

# 1 Photobiomodulation in Dentistry: Current Evidence and Future Directions

## 2

### 3 Abstract

4 Photobiomodulation (PBM), also known as low-level laser therapy, is a non-invasive therapeutic  
5 approach increasingly utilized in dentistry for pain reduction, inflammation control, and tissue healing.  
6 Recent advancements in laser technology have broadened its applications across multiple dental  
7 specialties, including periodontics, orthodontics, implantology, and oral surgery, with high patient  
8 acceptance due to its non-invasive nature. This review synthesizes current evidence on the biological  
9 mechanisms and clinical applications of PBM in dental practice. A comprehensive literature review was  
10 conducted to examine the cellular and molecular effects of PBM, alongside its clinical outcomes in pain  
11 management, bone healing, soft tissue regeneration, and tooth sensitivity management. Findings  
12 indicate that PBM effectively alleviates postoperative and procedural pain, modulates inflammatory  
13 responses, and promotes wound healing, thereby enhancing patient comfort. At the cellular level, these  
14 effects are mediated by mitochondrial activation, increased adenosine triphosphate production, and  
15 regulation of inflammatory mediators. PBM also supports bone regeneration and osseointegration by  
16 stimulating cellular proliferation, differentiation, and the activation of redox-sensitive transcription factors  
17 involved in osteogenesis. Despite its demonstrated clinical benefits, the absence of standardized  
18 treatment protocols limits widespread routine implementation. Future well-designed clinical studies are  
19 necessary to optimize PBM parameters and confirm its long-term efficacy in dental practice. Overall,  
20 photobiomodulation represents a promising adjunctive therapy that complements conventional dental  
21 treatments and contributes to improved patient outcomes.

## 22

### 23 Keywords

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25 Photobiomodulation, Low-level laser therapy, Dentistry, Pain management, Inflammation, Tissue  
26 healing, Bone regeneration, Soft tissue regeneration, Tooth sensitivity, Mitochondrial activation,  
27 Osteogenesis, Clinical outcomes, Osseointegration, Redox signaling, Wound healing

## 28 1. Introduction

29 The therapeutic use of light has been documented since ancient times and advanced markedly with the  
30 development of laser technology. The development of high power lasers transformed multiple medical  
31 and surgical fields, while improvements in light-based devices enabled non-surgical treatments using  
32 controlled wavelength and doses. One innovative approach is LLT, also known as low-intensity laser  
33 therapy, low-power laser therapy, Photobiostimulation and Photobiomodulation. Photobiomodulation is  
34 a non-invasive treatment that uses specific light wavelengths between 650 and 1000nm to promote  
35 tissue repair, reduce inflammation and relieve pain.<sup>1</sup> In dentistry, photobiomodulation is applied at the  
36 cellular level to stimulate differentiation, enhance alveolar bone replication, promote soft tissues  
37 regeneration and reduce postoperative pain, thereby improving periodontal treatment efficacy and  
38 patient comfort.<sup>2</sup> Studies also show it can relieve orthodontic pain and accelerate tooth movement,

39 potentially shortening treatment time. It also promotes bone healing around extraction sites and dental  
40 implants by enhancing blood flow and stimulating bone formation.<sup>3,4</sup> In the field of  
41 dentistry, Temporomadibular disorders commonly present with pain that affects approximately 10% of  
42 adults and impairs quality of life along with physical therapy, numerous studies have provided scientific  
43 evidence supporting the use of laser therapy for managing these conditions.<sup>5</sup> Most research has focused  
44 on pain reduction and improvements in the mandibular movement. The analgesic and anti-inflammatory  
45 effects of low level laser therapy, can reduce pain and muscle sensitivity while improving muscle  
46 performance. When combined with speech-language-hearing therapy, it may enhance mandibular  
47 movement, improve chewing function and promote balance within the stomatognathic  
48 system.[6] Scientific interest in photobiomodulation in dentistry has increased with advances in laser  
49 technology, particularly for promoting oral tissues healing, pain control and adjunctive therapeutic  
50 procedures. Clinical outcomes demonstrate high patient acceptance due to its noninvasive, atraumatic  
51 nature and lack of adverse effects. These advantages, together with rapid recovery and minimal patient  
52 cooperation requirements, have supported its expanding use across dental specialties, including in  
53 pediatric and special-needs populations.<sup>7</sup>

## 54 2. Mechanisms of Action

55 Photobiomodulation (PBM) also known as Low Level Laser Therapy (LLLT) is a non-invasive way to  
56 reduce inflammation and pain thus contributing to enhanced tissue repair. This is achieved by the  
57 modulation of cells and tissues to enhance stem cell differentiation and induce cell proliferation.

