease in children. periodontal disease in children, gingivitis, periodontitis, gingival disease and its prevalence, periodontal disease and its prevalence. The articles were selected of past 40 years. From this search, the articles which matched these criteria and keywords were selected for review purpose.

2.1 Management of Periodontal Disease in Pediatric and Adolescent Patients

2.1.1 Plaque control

Plaque control program should be planned depending on the chronological age and manual dexterity [64]. Scrub technique of tooth brushing technique has been found to be most effective in children but the modified Bass technique and can also be taught to adolescents [64,65,66]. Since periodontal lesions predominantly being interdentally, older individuals can be advised use of interdental cleaning aids [67].

2.1.2 Nonsurgical periodontal therapy

Microbial plaque load which is significantly reduced after scaling and root planing brings down the level of inflammatory cytokines which ultimately reduce inflammation, bleeding on probing, and probing depths in LAgP [68,69]. In LAgP, however motile organisms (A. actinomycetemcomitans)invade the soft tissues of the periodontium causing recolonization leading to a poor clinical response to therapy [70]. Scaling and root planing, tongue brushing with 1% chlorhexidine for 1 min, mouth rinsing with 0.2% chlorhexidine for 2 min, and irrigation of periodontal pockets with 1% chlorhexidine was found to result in better response in clinical outcomes in cases of LAgP [71].

2.1.3 Systemic antimicrobial therapy

Α. actinomycetemcomitans can recolonize periodontal pockets after scaling and root planing, therefore use of systemic antimicrobials are indicated in such cases [72]. The antibiotics are commonly administered for a period between 7 and 14 days. The dosage should be adjusted as per the child weight and age. Tetracycline cause tooth discoloration in calcifying teeth and photosensitivity and depress skeletal growth in children therefore they are not recommended in children of <8 years of age. Also it is not recommended for pregnant and lactating women. As per adult doses and individuals weighing 40 kg and more, metronidazole and amoxicillin 250-500 mg each, given TID for 8 days is appropriate for most cases of severe periodontitis [73]. Metronidazole and ciprofloxacin 500 mg each, given bid for 8 days is effective against some enteric Gram-negative rods not affected by the metronidazole-amoxicillin regimen Clindamycin little impact has verv on Α. actinomycetemcomitans [74]. As per adult doses and individuals weighing 40 kg and more, amoxicillin 500 mg and potassium clavulanate 125 mg bid have shown good results in AP owing to potassium clavulanate being an inhibitor of beta-lactamase and thus overcoming the bacterial resistance to amoxicillin [73].

2.2 Surgical Therapy

Surgical therapy is carried out to facilitate more accessibility and also the instrumentation of affected sites to remove tissue which are invaded by A. actinomycetemcomitans. At the same time, osseous defects which results from the disease process can be subjected to regenerative procedures to improve osseous support and also to enhance the prognosis of the teeth. Open flap debridement along with systemic antimicrobial therapy in LAgP has found to significantly reduce probing depths and improve attachment levels in affected teeth. With defect morphology being favorable, Guided Tissue Regeneration has shown good results in the treatment of LAP as compared to when osseous grafting is performed [75-81].

2.3 Maintenance

A periodontal examination needs to be carried out during each visit to monitor the stability of attachment levels and check for inflammation. Radiographic evaluation can be carried out in high risk teeth with de plaquing and occlusion adjustments performed when indicated.

3. CONCLUSION

Aggressive periodontitis is associated with systemic conditions which commonly affect children and adolescents. Moreover incipient form of chronic periodontitis may be more common in children. Early diagnosis is needed for early management in these conditions. Aggressive periodontitis has a characteristic pathophysiology in its onset and progression. This review presents evidence obtained from previous investigations carried out regarding periodontitis in children. It also explains about the detailed Management of Periodontal Disease in Pediatric and Adolescent Patients.

