



Optimizing Palatal Scar Healing Post Cleft Palate Repair- A Comprehensive Literature Review

Siddhant Thorat, Navaneetham R

Saveetha Dental College

(Received: 25 August 2025 Revised: 27 September 2025 Accepted: 14 October 2025)

KEYWORDS

Cleft palate repair, palatal scarring, wound healing, laser therapy, regenerative medicine, surgical techniques, scar management

ABSTRACT: Cleft palate is a congenital condition requiring surgical intervention to restore oral function, speech, and aesthetics. Despite advances in surgical techniques, post-surgical scarring remains a major challenge, often leading to speech impairment, restricted maxillofacial growth, and velopharyngeal dysfunction. Excessive fibrosis and contracture can compromise palatal mobility, affecting long-term outcomes. To mitigate these effects, various scar modulation strategies have been explored. Surgical refinements like Z-plasty, Furlow palatoplasty, and tension-free closures improve tissue flexibility, while pharmacological agents such as topical silicone, corticosteroids, and botulinum toxin help control fibroblast activity and prevent hypertrophic scarring. Additionally, laser therapy, platelet-rich plasma (PRP), stem cell applications, and mechanical therapies offer promising approaches to enhance healing and scar remodeling. Recent advances in molecular biology and regenerative medicine have deepened the understanding of scar pathophysiology, paving the way for biologically driven interventions. Early rehabilitation protocols and individualized treatment planning are critical in optimizing both aesthetic and functional recovery. This review aims to comprehensively evaluate current and emerging strategies for minimizing palatal scarring, highlighting future directions for improving post-surgical outcomes in cleft palate repair.

1. Introduction

Cleft palate is among the most prevalent congenital craniofacial anomalies with the global prevalence of cleft palate estimated ranging from approximately 1 in 500 to 1 in 1,000 live births, depending on geographic region and ethnic background¹. While modern surgical techniques and multidisciplinary care have significantly improved outcomes for affected individuals, challenges related to repair and subsequent healing persist, necessitating ongoing clinical attention. The standard approach to managing cleft palate involves early surgical intervention, typically within the first year of life, to separate the oral and nasal cavities, thereby enabling proper feeding and the development of normal speech^{1,2}. Over the years, several surgical modalities have evolved- from traditional one-stage repairs to more sophisticated techniques such as the Furlow double-opposing Z-plasty, which aim to achieve tension-free closure of the palatal tissues. Despite these improvements, surgical repair of cleft palate is not without its drawbacks³. A major

postoperative complication is the development of palatal scarring⁴. This literature review examines current strategies aimed at improving palatal scar healing following cleft palate repair. It evaluates various approaches- including surgical refinements, pharmacological interventions, laser therapies, regenerative medicine, and rehabilitative protocols- to assess their effectiveness and limitations. Enhancing scar healing is crucial for optimal maxillary development, improved orthodontic outcomes, and a better quality of life for individuals with cleft palate.

2. Objectives

Palatal wound healing

Wound healing is a biological process aimed at restoring the integrity and function of damaged tissue. It proceeds through overlapping phases: inflammation, tissue formation, and tissue remodeling. During inflammation, bleeding is controlled (hemostasis), and bacteria and debris are cleared. This is followed by the formation of



new tissue and wound contraction⁵. Finally, tissue remodeling strengthens the newly formed tissue, often resulting in scar formation. Intraoral wounds, such as those in the mouth, generally heal faster and with less scarring than skin wounds. This is partly due to lower levels of pro-inflammatory and pro-fibrotic cytokines, as well as the presence of saliva, which contains growth factors like epidermal growth factor (EGF)⁶. Fibroblast differences between skin and oral tissues also contribute to this enhanced healing.⁷

Within the oral cavity, buccal and palatal mucosa show different healing behaviors. Buccal mucosa is more flexible and elastic, while palatal mucosa is a mucoperiosteum- firm, cornified, and tightly bound to the bone⁸. These structural differences influence the wound healing outcomes⁹. Nonetheless, both follow a general process similar to that of skin. It is important to note that wound healing phases overlap in time and progress unevenly across the wound, with faster healing at the edges than in the center¹⁰.

Wound contraction and scarring are key factors contributing to growth disturbances after cleft palate repair. Wound contraction, responsible for 80-90% of closure, involves approximation of wound edges to reduce infection and dehydration risks¹¹. Two main theories attribute contraction to either fibroblast-generated tension or myofibroblast-mediated contraction via alpha-smooth muscle actin (ASMA)¹². The consensus theory integrates both, suggesting fibroblasts initiate tension that promotes myofibroblast differentiation, particularly under TGF- β 1 influence.¹³ Myofibroblasts then drive coordinated contraction of the matrix. Concurrently, matrix metalloproteinases (MMPs) mediate collagen remodeling, replacing type III with type I collagen aligned to tension vectors. This remodeling initiates scar formation. Scar tissue is characterized by reduced vascularity and cell density due to apoptosis of fibroblasts and endothelial cells. Lack of elastin resynthesis and aligned collagen fibers result in stiff, inelastic scar tissue. In the palate, scar adhesion to bone further impairs maxillary growth¹⁴.

