

Application of Low Level Laser in Temporomandibular Disorders

Arash Rahimi¹, Sepideh Rabiei², Seyed Masood Mojahedi³, Emad Kosarieh⁴

¹Candidate of Lasers in Dentistry, RWTH University, Aachen, Germany

²Candidate of Community Oral Health, Tehran University of Medical Sciences, Iran

³RWTH University, Aachen, Germany

⁴Candidate of Laser in Dentistry, RWTH University, Aachen, Germany

Abstract:

Introduction: The American Academy of Orofacial pain defined temporomandibular disorders (TMDs) as "a collective term that embraces a number of clinical problems that involve the masticatory muscles, the TMJ (Temporomandibular Joint), and the associated structures". Pain and dysfunctional symptoms or signs such as limitations in opening, asymmetric jaw movements and TMJ sounds are the most common findings. TMD is a prevalent disease that is most common among 20 to 40 year-olds; not a disease of senility. Researches show that about 75% of the population has one sign of TMD, and approximately 33% has at least one TMD symptom. They reported that 3.6% - 7% of the population has severe TMD problems that cause patients to seek treatment. Low level laser is a conservative treatment method that has been introduced in recent years. The purpose of this article is to review the related investigations and introduce the applications of low level lasers in TMD treatment.

Method: Electronic data bases were searched and hand search of published articles and texts was done.

Result and Conclusion: Laser application can be beneficial in different ways for TMD problems. In many cases occlusal adjustment and taking impression for splint is necessary but the pain prevents conventional treatment. Pain relief can be achieved by irradiating the joint and tender points; musculature will be relaxed, and treatment can begin. It has been revealed in recent clinical experiences and clinical studies that for myogenic conditions rather high doses are needed and that the energy density itself is an important factor. The disagreement in result of older studies may be related to this fact.

Keyword: low level laser therapy; temporomandibular disorders; orofacial pain

Please cite this article as follows:

Rahimi A, Rabiei S, Mojahedi SM, Kosarieh E. Application of low level laser in temporomandibular disorders: J Lasers Med Sci 2011;2(4):165-70

*Corresponding Author: Arash Rahimi D.D.S, M.S, MSc candidate of laser in dentistry, RWTH University, AACHEN, GERMANY. Tel: (GER) +49-1639441100 (IR) +98-9125068817, Email: dr.arash_dds@yahoo.com

Temporomandibular Disorders (TMDs)

TMJ dysfunction syndrome is a common TMJ disorder which is also referred to as mandibular pain-dysfunction syndrome, arthrosistemporomandibularis, TMJ arthrosis, myofacial pain

syndrom and Temporomandibular Disorders (TMDs) (1-3). The American Academy of Orofacial pain has defined Temporomandibular disorders (TMDs) as "a collective term that embraces a number of clinical problems that involve the masticatory muscles, the TMJ, and the associated structures" (3-5).

TMDs is a prevalent disease that is most common among 20 to 40 year-olds; not a disease of senility (2,5-8). Researches show that about 75% of the population has one sign of TMD, and approximately 33% has at least one TMD symptom. They reported that 3.6% - 7% of the population has severe TMD problems that cause patients to seek treatment (5,7-9). It is most prevalent among female patients and female patients request treatment more than men (2,5,7,10). Moreover, it is less probable that female patients relieve from TMD symptoms than men (5, 11, 12). Although there are many hypotheses, the underlying reason for this gender difference is not clear (13). The early phase with problems in coordination such as clicking, subluxation and recurrent dislocation is most common in 20 to 30 year-old female patients and the later phase with limitations is most prevalent in 30 to 50 year-olds (2, 10)

Sign and symptoms could be rarely bilateral, but the unilateral ones are common including muscular tenderness, limited motion, and a dull aching pain in the periarticular area often referring to the ear, face, head, neck, and shoulders which exacerbate by function. The first manifestations of the syndrome are usually functional disorders in coordination with the masticatory muscles with symptoms of clicking, without pain, subluxation, or recurrent dislocation. In clinical examination, hyper mobility of the joints, tendency to protrude the mandible or both could be seen during the initial opening movement. In most patients these symptoms would be followed by spasms of the masticatory muscles with pain during movement of the joint, particularly during mastication (2, 14). TMDs can lead to further symptoms such as nonotologicotalgia, dizziness, tinnitus, neck pain, and toothache or contribute to migraine and tension headaches, myofacial pain and many other pain complaints (5, 8).