DISCLAIMER

The products used for this research are commonly and predominantly use products in our area of research and country. There is absolutely no conflict of interest between the authors and producers of the products because we do not intend to use these products as an avenue for any litigation but for the advancement of knowledge. Also, the research was not funded by the producing company rather it was funded by personal efforts of the authors.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- 1. Harokopakis-Hajishengallis E. Physiologic root resorption in primary teeth: molecular and histological events. Journal ofOral Science. 2007;49(1):1–12.
- Pihlstrom BL, Michalowicz BS, Johnson NW. Periodontal diseases. The Lancet.. 2005;366(9499):1809–1820.
- 3. Oh TJ, Eber R, Wang HL. Periodontal diseases in the child and adolescent,"

Journal of Clinical Periodontology. 2002;29(5):400–410.

- 4. Pinkham JR, Casamassimo PS, Fields HW, McTigue DJ, Nowak A. Pediatric Dentistry, Elsevier Saunders;2005.
- Davies KR, Schneider GB, Southard et al TE. Deciduous canine and permanent lateral incisor differential root resorption. American Journal of Orthodontics and Dentofacial Orthopedics. 2001;120(4):339– 347.
- Lee A, Schneider G, Finkelstein M, Southard T. Root resorption: the possible role of extracellular matrix proteins. 2004;126(2):173–177.
- 7. Bosshardt DD, Degen T, Lang NP. Sequence of protein expression of bone sialoprotein and osteopontin at the developing interface between repair cementum and dentin in human deciduous teeth.
- Gunsolley JC, Califano JV, Koertge TE, Burmeister JA, Cooper LC, Schenkein HA. Longitudinal assessment of early onset periodontitis. J Periodontol. 1995;66:321-8.
- Brown LJ, Albandar JM, Brunelle JA, Löe H. Early onset periodontitis: Progression of attachment loss during 6 years. J Periodontol 1996;67:968-75.
- 10. Baer PN. The case for periodontosis as a clinical entity. J Periodontol. 1971;42:516-20.
- Sjödin B, Matsson L, Unell L, Egelberg J. Marginal bone loss in the primary dentition of patients with juvenile periodontitis. J Clin Periodontol. 1993;20:32-6.
- 12. Löe H, Brown LJ. Early onset periodontitis in the United States of America. J Periodontol. 1991;62:608-16.
- Boughman JA, Halloran SL, Roulston D, et al. An auto-somal dominant form of juvenile periodontitis: Its locali-zation to chromosome 4 and linkage to dentinogenesis imperfecta and Gc. J Craniofac Genet Dev Biol. 1986;6:341-50.
- Marazita ML, Burmeister JA, Gunsolley JC, Koertge TE, Lake K, Schenkein HA. Evidence for autosomal domi-nant inheritance and race-specific heterogeneity in early onset periodontitis. J Periodontol. 1994;65:623-30.
- Hart TC, Marazita ML, McCanna KM, Schenkein HA, Diehl SR. Reevaluation of the chromosome 4q candidate region for early onset periodontitis. Hum Genet. 1993;91:416-22.

- 16. Saxén L. Juvenile periodontitis. J Clin Periodontol. 1980; 7:1-19.
- 17. Saxén L. Prevalence of juvenile periodontitis in Finland. J Clin Periodontol. 1980;7:177-86.
- 18. Kronauer E, Borsa G, Lang NP. juvenile Prevalence of incipient age periodontitis at 16 years in Switzerland. J Clin Periodontol. 1986; 13:103-8.
- 19. Harley AF, Floyd PD. Prevalence of juvenile periodontitis in schoolchildren in Lagos, Nigeria. Community Dent Oral Epidemiol.1988;16:299-301.
- 20. Neely AL. Prevalence of juvenile periodontitis in a cir-cumpubertal population. J Clin Periodontol. 1992; 19:367-72
- 21. Butler J. A familial pattern of juvenile periodontitis (periodontosis) J Periodontol 1969;40:115-8.
- Albandar JM, Brown LJ, Brunelle JA, Löe H. Gingival state and dental calculus in early-onset periodontitis. J Periodontol. 1996;67:953-9.
- 23. Burmeister JA, Best AM, Palcanis KG, Caine FA, Ranney RR. Localized juvenile periodontitis and generalized severe periodontitis: Clinical findings. J Clin Periodontol. 1984;11:181-92.
- 24. Haraszthy V, Hariharan G, Tinoco E, et al. Evidence for the role of highly leukotoxic Actinobacillus actinomycetemcomitans in the pathogenesis of localized and other forms of early-onset periodontitis J Periodontol. 2000;71:912-22.
- Kornman KS, Robertson PB. Clinical and microbiological evaluation of therapy for juvenile periodontitis. J Perio-dontol. 1985;56:443-6.
- 26. Genco RJ, Zambon JJ, Christersson LA. The origin of periodontal infections. Adv Dent Res. 1988;2:245-59.
- 27. Zambon JJ. Actinobacillus actinomycetemcomitans in hu-man periodontal disease. J Clin Periodontol. 1985;12:1-20.
- Moore WEC, Holdeman LV, Cato EP, et al. Comparative bacteriology of juvenile periodontitis. Infect Immun. 1985;48:507-19.
- 29. Han N, Xiao X, Zhang L, et al. Bacteriological study of juvenile periodontitis in China. J Periodont Res. 1991;26: 409-14.