Palatal Scarring in Cleft Palate Patients: Composition and Complications

Scarring of the palatal tissue following cleft palate repair is an unavoidable outcome of surgical intervention, as the body's natural healing process results in fibrotic tissue

formation. The primary goal of cleft palate closure is to restore both structure and function, allowing for improved speech, feeding, and facial growth. However, the healing process initiates a complex inflammatory response that involves fibroblast activation, extracellular matrix remodeling, and collagen deposition, leading to the development of scar tissue.¹⁵ The severity of fibrosis depends on multiple factors, including the surgical approach used, the patient's biological healing capacity, and the age at which the surgery is performed.¹⁶ A study by Pilmane et al highlights the scar's anti-inflammatory profile showing distinct cellular changes, including increased IL-1 α , IL-10, PAX-9, MSX-1, and RYK expression.¹⁷

The structural composition of palatal scar tissue differs markedly from that of healthy palatal mucosa. The normal mucosa comprises an extracellular matrix that balances Type I and Type III collagen, along with elastin fibers that provide resilience and adaptability.^{18,19} In contrast, scar tissue exhibits an excessive accumulation of Type I collagen while having significantly reduced amounts of Type III collagen and elastin. This compositional imbalance leads to the formation of a dense, rigid tissue structure that lacks the flexibility of normal mucosa, thereby restricting movement and elasticity¹⁸⁻²⁰. Additionally, myofibroblasts, which are specialized contractile cells responsible for wound closure, become hyperactive in palatal wounds, generating excessive contractile forces that further contribute to fibrosis and restriction of maxillary growth²¹.

While these cells are essential for normal wound healing, their persistent activity in the palatal region can lead to functional deficits. The long-term consequences of excessive palatal scarring extend beyond localized tissue stiffness. One of the most significant complications is velopharyngeal dysfunction (VPD), which arises when the scarred soft palate loses its ability to close effectively against the pharyngeal wall. This results in nasal air escape during speech, leading to hypernasality, articulation difficulties, and impaired resonance.^{22,23} Patients with severe VPD may require secondary corrective surgeries, such as pharyngeal flaps or sphincteroplasty, in addition to extensive speech therapy.²⁴



Challenges and Implications of Palatal Scarring in Orthodontic Treatment for Cleft Patients

1. Impact of Palatal Scarring on Orthodontic Treatment

Orthodontic management of cleft patients is often complicated by the presence of palatal fibrosis, which creates resistance to the forces applied for dental and skeletal correction. One of the most commonly used techniques, rapid maxillary expansion (RME), depends on the successful separation of the midpalatal suture to increase the transverse dimension of the maxilla.²⁵ However, severe scarring makes the palatal tissue rigid, hindering suture opening and reducing the effectiveness of conventional RME. In such cases, surgically assisted rapid palatal expansion (SARPE) or distraction osteogenesis may be required to achieve the necessary skeletal widening.²⁶

Additionally, scarring in cleft patients often leads to compromised periodontal health. The altered vascularization within fibrotic tissue reduces its ability to support tooth movement, leading to slower orthodontic progress and an increased risk of root resorption and bone loss.^{27, 28} Furthermore, cleft patients frequently present with missing, malformed, or ectopically erupted teeth, further complicating orthodontic mechanics and necessitating a multidisciplinary approach involving orthodontists, prosthodontists, and maxillofacial surgeons.²⁹ (Fig 1)



Figure 1: Palatal scarring post cleft palate surgery

2. Effects of Palatal Fibrosis on Maxillary Growth

Palatal fibrosis significantly restricts maxillary growth, particularly in the transverse and sagittal planes. The inflexibility of scar tissue prevents normal palatal expansion, leading to a constricted dental arch. As a result, many cleft patients develop maxillary hypoplasia, which is associated with Class III skeletal malocclusion, posterior crossbites and midfacial deficiency³⁰. These craniofacial discrepancies are especially concerning in growing children, as they influence both functional and aesthetic development. The underdevelopment of the maxilla can cause long-term dental malocclusions, often requiring early orthodontic and surgical interventions to minimize its effects.³¹

3. Psychosocial and Functional Implications of Palatal Scarring

Beyond the physical and orthodontic challenges, palatal scarring also has profound psychosocial effects. Malocclusion and midfacial hypoplasia can significantly impact facial esthetics, leading to self-esteem issues and social anxiety, particularly during adolescence.^{31,32} Additionally, speech difficulties caused by velopharyngeal dysfunction (VPD) can further contribute to communication challenges, negatively affecting academic performance and social interactions. Addressing these concerns through early intervention is essential to ensure both functional and psychological well-being.³³