TMD symptoms show fluctuations related to masticatory muscle tensions, tooth clenching, grinding, and other oral parafunctional habits or psychosocial factors such as worriedness, stress, irritation, frustration, and depression (5).

The most common TMD complaints in patients who come to dental offices for treatment, is functional disorders of masticatory muscles and pain (15-20). The range of mouth opening in these patients is also decreased, making it difficult to have

access to the oral cavity, resulting in complications in the rehabilitative treatment (9, 57, 62).

Thorough 1900s, many different curing methods were identified, and today there exists a large number of conservative therapies, such as anti-inflammatory analgesic medication, occlusal splint, acupuncture, ultrasound, electrotherapy and low level laser to decrease pain in TMD patients (21, 22).

Low Level Laser Therapy (LLLT)

LASER (Light Amplification by Stimulated Emission of Radiation) devices are routinely used in physical therapy (23, 24).

Low level laser therapy (LLLT), also known as low-level therapy, laser biostimulation or soft laser therapy can be used in dentistry for different purposes in soft tissues, hard tissues, and pain reduction.

Although literature for more than thirty years has shown that use of LLLT in health care is beneficial with so many researches on this topic in dentistry, opinions differ because of the different methodologies and doses (25-33-35)

The lasers that are used in these fields, may have visible wavelengths (red), or invisible infrared wavelengths (from 650 to 1,000 nm). Helium-Neon lasers (632.5nm) were used before, but semi-conductor diode types (635 to 650 and 803 to 980 nm) like Gallium-Arsenide (GaAs) lasers and Aluminum-Gallium-Arsenide (AlGaAs) lasers are used nowadays. These near-infrared lasers could be used at high powers for soft tissue surgeries and bactericidal effects, or at low power for effective treatment of pain and reducing of inflammation and tissue repair promotion with greater tissue penetration than visible lasers (33, 35, 36). Since wavelength is the most important factor in any type of phototherapy, it is crucial to choose the effective wave length with desirable effects in living tissue (35). The typical power output for a low level laser device is between 10 – 500 mW, total irradiances at any point are several joules and the spot area ranges from a few up to 10cm². Applications can last from a few seconds up to 15 min, and can be either local or distal. LLLT does not have significant thermal effects on dental tissues and consequent therapeutic effects (23, 24, 35, 37). The penetration of LLL in soft and hard

tissue ranges from 3 mm to up to 15 mm, because the wavelengths which are used for LLLT have poor absorption in water (35).

Low Level Laser Mechanisms of Action

The low level laser therapy has complex mechanisms, but it is essentially related to the absorption of particular visible red and near-infrared wavelength in photoreceptors within sub-cellular components and particularly the electron transport (respiratory) chain within the membranes of mitochondria (35,38,39).

The three main effects of laser light used in LLLT on tissue include: 1- analgesic, 2- biostimulating, 3- anti-inflammatory effects.

Analgesic Effect

LLLT increases metabolism of endorphins, acetylcholine, serotonin, and cortisol, it also changes nerve impulse stimulation and transmission, and reduces stimulation and perception of pain as its result. Laser radiation causes hyperpolarization of the membrane which needs greater stimulation to trigger the cellular action potential. Moreover, the analgesic effect of laser radiation has correlation with the increase of beta endorphin in cerebrospinal fluid and normalization of the telethermographic state of the inflamed tissue (dependent on the infrared effect of the 904nm radiation) (33, 40, 41).

Biostimulating Effect

LLLT rises cell activity and adenosine triphosphate (ATP) production. It also causes increased release of growth factors, cytokines, and accelerates replication mechanisms that result in promotion in cell repair processes and declines the oxidative phase. Laser light can also react with beta growth factors and many different oxygenated molecules (33, 42,43). Cellular oxidative state alters these factors; indeed, the effects of photobiostimulation depend on the pretreatment cell state; and are therefore, unpredictable (33).