- 30. Daniel MA, Van Dyke TE. Alterations in phagocyte func-tion and periodontal infection. J Periodontol. 1996;67:1070-5.
- 31. Dennison DK, Van Dyke TE. The acute inflammatory response and the role of phagocytic cells in periodontal health and disease. Periodontol. 1997;14:54-78.
- 32. Van Dyke TE, Lester MA, Shapira L. The role of host response in periodontal disease progression: Implications for future treatment strategies. J Periodontol. 1993;64:792-806.
- Genco RJ, Van Dyke TE, Levine MJ, Nelson RD, Wilson ME. Molecular factors influencing neutrophil defects in periodontal disease. J Dent Res. 1986;65:1379-91.
- 34. Van Dyke TE, Levine MJ, Genco RJ. Neutrophil func-tion and oral disease. J Oral Pathol. 1985;14:95-120.
- 35. Van Dyke TE, Hoop GA. Neutrophil function and oral disease. Crit Rev Oral Biol Med. 1990;1:117-33.
- Van Dyke TE, Zinney W, Winkel K, Taufig A, Offenbacher S, Arnold RR. Neutrophil function in localized juvenile periodontitis: Phagocytosis, superoxide production and specific granule release. J Periodontol. 1986;57:703-8.
- Cogen RB, Roseman JM, Al-Joburi W, et al. Host factors in juvenile periodontitis J Dent Res 1986;65:394-9.
- Kalmar JR, Arnold RR, Van Dyke TE. Direct interaction of Actinobacillus actinomycetemcomitans with normal and defective (LJP) neutrophils. J Periodont Res. 1987;22:179-81.
- Asman B, Bergström K, Wijkaner P, Lockowandt B. Influence of plasma components on luminol-enhanced chemiluminescence from peripheral granulocytes in juve-nile periodontitis. J Clin Periodontol. 1986;13:850-5.
- 40. Åsman B, Bergström K, Wijkander P, Lockowandt B. Peripheral PMN cell activity in relation to treatment of juvenile periodontitis. Scand J Dent Res. 1988;96:418-20.
- Zafiropoulos GG, Flores-de-Jacoby L, Czerch W, Kolb G, Markitzu A, Havemann K. Neutrophil function in pa-tients with localized juvenile periodontitis and rapidly progressive periodontitis. J Biol Buccale. 1988;16:151-6.
- 42. Zafiropoulos GG, Flores-de-Jacoby L, Plate VM, Eckle I, Kolb G. Polymorphonuclear neutrophil

chemilumines-cence in periodontal disease. J Clin Periodontol. 1991;18:634-9.