Various treatment modalities for enhancement of palatal scar healing following cleft palate repair

1. Surgical Techniques & Modifications

In cleft palate repair, various surgical techniques have been developed to optimize functional outcomes and minimize scarring. Various palatal flaps are used to manage palatal scarring after cleft palate surgery, depending on the severity of fibrosis and functional impairment. Furlow's double-opposing Z-plasty is effective for mild scarring, improving velopharyngeal function by lengthening the soft palate by enhancing soft palate mobility by lengthening the palate and reorienting the palatal muscles, thereby reducing velopharyngeal insufficiency. Additionally, the V-Y pushback palatoplasty is employed to lengthen the palate and improve speech outcomes^{34,35}. Buccal myomucosal and tongue flaps provide well-vascularized tissue for



moderate to severe fibrosis or fistula closure, with tongue flaps requiring staged procedures³⁶ (Figure 2, 3). Radial forearm free flaps (RFFF) are used for extensive tissue deficiencies, offering pliable, vascularized tissue but requiring microvascular surgery.³⁷ Pharyngeal flaps are employed in cases of severe velopharyngeal dysfunction (VPD) to improve speech but may pose airway risks³⁸. The Bardach two-flap palatoplasty, introduced in 1967, enables complete closure of palatal clefts with two-layer closure of the hard palate and three-layer closure of the soft palate. This technique minimizes exposed bone, enhances soft palate mobility, and achieves normal speech in approximately 75–80% of patients, with an oronasal fistula rate averaging around 5.2%³⁹. The choice of flap depends on defect size, functional needs, and long-term rehabilitation goals. The use of fine sutures and tissue adhesives in these procedures minimizes tissue trauma and inflammation, promoting better healing with reduced scar formation⁴⁰. Additional surgical techniques to enhance palatal scar healing include the Modified Sommerlad-Furlow technique for wide clefts, vomer flap surgery for nasal lining reconstruction, and distraction osteogenesis to reduce soft tissue tension⁴¹. Two-stage palate repair minimizes healing tension, aiding maxillary growth and reducing scarring⁴². Collectively, these surgical strategies aim to improve both the functional and aesthetic results of cleft palate repair.

Pharmacological Approaches

To enhance scar healing and minimize hypertrophic scar formation following cleft palate surgery, several pharmacological interventions specifically target myofibroblast activity and the development of fibrous tissue. Topical Silicone gels and sheets are widely recognized as effective treatments for scar management. Silicone products work by creating a hydrated environment over the scar, which promotes healing and can help flatten and soften the scar tissue. They function by reducing collagen production, thereby preventing excessive fibrosis. The use of silicone is often recommended for both prevention and treatment of hypertrophic scars and keloids, making it a cornerstone in post-surgical scar care⁴³. Corticosteroids (e.g., Triamcinolone Acetonide- TAC) are also commonly used in the management of hypertrophic scars. Administered via intralesional injections, TAC helps alleviate inflammation and directly inhibits fibroblast proliferation, crucial factors in the development of

fibrous tissue. Additionally, corticosteroids reduce collagen synthesis and promote the activity of collagenase, which breaks down collagen⁴⁴. A notable study has shown that combining TAC with laser-assisted delivery can significantly improve scar texture over a six-month period, highlighting the potential for enhanced treatment outcomes through this combined approach.⁴⁵ 5-Fluorouracil (5-FU): This chemotherapeutic agent has been utilized for intralesional injections in treating various types of pathological scarring, including hypertrophic scars and keloids. 5-FU functions by inhibiting fibroblast proliferation and collagen synthesis, effectively reducing the amount of fibrous tissue within scars. Although the evidence supporting its use specifically for cleft palate scars is limited, its successful application in other types of scars suggests potential benefits in this area as well.⁴⁶

Botulinum Toxin (Botox): Botox has gained attention for its potential role in scar management. By reducing muscular tension around the scar, Botox can lead to improved flexibility and appearance of the scar tissue. This treatment works by inhibiting the release of acetylcholine, which diminishes muscle contractions that might exacerbate scar formation. Moreover, studies suggest that Botox may also reduce fibroblast activity, resulting in less collagen deposition, thereby contributing to smoother scars.⁴⁷ Various laser modalities, such as fractional laser resurfacing, are employed to treat scars by promoting collagen remodeling and improving overall skin texture. Laser treatments enhance the absorption of topical agents, such as TAC, making them more effective in reducing scar thickness and improving aesthetics. The precision of laser therapy allows for targeted treatment, minimizing damage to surrounding tissues while effectively addressing scar issues.⁴⁸ Administration of interferon proteins have shown promise in the management of hypertrophic scars and keloids.