58 Photobiomodulation (PBM) uses class III, low-level lasers with 500mW output power. Laser and LED  
59 light stimulates the cell membrane and mitochondrial photoreceptors synthesis of ATP.<sup>8</sup>

60 Several theories on mechanisms of action of Photobiomodulation have been proposed:

- 61 1. This theory proposes the photochemical interaction with the target cell, the photons from the  
62 PBM are absorbed by the chromophores of the mitochondria. Unit IV of the mitochondrial  
63 respiratory chain hosts enzyme cytochrome c oxidase which is the primary chromophore which  
64 absorbs red light resulting in activity of different signaling molecules like reactive oxygen species  
65 (ROS), adenosine triphosphate (ATP), nitric oxide, calcium ions and many others. The enzyme  
66 cytochrome c oxidase carries the electrons from the higher energy orbits to the ultimate electron  
67 acceptor. This leads to creation of a proton gradient, which increases production of ATP.<sup>8</sup>
- 68 2. This theory introduces the direct or indirect photon radiation effects on the genome pool and  
69 DNA. PBM produces ROS indirectly, by low laser radiation. ROS is a free radical, and is  
70 cytotoxic in high levels, but in low levels, it is beneficial in healing tissues and relieving pain. ROS  
71 production is regulated by the mitochondrial membrane potential which is kept in check by the  
72 Cyclooxygenase enzymes. ROS helps in proliferation and differentiation of stem cells. ROS with  
73 cytokines and growth factors helps in healing the damaged tissues after Low level Laser therapy.  
74 This is achieved by moving the satellite cells to the site of injury.<sup>8</sup>
- 75 3. This theory is about the light and heat-gated channels. It suggests that there is a heightened  
76 activity of the plasma membrane in the red light spectrum (600-810 nm) of laser irradiation,  
77 When the wavelength is high and the cytochrome c oxidase is out of range (980-1064 nm).

83 During the low level radiation, the chromophore is activated when NO is displaced and then  
84 bound to the chromophore at the copper and heme centers. The activated chromophore might  
85 stop the oxygen supply to cytochrome c oxidase which was producing a large number of ATPs  
86 due to increased mitochondrial activity, leading to the shift to oxidative phosphorylation from  
87 glycolysis. This leads to increased differentiation of stem cells promoting osteogenesis. ROS  
88 produced during PBM also contributes to the cell differentiation. The gated ion channels which  
89 are sensitive to light allow  $\text{Ca}^{2+}$  to react with NO, cAMP and ROS resulting in activation of  
90 transcription factors.<sup>8</sup>

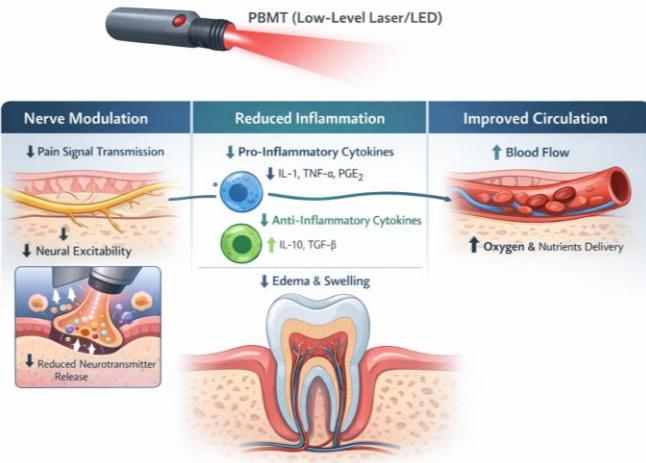
91 Activation of redox-sensitive transcription factors are triggered by ROS. Factors like NF- $\kappa$ B influence  
92 inflammation and bone remodeling by regulating the associated genes. PBM affects the receptor  
93 activator of NF- $\kappa$ B Ligand (RANKL) and Osteoprotegerin (OPG) ratio influencing osteogenesis. This  
94 ratio was seen elevated with 780 nm wavelength of light in cells derived from human alveolar bone.<sup>9</sup>