Anti-inflammatory Effect

LLLT alters blood flow and induces angiogenesis. Furthermore, it increases lymph drainage and

consequently inflammation decreases. In vivo studies on animal models demonstrate both a microcirculation effect that increases production of blood corpuscles, as well as an efficient anti-edemic action that encourages early drainage of the interstitial fluid because of increased lymphatic peristalsis (33,44). Furthermore, the biochemical effect of the laser light can stimulate the production of vascular endothelial growth factor and the conversion of adenosine-monophosphatase into nitric oxide, which improves vessel growth (33,45).

A number of cell systems are involved in the mechanisms possibly underlying the biostimulation of damaged tissue, including fibroblast, macrophages, lymphocytes, epithelial cell, and endothelia (33,46,47). The primary mechanisms of laser assisted therapy are mainly attributable to the interaction between photons and cell molecules that transform the photonic light energy into biochemical energy, whereas the secondary mechanisms are attributable to chemical changes induced by the primary effects. The photochemical effect occurs when the laser light is emitted at low power for prolonged periods of time, and is dependent upon the wavelength, dose, and mode of operation (33,40,41,48-51).

Low Level Laser Therapy in TMD

Laser therapy has been used for analgesic and anti-inflammatory purposes and for muscle relaxation in medical and dental fields (52,53).

In dentistry, laser is used in patients with TMDs, mainly for referring pain to the masticatory, neck, and shoulder muscles. The pain may be the result of malocclusion or psychological and emotional disorders (53, 54).

LLLT is a noninvasive, reversible therapy without any known side effects. Thus, it is quite proper for TMD treatment (55-57). Several studies have reported the positive effect of LLLT in TMD (57-59). Kulekcioglu et al. showed that, after 15 sessions of LLLT, the TMD of both myogenic and arthrogenic origins causes response to therapy with a significant reduction in pain, improvement in mouth opening and lateral movement, and diminished number of trigger points (57,58,60). Results differ from one study to the other, because there are a large range of treatment parameters in therapy (i.e., wavelength, fluence, intensity,

exposure time, total duration of treatment, etc.) (57,61).

In many cases, occlusal adjustment and taking impression for splint is necessary, but the pain prevents the conventional treatment methods. Pain relief can be achieved by irradiating the joint and tender points, musculature will be relaxed, and treatment can begin.

In a research by Hansson, he evaluated the effects of GaAs laser on TMJ arthritis. He emphasizes that lasers are not an alternative to the conventional treatment, but it helps to fasten healing and reduce inflammation more quickly (66). Bezuur and Hansson also treated a group of 27 patients suffering from long-term TMJ problems with a GaAs laser. The treatment was performed over the joint on five consecutive days. 80% of the 15 patients with arthrogenous pain reported total pain relief. The maximum jaw-opening ability increased during the treatment period and continued to increase during the year that the group was monitored. The group suffering from myogenic problems also improved both in terms of pain and jaw opening ability (67). In another study, Kim et al. divided a group of 36 patients with TMJ problem into three therapy groups. The patients were treated either with bite splints, GaAlAs laser or laser acupuncture. The treatment results were compared after two and four weeks with the status of patients before treatment. The following conclusions were drawn. The patients' subjective problems were declined in both the bite splint and laser treatment groups, but improvement in the laser group was much greater than the bite splint group. Clinically noticeable symptoms showed a significant reduction in all groups, but the group treated with laser light responded faster to treatment than the other groups. In this study, laser treatment was more beneficial than bite splints, while laser acupuncture produced the poorest results (69).

It has been revealed in recent clinical experiences and clinical studies that for myogenic conditions rather high doses are needed and that the energy density itself is an important factor. The disagreement in results of older studies may be related to the above fact (63). In cases of trismus, tender points and muscle attachments should be treated. Usually 6 -10 J per point is a good start, but occasionally higher energies are required (64). Even a more peripheral muscle attachment

should be palpated and irradiated during subsequent treatment. This treatment in TMD patients should not be quitted after pain relief, but should be continued at longer intervals.

Arthritis and arthrosis need lower energies, because they have superficial location. 4-6 J per session is suggested for these problems (63). In a research including 75 cases, Bradly et al. found LLLT as effective as mono-therapy in acute joint pain curing. In more chronic cases, without noticeable bone changes on X-ray, LLLT was used as an adjunct treatment to splints. LLLT is almost as effective as intra-articular steroids for joint pain (65, 68).

Several studies have indicated that additional irradiation of the stellate ganglion in patients with pain is effective and can be recommended for TMD treatment as well (70, 73).