- 43. Shapira L, Borinski R, Sela MN, Soskolne A. Superoxide formation and chemiluminescence of peripheral polymorphonuclear leukocytes in rapidly progressive periodontitis patients. J Clin Periodontol. 1991;18:44-8.
- 44. Nemoto E, Nakamura M, Shoji S, Horiuchi H. Circulating promyelocytes and low levels of CD16 expression on polymorphonuclear leukocytes accompany early onset periodontitis. Infect Immun. 1997;65:3906-12.
- Offenbacher S, Scott SS, Odle BM, Wilson-Burrows C, Van Dyke TE. Depressed leukotriene B4 chemotactic response of neutrophils from localized juvenile periodontitis patients. J Periodontol. 1987;58:602-6.
- 46. Van Dyke TE, Offenbacher S, Kalmar J, Arnold RR. Neu-trophil defects and host parasite interactions in the patho-genesis of localized juvenile periodontitis. Adv Dent Res. 1988;2:354-8.
- 47. Agarwal S, Reynolds MA, Duckett LD, Suzuki JB. Altered free cytosolic calcium changes and neutrophil chemotaxisin patients with juvenile periodontitis. Adv Dent Res. 1989;24:149-54.
- 48. Daniel MA, McDonald G, Offenbacher S, Van Dyke TE. Defective chemotaxis and calcium response in localized juvenile periodontitis neutrophils. J Periodontol. 1993;64:617-21.
- 49. Hurttia HM, Pelto LM, Leino L. Evidence of an associa-tion between functional abnormalities and defective dia-cyl-glycerol kinase activity in peripheral blood neutrophils from patients with localized juvenile periodontitis. J Periodont Res. 1997;32:401-7.
- Kurihara H, Murayama Y, Warbington ML, Champagne C, Van Dyke TE. Depressed protein kinase C (PKC) activity of neutrophils in localized juvenile periodontitis. Infect Immun. 1993;61:3137-42.
- 51. Leino L, Hurttia H, Peltonen E. Diacylglycerol in peri-pheral blood neutrophils from patients with localized juvenile periodontitis. J Periodont Res. 1994;29:334-8.
- 52. Tyagi SR, Uhlinger DJ, Lambeth JD, Champagne C, Van Dyke TE. Altered diacylglycerol level and metabolism in

localized juvenile periodontitis neutrophils. Infect Immun. 1992;60:2481-7.

- 53. Genco RJ, Van Dyke TE, Levine MJ, Nelson RD, Wilson ME. Molecular factors influencing neutrophil defects in periodontal disease. J Dent Res. 1986;65:1379-91.
- 54. Van Dyke TE, Levine MJ, Genco RJ. Neutrophil func-tion and oral disease. J Oral Pathol. 1985;14:95-120.
- 55. Van Dyke TE, Hoop GA. Neutrophil function and oral disease. Crit Rev Oral Biol Med. 1990;1:117-33.
- Agarwal S, Suzuki J. Altered neutrophil function in local-ized juvenile periodontitis: Intrinsic cellular defect or effect of immune mediators? J Periodont Res. 1991;26:276-8.
- 57. Van Dyke TE, Levine MJ, Genco RJ. Periodontal diseases and neutrophil abnormalities. In: Genco RJ, Mergenhagen SE, eds. Host-Parasite Interactions in Periodontal Diseases.Washington, DC: American Society for Microbiology. 1982;235-45.
- Van Dyke T. The role of neutrophils in host defense to periodontal infections. In: Hamada S, Holt S, McGhee J, eds. Periodontal Disease: Pathogens and Host Immune Responses. Tokyo: Quintessence Publishing Co. 1991;251-61.
- 59. Van Dyke T, Levine M, Tabak L, Genco R. Reduced che-motactic peptide binding in juvenile periodontitis: A model for neutrophil function. Biochem Biophys Res Commun. 1981;100:1278-84.
- 60. Van Dyke T, Levine M, Tabak L, Genco R. Juvenile perio-dontitis as a model for neutrophil function: Reduced binding of complement chemotactic fragment, C5a. J Dent Res. 1983;62:870-2.
- 61. Van Dyke T, Schweinebraten M, Cianciola U, Offenbacher S, Genco R. Neutrophil chemotaxis in families with local-ized juvenile periodontitis. J Periodont Res. 1985;20:503-14.
- Van Dyke TE, Wilson-Burrows C, Offenbacher S, Hensen P. Association of an abnormality of neutrophil chemotaxis in human periodontal disease with a cell surface protein. Infect Immun. 1987;55:2262-7.
- 63. Van Dyke TE, Warbington M, Gardner M, Offenbacher S. Neutrophil surface protein markers as indicators of de-fective chemotaxis in LJP. J Periodontol. 1990;61:180.