### 01 **3. Clinical Applications in Dentistry**

#### 02 **a. Pain Reduction and Analgesia**

03 Photobiomodulation therapy (PBMT) is a rapidly growing non-invasive modality for pain management in  
04 dentistry. It involves the use of low-level lasers or light-emitting diode (LED) devices to deliver precise  
05 wavelengths of light to target tissues. PBMT modulates nerve function, reduces inflammation, and  
06 improves local blood flow, thereby decreasing nociceptive signaling and enhancing patient comfort.<sup>10</sup>

07 Clinical evidence shows that PBMT effectively reduces pain perception during routine dental procedures  
08 such as local anesthetic injections, scaling, and minor oral surgeries. Patients report lower discomfort  
09 scores, faster recovery, and reduced need for pharmacologic analgesics. Postoperative pain reduction is  
10 attributed to decreased pro-inflammatory cytokines, enhanced ATP production in cells, and improved  
11 microcirculation at the surgical site.<sup>11</sup> Figure 1.



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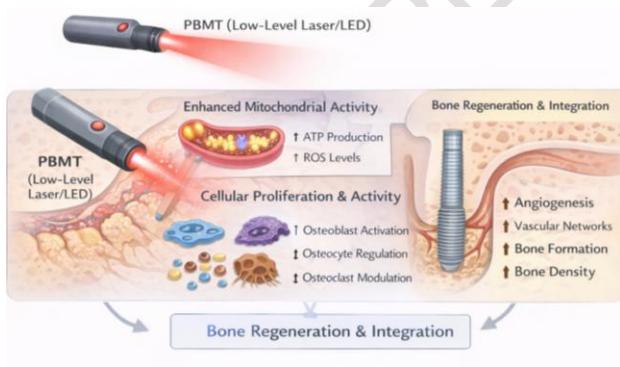
13 *Figure 1. Mechanisms of photobiomodulation-induced analgesia in dental tissues.*

14

15 Furthermore, PBMT has been evaluated for the management of temporomandibular joint (TMJ)  
 16 disorders and oral mucositis, showing promising results in reducing chronic pain and inflammation while  
 17 promoting tissue repair. Standardization of parameters, including wavelength, dose, and exposure time,  
 18 is critical to ensure reproducible analgesic outcomes.<sup>10,11</sup>

### 19 **b. Bone Healing and Osseointegration**

20 PBMT significantly contributes to bone regeneration and implant osseointegration by stimulating  
 21 osteoblastic proliferation, enhancing angiogenesis, and increasing mitochondrial activity in bone cells.<sup>12</sup>  
 22 Clinical studies report accelerated early-stage implant stabilization and improved bone density in peri-  
 23 implant regions, indicating faster and more effective osseointegration.<sup>13</sup> Figure 2.



24

25 *Figure 2. Effect of photobiomodulation on cellular and molecular pathways involved in bone healing and  
 26 osseointegration.*

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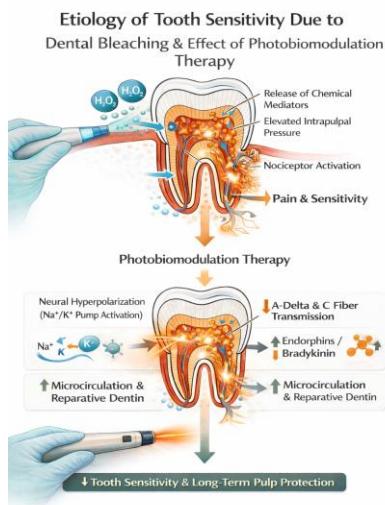
28 PBMT also promotes the healing of periapical lesions after endodontic therapy, supporting bone  
29 regeneration and reducing inflammatory response.<sup>14</sup> Radiographic studies confirm increased bone fill  
30 and improved tissue quality in PBMT-treated sites compared to control groups. When combined with  
31 guided bone regeneration (GBR) techniques and biomaterials, PBMT enhances osteoconductive and  
32 osteoinductive effects, facilitating graft integration and mineralization.<sup>15</sup> The clinical application of PBMT  
33 in regenerative dentistry extends to dental implantology, periodontal therapy, and maxillofacial  
34 reconstruction. Optimization of treatment parameters—such as wavelength (typically 600–1000 nm),  
35 energy density, irradiation time, and frequency—is essential to maximize therapeutic benefits while  
36 avoiding tissue overstimulation.