Occlusal adjustment and splint is very helpful in TMD patients. LLLT reduces palpation tenderness and improves microcirculation in the tense muscle. Hence, pain reduction, and consequently increase in mouth opening contributes to impression and occlusal adjustment leading to more rapid success (63).

References

1. Wang K. A report of 22 cases of temporomandibular joint dysfunction syndrome treated with acupuncture and Laser radiation. *J tradit chin Med* 1992; 12(2):116-8.
2. Hertling D, Kessler RM. Management of common musculoskeletal disorders: physical therapy principles and methods. 4th, ed. USA: Lippincott Williams and Wilkins 2006.
3. de leeuw R. Orofacial Pain: Guidelines for Assessment, Diagnosis, and Management. 4th ed. Chicago: Quintessence, 2008. p.131.
4. McNeill C. History and evolution of TMD concept. *Oral sury Med oral pathol oral Radiol Endod* 1997;83:51-60.
5. Wright EF. Manual of temporomandibular disorders. 2nd ed. Wiley- Blackwell; 2010.
6. Schwarts L, ed: Disorders of the temporomandibular joint. Philadelphia WB saunders, 1959; 65-6.
7. Okeson JP. American academy of orofacial pain: guidelines for assessment, diagnosis and management. 4th ed. Chicago: Quintessence; 2008. 131-133, 158p.
8. Friction J. Myogenous temporomandibular disorders: diagnostic and management consideration. *Dent Clin North Am* 2007;51(1): 61-83.
9. Yap AUJ, Ho VCL. Temporomandibular disorders-An