Figure 2: Tongue flap for palatal cleft closure

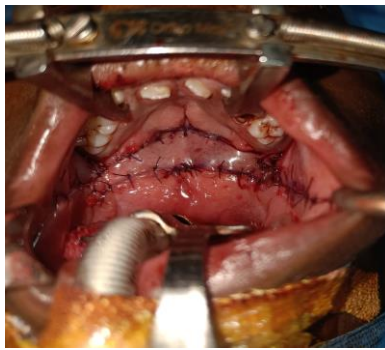


Figure 3: Bilateral buccinator myomucosal flap for palatal cleft closure

Interferons modulate fibroblast activity and can inhibit the proliferation of myofibroblasts, which are responsible for excessive collagen production in scars. Their use in scar management is still being explored, but they offer a potential avenue for reducing scar formation. Collectively, these pharmacological and non-pharmacological approaches aim to modulate the wound healing process, reduce excessive fibrosis, and enhance both the aesthetic and functional outcomes of scar formation following cleft palate surgery. Each method can be tailored to individual patient needs, and often they are used in combination to achieve the best possible results in scar management.⁴⁹

3. Laser Therapy & Light-Based Treatments

Laser therapy and light-based treatments have become integral in managing and improving scar appearance through various mechanisms. Ablative lasers, such as fractional CO₂ and Er:YAG lasers, work by removing the epidermis, leading to collagen remodeling and improved scar texture⁴⁸. However, these treatments require longer healing times and carry a higher risk profile compared to non-ablative therapies. Non-ablative lasers, like the 585

nm pulsed dye laser (PDL), target vascular components in early scars, reducing redness and preventing hypertrophic changes. Haoshu Chi et. Al studied the effects of PDL on cleft lip scars. Its efficacy on palatal scarring is yet to be studied⁵⁰. These treatments are associated with shorter recovery periods but may offer only minor improvements in the cosmetic appearance of atrophic and acne scars. Low level laser therapy (LLLT) has been explored for its potential to aid in wound healing and reduce fibrosis post-surgery in a systematic review by Seyyedi SA et. al⁵¹ While some studies suggest benefits, further research is needed to establish its efficacy in palatal scar management. Collectively, these laser-based interventions aim to modulate the wound healing process, enhance collagen remodeling, and improve the aesthetic and functional outcomes of scar formation. Laser therapy for cleft palate scars benefits from early initiation (as early as one month post-op) for better outcomes⁵². Combining modalities like PDL and fractional CO₂ lasers enhances scar texture and pliability. Intense Pulsed Light (IPL) used with lattice CO₂ lasers further improves scar evolution. Low-Level Laser Therapy (LLLT) shows promise in reducing fibrosis and accelerating wound healing. These interventions also contribute to improved functional outcomes and patient satisfaction.⁵³

4. Mechanical & Pressure Therapy

Mechanical and pressure therapies are pivotal in scar management, aiming to enhance scar appearance and functionality. Microneedling, or collagen induction therapy, employs fine needles to create controlled micro-injuries in the skin, stimulating collagen production and remodeling, which can improve the texture and appearance of scars^{54,55} Pressure dressings, often utilized in burn care, apply consistent pressure to scar tissue, reducing blood flow and oxygenation, thereby minimizing hypertrophic scar formation and enhancing scar pliability⁵⁶. Massage therapy complements this approach by mechanically disrupting fibrotic tissue, promoting flexibility and softness in the scarred area⁵⁷ In the context of cleft palate treatment, orthodontic and prosthetic interventions, such as palatal expanders and obturators, are employed to assist in palatal expansion and minimize post-surgical contractures, thereby improving both functional outcomes and aesthetic appearance⁵⁸



5. Regenerative & Biological Therapies

Emerging therapeutic approaches in scar management focus on enhancing tissue regeneration and reducing fibrosis through various modalities. Platelet-Rich Plasma (PRP) therapy involves concentrating platelets from the patient's own blood to deliver growth factors that promote faster, high-quality healing with reduced fibrosis⁵⁹. This technique has been explored for its potential in improving skin texture and elasticity. Stem cell therapy, particularly using mesenchymal stem cells, is under investigation for its ability to modulate scarring and enhance tissue regeneration.⁶⁰ These cells can differentiate into various cell types and secrete bioactive molecules that influence the wound healing process, potentially leading to improved outcomes in scar appearance and function⁶¹. Hyaluronic acid-based injections are utilized to improve tissue hydration and elasticity, thereby reducing scar rigidity. By attracting water molecules, hyaluronic acid maintains tissue turgor and flexibility, which can soften scar tissue and improve its appearance⁶². FTY720 nanofiber scaffolds enhance oral wound healing and prevent ONF formation by promoting pro-regenerative immune infiltration, modulating interleukin expression, upregulating Sox2, and stimulating keratinocyte migration and proliferation. Future research will explore the underlying mechanisms, supporting FTY720 patches as a potential strategy to prevent ONF in cleft palate repair.⁶³ Collectively, these therapies aim to synergistically enhance the wound healing process, minimize fibrosis, and improve the aesthetic and functional outcomes of scar management.