37 Emerging research explores the synergistic effects of PBMT with stem cell therapy, growth factors, and  
38 bioactive scaffolds, opening new avenues in tissue engineering and regenerative dental  
39 medicine. Standardized clinical protocols and large-scale randomized controlled trials are required to  
40 establish robust evidence for PBMT's efficacy and long-term outcomes in dental pain management and  
41 bone regeneration.<sup>15</sup>

#### 42 **c. Tooth Sensitivity**

43 Tooth sensitivity refers to short, transient and sharp pain response to various factors such as  
44 mechanical factors like attrition and abrasion, chemical factors like erosion and biological factors  
45 causing gingival recession classified as non-carious cervical lesions of multifactorial etiology.<sup>16</sup>

46 Apart from that, tooth sensitivity is common adverse effect associated with Dental bleaching procedures,  
47 primarily linked to diffusion of hydrogen peroxide through enamel and dentin into pulp leading to acute  
48 and transient inflammatory responses like nerve irritation and pulpal inflammation due to release of  
49 chemical mediators altering the microcirculation, elevate pressure on peripheral nerve fibers and  
50 stimulate nociceptors.<sup>17</sup>



51

52 *Figure 3. Etiology of Tooth Sensitivity Due to Dental Bleaching & Effect of PBM Therapy*

53 Clinical studies explained different mechanisms of action for treating tooth sensitivity with the use of  
54 LLLT such as Neural modulation, Biochemical analgesia, Anti-inflammatory & vascular effects, and  
55 Regenerative effectsFigure 3.<sup>16</sup> Clinical application of laser therapy for management of dentin  
56 hypersensitivity requires certain parameters to avoid dentin erosion and thermal damage, previous  
57 studies have demonstrated power setting above 0.75 W may cause surface charring. However, an  
58 ER,CR:YSGG laser operated at 0.25 W ensures safety and efficacy.<sup>18</sup> Beyond its established  
59 therapeutic role, PBM has been proposed as prophylactically to minimize sensitivity associated with  
60 dental bleaching.

61 **d. Preventive Use of PBM Prior to Bleaching Procedures**

62 While most available studies mostly focused on management of post-bleaching hypersensitivity, the use  
63 of PBM as prophylactic before bleaching treatment was proposed as a preventive approach especially  
64 with in-office bleaching. The findings suggest that PBM-induced pulp biostimulation is characterized by  
65 increased odontoblastic activity and decreased A $\delta$  and C nerve fibers excitability. This approach  
66 highlights PBM's potential to improve patient comfort during aesthetic dental treatments.<sup>19</sup>

67 Although many studies illustrate the positive role of PBM in reducing dentin sensitivity, treatment efficacy  
68 depends on various parameters such as site, wavelength, power density, absorption rate and exposure  
69 time.

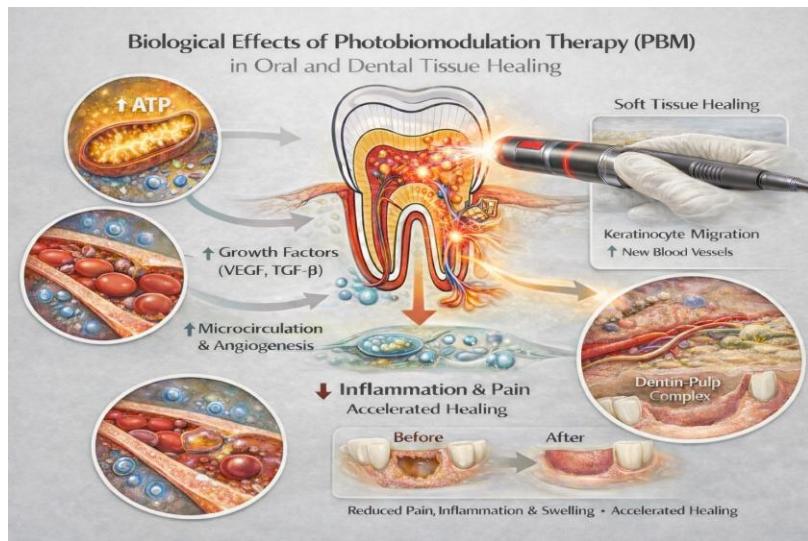
70 Despite these encouraging findings, inconsistencies in laser parameters and treatment protocols across  
71 studies require further high-quality randomized clinical trials to establish clinical guidelines.