- overview. Singapore Med J 1999;40(3):179-82.
10. Schawartz L, chaves CM. Facial pain and mandibular dysfunction. Philadelphia:WB sanders; 1968.
 11. Egermark I, Carlsson GE, Magnusson T. A 20 year longirudinal study of subjective symptoms of temporomandibular disorders from childhood to adulthood. Acta Odontol Scand 2001;59(1):40-8.
 12. Wanman A. Longitudinal course of symptoms of craniomandibular disorders in men and female patients: A lo-year follow up study of an epidemiologic sample. Acta odontol Scand 1996; 54(6): 337-42.
 13. Shinal RM, Fillingim RB. Overview of orofacial pain: Epidemiology and geder differences in orofacial pain. Dent Clin North AM 2007;51(1):1-18.
 14. Somers N. An approach to the management of Temporomandibular joint dysfunction Aust dent J 1978;23(1):37-41.
 15. Schiffman EL, Friction JR, Haley DP, Shapiro BL. The prevalence and treatment needs of subjects with TMD. Am J Dent Assoc 1990; 120(3):295-303.
 16. McCreary CP, Clark GT, Merril RL, Flack v, Oakley ME. Psychological distress and diagnostic subgroups of TMD patients. Pain 1991;44(1):29-34.
 17. JJeffrey P. Okeson: Management of TMD and occlusion, Mosby, 6th edition.2007.195-220.
 18. Rugh JD, Solberg WK. Oral health status in the United States: temporomandibular disorders. J Dent Educ 1985;49(6):398- 406.
 19. Emshoff R, Bosch R, Pumpel E. Schöning H, Strobl H. Low-level laser therapy for treatment of temporomandibular joint pain: a double-blind and placebo-controlled trial. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 2008;105(4):452-6.
 20. Bjordal JM, Couppe C, Chow RT, Tuner J, Ljunggren EA. A systematic review of low level laser therapy with locastion-specific doses for pain from chronic joint disorders. Aust J Physiother 2003;49:107-16.
 21. Ta LE, Dionne RA. Treatment of painful temporomandibular joints with a cyclooxygenase-2 inhibioter: a randomized placebo-controlled comparison of celecoxib to naproxen. Pain 2004; 111(1-2):13-21.
 22. Marini I, Gatto MR, Bonetti GA. Effects of superpulsed low level laser therapy on temporomandibular joint pain, Clin J pain 2010; 26(7): 611-6.
 23. Tuner J, Hode L laser therapy. Clinical practice and scientific background. Prima books 2002.82-3
 24. Baratto L, Calza L, Capra R, Gallamini M, Giardino L, Giuliani A, et al. Ultra low level laser therapy . laser med sci 2011;26(1):103-12.
 25. Pinheiro AL, Meireles GC, de Barros Vieira AL, Almeida D, Carvalho CM, dos Santos JN. Phototherapy improves healing of cutaneous wounds in nourished and umdernourishedwistar rats. Braz Dent J 2004;15:21-8.
 26. Conlan MJ, Rapley JW, Cobb CM. Biostimulation of wound healing by low-energy laser irradiation-A review. J Clin Periodontol 1996;23(5):492-6.
 27. Mester E. Clinical results of laser stimulation and experimental studies on the mechanism of action. Minerva Med 1981;72(33):2195-9.
 28. Schindl A, Schindl M, Pernerstorfer-Schön H, Schindl L.Low intensity Laser therapy:A review. J Investig Med 2000;48:312-26.
 29. Mendez TM, Pinheiro AL, Pacheco MT, Nascimento PM, Ramalho LM. Dose and wavelength of laser light have influence on the repair of cutaneous wounds. J Clin Laser Med Surg 2004;22:19-25.
 30. Benedicenti A. atlante di laser . villa carcina, Italy:Teamwork, 2005.p 16.
 31. Walsh LJ. The current status of low level laser therapy in dentistry. Part 1. soft tissue applications. Aust Dent J 1997;42(4):247-54.
 32. Mester E, Mester AF, Mester A. The biomedical effects of laser application . laser surg Med 1985;5(1):31-9.
 33. Giovanni Olivi . Laser dentistry A user guid. Qb pub 2011. 155-7.
 34. Walsh L J: the current status of low level laser therapy in dentistry . I. soft tissue applications. Aust Dent J 42:247-254, 1997.
 35. A. Moritz . Oral laser application. QV pub. 2006. 521-535.
 36. Walsh LJ. The current status of laser application in dentistry. Aust Dent J 2003;48(3):146-55.
 37. Sandford MA, Walsh L J: thermal effects during desensitization of teeth with GaAlAs lasers . periodontal 1994:5-30.
 38. Karu T. Photobiology of low- power laser effects. Health Phys. 198;56(5):691-704.
 39. Karu T: photobiology of low- power laser therapy: Harwood Academicpublishers, London 1989: 2:42.
 40. BenedicentiA.Lepossibilitadella laser terapianella- curadellanevralgia del trigemino. Parodontolstomatolnuova 1979;3:21.
 41. Benedicenti A. biostimolazione con laser a semicond uttore:parodontolstomatolnuova 1978;3:49.
 42. KaruTL. Molecular mechanism of therapeutic effect of low intensity laser radiation. Laser life Sci 1988;2:53-74.
 43. Pinheiro AL, Gerbi ME. Photoengineering of bone repair processes. Photomed Laser Surg 2006;24(2):169-78.
 44. Karu TL. Photobiological fundamentals of low power laser therapy. J Quant electron 1987;10:1703-17.
 45. WildenL, Karthein R. Import of radiation phenomena of electrons and therapeutic low level laser in regard to the mitochondrial energy transfer. J Clin Laser Med surg 1998;16(3):159-65.
 46. Vacca RA, Marra E, Passarella S, Petragallo VA, Greco M. Increase in cytosolic and mitochondrial protein synthesis in rat hepatocytes irradiated in vitro be He-Ne laser. J Photochem Photobiol B 1996;34(2-3):197-202.
 47. Moreira ME. Reduction in the mitochondrial energy