6. Post-Surgical Rehabilitation & Therapy

Integrating speech therapy and myofunctional exercises into rehabilitation protocols can significantly enhance oral function and alleviate tension on scarred tissues. These interventions focus on strengthening the orofacial musculature, optimizing speech articulation, and improving swallowing efficiency, thereby reducing undue stress on healing tissues⁶⁴. Concurrently, scar massage and physiotherapy are vital in managing scar tissue by breaking down adhesions, enhancing

Modality	Subtypes / Techniques	Mechanism / Purpose	Key Benefits / Outcomes
1. Surgical Techniques	<ul style="list-style-type: none"> - Furlow Z-Plasty - V-Y Pushback - Bardach Two-Flap -Buccal & Tongue Flaps - RFFF - Pharyngeal Flaps 	Reorientation of palatal muscles, lengthening palate, improving tissue vascularity and closure tension	Reduces VPD, improves mobility and speech, decreases scarring, improves healing
2. Pharmacological Approaches	<ul style="list-style-type: none"> - Topical Silicone - Corticosteroids (TAC) - 5-Fluorouracil (5-FU) - Botulinum Toxin - Interferons 	Modulate fibroblast activity, reduce collagen synthesis, control inflammation	Decrease hypertrophic scarring, improve scar texture, enhance drug delivery when combined
3. Laser Therapy	<ul style="list-style-type: none"> - Fractional CO₂ Laser - Er:YAG - Pulsed Dye Laser (PDL) - Low-Level Laser Therapy (LLLT) 	Stimulates collagen remodeling, improves vascular response, enhances absorption of topical agents	Enhances scar aesthetics, reduces thickness and redness, shortens recovery
4. Mechanical & Pressure Therapy	<ul style="list-style-type: none"> - Microneedling - Pressure Dressings - Scar Massage - Palatal Expanders / Obturators 	Induce collagen remodeling, break down fibrotic tissue, maintain tissue flexibility	Improves scar pliability and texture, prevents contracture, supports palatal growth
5. Regenerative Therapies	<ul style="list-style-type: none"> - PRP - Stem Cells - Hyaluronic Acid - FTY720 Scaffolds 	Stimulate tissue regeneration, modulate immune response, maintain hydration	Accelerate healing, reduce fibrosis, improve elasticity and cellular organization
6. Rehabilitative Therapies	<ul style="list-style-type: none"> - Speech Therapy - Myofunctional Exercises - Transverse Friction Massage (TFM) 	Strengthen orofacial musculature, improve articulation, reduce tension on healing tissues	Improves oral function, reduces contracture, enhances patient recovery and quality of life

Figure 4: Palatal scarring post cleft palate surgery

flexibility, and minimizing the risk of contracture formation. For instance, transverse friction massage (TFM) has been shown to influence neuromotor mechanisms by altering electromechanical delay and force transmission, suggesting a reduction in active muscle stiffness⁶⁵. Collectively, these therapeutic strategies contribute to improved functional outcomes and aesthetic results in patients undergoing rehabilitation for conditions involving scar tissue.

Conclusion

Palatal scarring following cleft palate repair presents a significant challenge due to its potential to impair speech, hinder maxillary growth, and negatively affect both function and aesthetics. To address this complex issue, a multimodal approach that combines surgical precision, pharmacologic interventions, laser therapy, and rehabilitative strategies has emerged as the most effective pathway for improving scar outcomes. Early identification of scar type and timely intervention are critical, as they allow for treatment plans to be customized according to the individual's functional requirements and tissue response. Surgical techniques aimed at minimizing trauma, maintaining vascular integrity, and ensuring tension-free closure play a foundational role in preventing excessive scarring. Adjunctive pharmacologic agents—such as corticosteroids, botulinum toxin, and novel anti-fibrotic compounds—can modulate the wound healing process, reducing inflammation and fibroblast proliferation. Laser



therapies, including fractional CO₂ and pulsed dye lasers, contribute by promoting collagen remodeling, enhancing tissue pliability, and improving the overall appearance of the scar. Rehabilitation strategies, such as speech therapy, palatal stretching exercises, and myofunctional training, further aid in restoring oral function and enhancing scar mobility. In addition, advances in regenerative medicine—such as the use of growth factors, platelet-rich plasma (PRP), and mesenchymal stem cells—are paving the way for biological modulation of scar formation at the molecular level. This review consolidates current evidence on these diverse scar reduction strategies, evaluates their individual and combined effectiveness, and highlights future directions for optimizing outcomes in cleft palate patients. An integrative and patient-specific therapeutic model is key to addressing the multifactorial nature of palatal scarring and achieving optimal long-term functional and aesthetic results.