72 **e. Tissue regeneration and Healing**

73 Photobiomodulation (PBM), also known as low-level laser therapy (LLLT), is a non-invasive therapeutic  
74 approach that enhances tissue regeneration and wound healing regulating cellular and molecular  
75 processes rather than producing thermal effects [Figure 4]. When delivered within optimal therapeutic  
76 wavelengths, PBM initiates photochemical reactions that activate biological processes without causing  
77 structural damage to tissues which leads to regulated inflammation control, enhanced extracellular  
78 matrix formation, and improved cellular survival during wound healing.<sup>20,22</sup>

79 **f. PBM in soft tissue healing and Stem cell modulation**

80 Furthermore, the use of PBM in periodontal and mucogingival therapy is supported by its ability to  
81 promote soft tissue healing, reflecting clinical outcomes including reduced postoperative inflammation,  
82 decreasing pain level and improving esthetic outcomes.<sup>22,23</sup> [Figure 4]. Beyond soft tissue healing, PBM  
83 has also attracted significant interest in regenerative endodontics, PBM has demonstrated great  
84 potential in stem cells modulation and intracellular signaling in regeneration of pulp-dentin complexes.  
85 Lower doses of low-level laser therapy are associated with more efficacy in enhancing stem cell  
86 proliferation .<sup>21</sup> Figure 4.



88 *Figure 4. Biological Actions of PBM in Dental Tissues*

90 **g. Management of Alveolar Osteitis using LLLT**

91 PBM also showed great results in management of alveolar osteitis. Alveolar osteitis, also known as Dry  
 92 socket, is a post-operative complication marked by intense pain, dislodgement of blood clot and delayed  
 93 wound healing of the socket. LLLT enhances wound healing by promoting immune cell migration,  
 94 stimulating soft tissue repair and increasing blood flow. PBM shows greater outcomes compared to  
 95 conventional treatments such as Alveogyl during follow-up.<sup>24</sup>[Figure 2].

96 Despite substantial evidence supporting the regenerative benefits of PBM in dental application, its  
 97 clinical implementation remains restricted due to various challenges. The most critical limitation is the  
 98 absence of standardized treatment protocols and guidelines along with variability in laser wavelength  
 99 and parameters, which compromises outcome reliability and results comparability.

200 Future research is needed to establish evidence-based protocols and parameters conducting high-  
 201 quality randomized controlled trials to confirm long term safety and efficacy.

206 **4. Upcoming Trends and Future Directions**

208 **a. Home-Based Photobiomodulation Devices**

209 Many photobiomodulation (PBM) devices are currently available on the market for home use,  
 210 particularly for the treatment of inflammation, pain relief, recurrent herpetic infections, and as  
 211 supportive therapy following clinical procedures. The availability of these devices may add value

212 to conventional clinical tools by extending therapy beyond the dental office. However, further  
213 research is still required to establish robust evidence-based databases that address dosimetric  
214 accuracy, patient compliance, and safety monitoring, as variability in wavelength, irradiance, and  
215 exposure time remains a significant concern.<sup>25</sup>

216

217 **b. PBM Combined with Platelet Concentrates (PRP/PRF)**

218 Platelet-rich plasma (PRP) and platelet-rich fibrin (PRF) provide concentrated growth factors that  
219 enhance tissue healing. The use of PBM in combination with platelet concentrates may further  
220 stimulate growth factor activity, cellular proliferation, and tissue remodeling when applied  
221 alongside autologous biologics. Future research is expected to focus on optimizing combined  
222 treatment protocols and identifying clinical scenarios in which this synergistic approach offers the  
223 greatest therapeutic benefit.<sup>26</sup>

224

225 **c. PBM and Dental Pulp Mesenchymal Stem Cells**

226 Emerging evidence indicates that PBM can modulate the proliferation and differentiation of  
227 dental pulp mesenchymal stem cells, promoting osteogenic and odontogenic pathways. This  
228 interaction suggests a potential role for PBM in regenerative endodontics and tissue engineering.  
229 Future investigations will likely explore dose-dependent effects and translational applications  
230 related to pulp regeneration and dentin repair.<sup>27</sup>