- potential by inhibition of electron transport in human vascular smooth muscle cells. *AcadEmerg Med* 2001;8:548.
48. Simunovic Z, Ivankovich AD, Depolo A. Wound healing of animal and human body sport and traffic accident injuries using low-level laser therapy treatment: a randomized clinical study of seventy-four patients with control group. *J Clin laser Med surg* 2000;18(2):67-73.
 49. Kyzer MD, Aly AS, Davidson JM, Reinisch L, Ossoff RH. Sub ablation effects of the KTP laser on wound healing. *Laser surg Med* 1993; 13(1):62-71.
 50. Benedicenti A. atlante di laser . villa carcina, Italy:Teamwork, 2005.
 51. Giovanni Olivi, Fred s, Maria Daniela . Pediatric Laser dentistry.quintessence books. 2010.191-195.
 52. Nunez SC, Garcez AS, Suzuki SS, Ribeiro MS. Management of mouth opening in patients with temporomandibular disorders through low-level laser therapy and transcutaneous electrical neural stimulation. *Photomed Laser Surg*;24(1): 45-49.
 53. Rizzi EC, Issa JP, Dias FJ, Leao JC, Regalo SC, Siessere S, Watanabe IS, Iyomasa MM. Low level laser intensity application in masseter muscle for treatment purpose, *Photomed Laser Surg* 2010;28(Suppl 2):31-5.
 54. Friction JR. The relationship of temporomandibular disorders and fibromyalgia: implication for diagnosis and treatment. *Curr. Pain Headache Rep* 2004;8(5):355-63.
 55. Ferreira DM, Zângaro RA, Villaverde AB, Cury Y, Frigo L, Pico G, Longo I, Barbosa DG. Analgesic effect of He-Ne (632.8 nm) low-level laser therapy on acute inflammatory pain. *Photomed. Laser Surg* 2005;23(2):177-81.
 56. Bayat M, Vasheghani MM, Razavi N, Taheri S, Rakhshan M. Effect of low-level laser therapy on the healing of second-degree burns in rats: a histological and microbiological study.*J Photochem Photobiol B* 2005;78(2):171-7.
 57. Nunez SC, Garcez AS, Suzuki SS, Ribeiro MS. Management of mouth opening in patients with TMD through low level laser therapy and transcutaneous electrical neural stimulation. *Photomed Laser Surg* 2006;24(1):45-9.
 58. Kulekcioglu S, Sivrioglu K, Ozcan O, Parlak M. Effectivness of low-level laser therapy in temporomandibular disorder. *Scand J Rheumatol* 2003; 32(2):114-18.
 59. Ilbuldu E, Cakmak A, Disci R, Aydin R. Comparison of laser, dry needling, and placebo laser treatment in myofascial pain syndrome. *Photomed Laser Surg* 2004; 22(4):306-11.
 60. Bjordal JM, Couppe C, Chow RT. A systematic review of low level laser therapy with location-specific doses for pain from chronic joint disorders. *Aust J Physiother* 2003;49(3):107-16.
 61. Núñez SC, Nogueira GE, Ribeiro MS, Garcez AS, Lage-Marques JL. He-Ne laser effects on blood microcirculation during wound healing: a method of in vivo study through laser Doppler flowmetry. *Lasers Surg Med* 2004;35(5):363-8.
 62. Walker N, Bohannon RW, Cameron D. Discriminant validity of temporomandibular joint range of motion measurements obtained with a ruler.*J Orthop Sports Phys Ther* 2000;30(8):484-92.
 63. Tuner J. The new laser therapy hand book . prima book 2010:270-300.
 64. Tasaki . Application of low power laser therapy in closed lock TMD. *Laser surg Med* 1992;suppl 4:84.
 65. Bradley PF. Low intensity laser therapy for TMD. *Laser therapy* 1996;8(1):47.
 66. Hansson TL. Infrared laser in the treatment of craniomandibular disorders, arthrogenous pain., *J Prosthet Dent*;61(5):614-7.
 67. Bezuur NJ, Habets LL, Hansson TL. The effect of therapeutic laser treatment in patients with craniomandibular disorders. *J Craniomandib Disord* 1988;2(2):83-6.
 68. Badly P. Low intensity laser therapy for TMD. *Laser therapy* 1996;8((1):47.
 69. Kim,Ki-suk. Comparative study of the clinical effects of splint, laser acupuncture and laser therapy for TMD. *J dental college, seoul Nat Univ* 1988:1(12):195.
 70. Hashimoto T. Efficacy of laser irradiation on the area near the stellate ganglion is dose-dependent. *Laser therapy* 1997;9(1):7-12.
 71. Kemmotsu O. Laser therapy for pain attenuation. *Proc.2nd cong. World assn for laser therapy, Kansas city* 1998;p.7-8.
 72. Tsushima T. Effects of two-point linear polarized near-infrared irradiation in difficult TMD. *Proc. 2nd cong. World assn for laser therapy, Kansascity, 1998; p. 20-30.*
 73. Tamagawa S. Severe intractable facial pain attenuated by a combination of infrared diode low reactive level laser therapy and stellat ganglion block. *Laser therapy* 1996;8(2):158-9.