Ethics Approval

This study was approved by the [Committee], approval/exemption number [XXXX].

Competing Interests

The authors declare that they have no competing interests.

Data Availability

The data supporting the findings of this study are available from the corresponding author upon reasonable request.

Funding

This research received no external funding.

References

1. Global prevalence of cleft palate, cleft lip and cleft palate and lip: A comprehensive systematic review and meta-analysis. *Journal of Stomatology, Oral and Maxillofacial Surgery*. 2022 Apr 1;123(2):110–20.
2. Roberts MCT, Semb G, Shaw WC. Strategies for the Advancement of Surgical Methods in Cleft Lip and Palate. *The Cleft Palate-Craniofacial Journal* [Internet]. 1991 [cited 2025 Mar 12]; Available from: https://journals.sagepub.com/doi/10.1597/1545-1569_1991_028_0141_sftaos_2.3.co_2
3. Naidu P, Yao CA, Chong DK, Magee WP 3rd. Cleft Palate Repair: A History of Techniques and Variations. *Plast Reconstr Surg Glob Open*. 2022 Mar;10(3):e4019.
4. desJardins-Park HE, Mascharak S, Chinta MS, Wan DC, Longaker MT. The Spectrum of Scarring in Craniofacial Wound Repair. *Front Physiol*. 2019 Mar 29;10:322.
5. Website [Internet]. Available from: Schultz GS, Chin GA, Moldawer L, et al. Principles of Wound Healing. In: Fitridge R, Thompson M, editors. *Mechanisms of Vascular Disease: A Reference Book for Vascular Specialists* [Internet]. Adelaide (AU): University of Adelaide Press; 2011. 23. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK534261/>
6. Waasdorp M, Krom BP, Bikker FJ, van Zuijlen PPM, Niessen FB, Gibbs S. The Bigger Picture: Why Oral Mucosa Heals Better Than Skin. *Biomolecules* [Internet]. 2021 Aug 6;11(8). Available from: <http://dx.doi.org/10.3390/biom11081165>
7. Wight TN. Provisional matrix: A role for versican and hyaluronan. *Matrix Biol*. 2017 Jul;60-61:38–56.
8. Manogaran S, Ramadoss R, Hemashree, Selvam SP, Sundar S, Krishnasamy N. Stimuli responsive materials in the management of cleft lip and cleft palate – A systematic review. *Indian J Otolaryngol Head Neck Surg* [Internet]. 2025 Jul 30; Available from: <https://link.springer.com/10.1007/s12070-025-05907-5>
9. Squier CA, Kremer MJ. Biology of Oral Mucosa and Esophagus. *J Natl Cancer Inst Monogr*. 2001 Oct 1;2001(29):7–15.
10. Clark RAF, Henson PM. *The Molecular and Cellular Biology of Wound Repair*. Springer Science & Business Media; 2012. 601 p.
11. McGrath MH, Simon RH. Wound geometry and the kinetics of wound contraction. *Plast Reconstr Surg*. 1983 Jul;72(1):66–73.
12. Hinz B, Celetta G, Tomasek JJ, Gabbiani G, Chaponnier C. Alpha-smooth muscle actin