231

232 **d. PBM in Orthodontics, Temporomandibular Disorders, and Oral Surgery**

233 PBM has demonstrated consistent benefits in postoperative pain control and enhancement of  
234 healing following oral surgical procedures. However, evidence supporting its role in accelerating  
235 orthodontic tooth movement remains variable. Ongoing and future research aims to clarify its  
236 effectiveness in orthodontics and to establish standardized protocols for tooth movement  
237 acceleration and temporomandibular disorder management.<sup>28</sup>

238

239 **e. Inflammation Control and Soft Tissue Healing**

240 One of the most established applications of PBM is its ability to modulate inflammation and  
241 promote soft tissue healing. Through immunomodulatory and bioenergetic mechanisms, PBM  
242 supports wound repair and reduces postoperative morbidity. Future clinical integration may  
243 expand the use of PBM as a host-modulating adjunct in periodontal therapy and implant-related  
244 procedures.<sup>29</sup>

245

246 **f. PBM in Oral Oncology and Supportive Cancer Care**

247 PBM has gained strong clinical support for the prevention and management of  
248 oncotherapy-associated oral mucositis. Its inclusion in international clinical guidelines highlights  
249 its growing role in supportive cancer care. Future research is expected to refine dosing protocols,  
250 explore additional supportive oncologic applications, and facilitate broader clinical adoption.<sup>30</sup>

251

252 **g. PBM in Implant Dentistry and Peri-Implant Disease**

253 PBM is increasingly explored as an adjunctive therapy in implant dentistry to enhance soft tissue

256 healing and modulating peri-implant inflammation. Its non-thermal and non-destructive nature  
257 makes it particularly attractive for managing peri-implant conditions. Future studies may further  
258 clarify its role in supporting osseointegration and in antibiotic-sparing management of  
259 peri-implantitis.<sup>31</sup>

260  
261 Overall, photobiomodulation remains an actively evolving field in dentistry, with ongoing  
262 preclinical and clinical research continuously expanding its therapeutic indications and refining  
263 its clinical protocols.

## 264 5. Conclusion

265 In the Conclusion, Photobiomodulation is emerging as a promising alternative to conventional pain  
266 management in dentistry, with both lasers and light-emitting diodes showing potential for pain relief in  
267 dentistry and the other physical therapy but variability in study methods and treatment parameters limit  
268 its routine clinical use. Furthermore, we observe that more studies are needed before introducing the  
269 photobiomodulation in the clinical practise even in the routine dental extraction procedure.

## 272 273 Abbreviations:

274 **PBM** – Photobiomodulation

275 **PBMT** – Photobiomodulation Therapy

276 **LLLT** – Low-Level Laser Therapy

277 **LED** – Light-Emitting Diode

278 **ATP** – Adenosine Triphosphate

279 **ROS** – Reactive Oxygen Species

280 **NF-κB** – Nuclear Factor Kappa B

281 **AP-1** – Activator Protein-1

282 **VEGF** – Vascular Endothelial Growth Factor

283 **HIF-1α** – Hypoxia-Inducible Factor-1 Alpha

284 **TGF-β** – Transforming Growth Factor Beta

285 **NO** – Nitric Oxide

286 **Ca<sup>2+</sup>** – Calcium Ion

287 **cAMP** – Cyclic Adenosine Monophosphate  
288 **RANKL** – Receptor Activator of Nuclear Factor Kappa B Ligand  
289 **OPG** – Osteoprotegerin  
290 **TMJ** – Temporomandibular Joint  
291 **GBR** – Guided Bone Regeneration  
292 **PRP** – Platelet-Rich Plasma  
293 **PRF** – Platelet-Rich Fibrin  
294 **VAS** – Visual Analog Scale  
295 **GaAlAs** – Gallium–Aluminum–Arsenide Laser  
296 **ER,Cr:YSGG** – Erbium, Chromium-doped Yttrium Scandium Gallium Garnet Laser  
297 **MASCC/ISOO** – Multinational Association of Supportive Care in Cancer / International Society of Oral  
298 Oncology

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