- 64. Clerehugh V, Tugnait A. Diagnosis and management of periodontaldiseases in children and adolescents. Periodontol. 2000 2001;26:146–68.
- 65. Rugg-Gunn AJ, Macgregor ID, Edgar WM, Ferguson MW. Toothbrushing behaviour in relation to plaque and gingivitis in adolescent schoolchildren. J Periodontal Res. 1979;14:231-8.
- Sangnes G. Effectiveness of vertical and horizontal toothbrushing techniques in the removal of plaque. II. Comparison of brushing by six-year-old children and their parents. ASDC J Dent Child. 1974;41:119-23.
- Kinane DF. The role of interdental cleaning in effective plaque control: Need for interdental cleaning in primary and secondary prevention.In: Lang NP, Attström R, Löe H, editors. Proceedings of the European Workshop on Mechanical Plaque Control. Chicago: Quintessence. 1998;156-68.
- Doungudomdacha S, Rawlinson A, Walsh 68. TF, Douglas CW. Effect of non- surgical periodontal treatment on clinical parameters and the numbers of Porphyromonas gingivalis. Prevotella intermedia and Actinobacillus actinomycetemcomitans at adult periodontitis sites.J Clin Periodontol. 2001:28:437-45.
- 69. Haffajee AD, Cugini MA, Dibart S, Smith C, Kent RL Jr., Socransky SS. The effect of SRP on the clinical and microbiological parameters of periodontal diseases. J Clin Periodontol. 1997;24:324-34.
- Renvert S, Wikström M, Dahlén G, Slots J, Egelberg J. Effect of root debridement on the elimination of Actinobacillus actinomycetemcomitans and Bacteroides gingivalis from periodontal pockets. J Clin Periodonto.1990;17:345-50.
- Quirynen M, Mongardini C, Pauwels M, Bollen CM, Van Eldere J,van Steenberghe D. One stage full-versus partial-mouth disinfection inthe treatment of chronic adult or generalized early-onset periodontitis.II.

Long-term impact on microbial load. J Periodontol. 1999;70:646-56.

- 72. van Winkelhoff AJ, Tijhof CJ, de Graaff J. Microbiological and clinical results of metronidazole plus amoxicillin therapy in Actinobacillus actinomycetemcomitans associated periodontitis. J Periodontol1992;63:52-7.
- 73. Slots J, Ting M. Systemic antibiotics in the treatment of periodontaldisease. Periodontol. 2000 2002;28:106-76.
- 74. Walker C, Gordon J. The effect of clindamycin on the microbiotaassociated with refractory periodontitis. J Periodontol. 1990;61:692-8.
- 75. Sirirat M, Kasetsuwan J, Jeffcoat MK. Comparison between 2 surgical techniques for the treatment of early-onset periodontitis. J Periodontol. 1996;67:603-7.
- 76. Fritz ME, Alexander SA, Hameroff JA, Zinney WB, Van Dyke TE. Treatment of molar teeth with juvenile periodontitis. J Dent Res.1989;68:883.
- DiBattista P, Bissada NF, Ricchetti PA. Comparative effectiveness of various regenerative modalities for the treatment of localized juvenileperiodontitis. J Periodontol 1995;66:673-8.
- 78. American Academy of Pediatric Dentistry. Classification of periodontal diseases in infants, children, adolescents, and individuals with special health care needs. The Reference Manual of Pediatric Dentistry. Chicago, III.: American Academy of Pediatric Dentistry. 2020;418-32.
- 79. Dr. Divya Bhadran. "Periodontitis in Children And Adolescents. IOSR Journal of Dental and Medical Sciences (IOSR-JDMS). 2017;16(08):19–22.
- 80. Arul Pari et al., Gingival Diseases in Childhood Journal of Clinical and Diagnostic Research. 2014;8(10):ZE01-ZE04
- Arthi Lakshmi, Vishnu Rekha, Ditto Sharmin, Sankar Annamalai and Parisa Norouzi Baghkomeh. "Periodontal diseases in children – A literature review. International Journal of Current Research. 2017;9(05):51268-51275.

© 2021 Wasnik et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history: The peer review history for this paper can be accessed here: https://www.sdiarticle4.com/review-history/76444