- expression upregulates fibroblast contractile activity. *Mol Biol Cell*. 2001 Sep;12(9):2730–41.
13. Muddasani V, Kamalakannan SK, S H, Arun A, J K. The Use of Ultrasound-Guided 3D-Constructed Obturator Device in the Management of Cleft Lip and Palate: A Case Series. *Cureus*. 2024 Jul;16(7):e64948.
 14. Tomasek JJ, Gabbiani G, Hinz B, Chaponnier C, Brown RA. Myofibroblasts and mechano-regulation of connective tissue remodelling. *Nat Rev Mol Cell Biol*. 2002 May;3(5):349–63.
 15. Ishikawa H, Nakamura S, Misaki K, Kudoh M, Fukuda H, Yoshida S. Scar Tissue Distribution on Palates and its Relation to Maxillary Dental Arch Form. *The Cleft Palate-Craniofacial Journal* [Internet]. 1998 [cited 2025 Mar 28]; Available from: https://journals.sagepub.com/doi/10.1597/1545-1569_1998_035_0313_stdopa_2.3.co_2
 16. Muscle fibrosis in the soft palate: Delivery of cells, growth factors and anti-fibrotics. *Advanced Drug Delivery Reviews*. 2019 Jun 1;146:60–76.
 17. Pilmane M, Jain N, Nadzina E, Fedirko P, Sumeraga G. Immunohistochemical evaluation of the cleft-affected scar tissue three decades post-corrective surgery: A rare case report. *Acta Otolaryngologica Case Reports* [Internet]. 2022 Dec 31 [cited 2025 Apr 29]; Available from: <https://www.tandfonline.com/doi/abs/10.1080/23772484.2022.2146586>
 18. Website [Internet]. Available from: Zito PM, Jawad BA, Hohman MH, et al. Z-Plasty. [Updated 2023 Jul 5]. In: *StatPearls* [Internet]. Treasure Island (FL): StatPearls Publishing; 2025 Jan-. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK507775/>
 19. Frantz C, Stewart KM, Weaver VM. The extracellular matrix at a glance. *J Cell Sci*. 2010 Dec 15;123(Pt 24):4195–200.
 20. Egbunah UP, Zhu YT, Hauri DD, Ratz T, Thor ALI. The effect of collagen matrix graft on palatal fistula formation after cleft palate repair: A preliminary randomized controlled study. *J Cleft Lip Palate Craniofacial Anomalies*. 2024 Jan;11(1):21–9.
 21. Li B, Wang JHC. Fibroblasts and myofibroblasts in wound healing: force generation and measurement. *J Tissue Viability*. 2011 Nov;20(4):108–20.
 22. Asar A, Gaber R, Yehia M, El-Kassaby MAW. Treatment algorithm for velopharyngeal dysfunction in patients with cleft palate: a systematic review. *Br J Oral Maxillofac Surg*. 2023 May;61(4):259–66.
 23. Anishya D, Nagesh S. Assessment of Nasal Aesthetic Parameters in Patients with Unilateral Cleft Lip and Palate - A Retrospective Study. *Cleft Palate Craniofac J*. 2025 Sep;62(9):1496–503.
 24. Marsh JL. Management of velopharyngeal dysfunction: differential diagnosis for differential management. *J Craniofac Surg*. 2003 Sep;14(5):621–8; discussion 629.
 25. Khairi N, Halilah T, Khandakji M, Bartzela T. Rapid Maxillary Expansion Treatment in Patients with Cleft Lip and Palate: A Survey on Clinical Experience in the European Cleft Centers. *J Clin Med* [Internet]. 2023 Apr 27;12(9). Available from: <http://dx.doi.org/10.3390/jcm12093159>
 26. Kumar Shetty S, Neeraj, Kumar M, K. Madhur V. Surgically assisted rapid palatal expansion (SARPE): A literature review. *Sch J Dent Sci*. 2021 Jan 9;8(1):25–40.
 27. Wu Q, Li Z, Zhang Y, Peng X, Zhou X. Dental caries and periodontitis risk factors in cleft lip and palate patients. *Front Pediatr*. 2022;10:1092809.
 28. Yang C, Qian Y, Chen Z, Yang Y, Yu Q. Study on Tooth Movement After the Alveolar Bone Grafting in Patients With Unilateral Cleft Lip and Palate. *J Craniofac Surg*. 2019 Jun;30(4):e284–8.
 29. De Stefani A, Bruno G, Barone M, Gracco A. Orthodontic, Maxillofacial Surgery, and Prosthodontic Rehabilitation Supported by Miniscrew in a Patient with Cleft Lip and Palate. *Case Rep Dent*. 2021 Sep 25;2021:5540487.
 30. Yen SLK. Protocols for Late Maxillary Protraction in Cleft Lip and Palate Patients at



- Childrens Hospital Los Angeles. *Semin Orthod.* 2011 Jun 1;17(2):138–48.
31. Zhou C, Duan P, He H, Song J, Hu M, Liu Y, et al. Expert consensus on pediatric orthodontic therapies of malocclusions in children. *Int J Oral Sci.* 2024 Apr 16;16(1):32.
32. Effects of cleft lip and palate on children's psychological health: A systematic review. *Journal of Taibah University Medical Sciences.* 2018 Aug 1;13(4):311–8.
33. Barr L, Thibeault SL, Muntz H, de Serres L. Quality of life in children with velopharyngeal insufficiency. *Arch Otolaryngol Head Neck Surg.* 2007 Mar;133(3):224–9.
34. Website [Internet]. Available from: Almasri MA, editor. *Designing Strategies for Cleft Lip and Palate Care* [Internet]. InTech; 2017. Available from: <http://dx.doi.org/10.5772/62857>
35. Comparison of the effect of the rotation palatoplasty and V–Y pushback palatoplasty techniques on palate elongation with magnetic resonance imaging. *International Journal of Oral and Maxillofacial Surgery.* 2015 Jun 1;44(6):738–44.
36. Choi JM, Park H, Oh TS. Use of a buccinator myomucosal flap and bilateral pedicled buccal fat pad transfer in wide palatal fistula repair: a case report. *Arch Craniofac Surg.* 2021 Aug;22(4):209–13.
37. Microvascular radial forearm fasciocutaneous free flap in hard palate reconstruction. *European Journal of Surgical Oncology (EJSO).* 2005 Sep 1;31(7):784–91.
38. Lee YW, Bae YC, Park SM, Nam SB, Seo HJ, Kim GW. Outcomes of a superiorly-based pharyngeal flap for the correction of velopharyngeal dysfunction. *Arch Craniofac Surg.* 2020 Feb;21(1):22–6.
39. Two-Flap palatoplasty: Bardach's technique. *Operative Techniques in Plastic and Reconstructive Surgery.* 1995 Nov 1;2(4):211–4.
40. Dental materials for cleft palate repair. *Materials Science and Engineering: C.* 2016 Apr 1;61:1018–28.
41. Agrawal K. Cleft palate repair and variations. *Indian J Plast Surg.* 2009 Oct;42 Suppl(Suppl):S102–9.
42. Liao YF, Yang IY, Wang R, Yun C, Huang CS. Two-stage palate repair with delayed hard palate closure is related to favorable maxillary growth in unilateral cleft lip and palate. *Plast Reconstr Surg.* 2010 May;125(5):1503–10.
43. Alam M, Waibel J, Uebelhoer N, Arndt KA, Dover J, Donelan M, et al. *Treatment of Scars from Burns and Trauma.* McGraw Hill Professional; 2020. 200 p.
44. Jalali M, Bayat A. Current use of steroids in management of abnormal raised skin scars. *Surgeon.* 2007 Jun;5(3):175–80.
45. Leszczynski R, da Silva CAP, Kuczynski U, da Silva EMK. Laser therapy for treating hypertrophic and keloid scars. *Cochrane Libr* [Internet]. 2015 Apr 14; Available from: <https://doi.wiley.com/10.1002/14651858.CD011642>
46. Gauglitz GG, Korting HC, Pavicic T, Ruzicka T, Jeschke MG. Hypertrophic scarring and keloids: pathomechanisms and current and emerging treatment strategies. *Mol Med.* 2011 Jan-Feb;17(1-2):113–25.
47. Botulinum toxin to reduce cleft lip/palate scars after surgery and improves scar quality in children: A systematic review and meta-analysis. *Global Pediatrics.* 2023 Jun 1;4:100054.
48. Hasegawa T, Matsukura T, Mizuno Y, Suga Y, Ogawa H, Ikeda S. Clinical trial of a laser device called fractional photothermolysis system for acne scars. *J Dermatol.* 2006 Sep;33(9):623–7.
49. Effects of interferons on proliferation and collagen synthesis of rat palatal wound fibroblasts. *Archives of Oral Biology.* 1999 Jul 1;44(7):541–7.
50. Chi H, Peng H, Zhao X, Zhou G, Shen L, Cai M. The Effectiveness of 595-nm Pulsed Dye Laser for the Treatment of Bilateral Cleft-Lip Scars in Asian Patients: A 6-Month Prospective, Randomized,



- Self-Controlled Trial. *Adv Wound Care (New Rochelle)*. 2024 Jun;13(6):322–8.
51. Seyyedi SA, Taram S, Heydari M, Valizadeh R. Efficacy of low-level laser therapy in oral mucosal surgical wound healing: a systematic review and meta-analysis. *Open Exploration*. 2019 Feb 11;3(5):451–60.
52. Sun Y, Li Z, Qi X, Wang B, Yu N, Huang J, et al. Laser therapy for treating cleft lip or/and palate scarring-a systematic review and meta-analysis. *Lasers Med Sci*. 2024 Jun 20;39(1):160.
53. Murugesan D, Felicita AS, Jayaseelan VP. Genetic analysis of TPM1 gene polymorphism (rs11071720) in patients with non-syndromic cleft lip/palate in the south Indian population. A case control study. *Odovtos - Int J Dent Sci*. 2024 May 9;222–32.
54. Alster TS, Li MKY. Microneedling of Scars: A Large Prospective Study with Long-Term Follow-Up. *Plast Reconstr Surg*. 2020 Feb;145(2):358–64.
55. Bandral MR, Padgavankar PH, Japatti SR, Gir PJ, Siddegowda CY, Gir RJ. Clinical Evaluation of Microneedling Therapy in the Management of Facial Scar: A Prospective Randomized Study. *J Maxillofac Oral Surg*. 2019 Dec;18(4):572–8.
56. Bardach J, Morris HL. *Multidisciplinary Management of Cleft Lip and Palate*. W.B. Saunders Company; 1990. 904 p.
57. Plummer P. *Technic of Cleft-Lip and Cleft-Palate Massage*. *The Physical Therapy Review*. 1937 Mar 1;17(2):54–6.
58. Goyal M, Chopra R, Bansal K, Marwaha M. Role of obturators and other feeding interventions in patients with cleft lip and palate: a review. *Eur Arch Paediatr Dent*. 2014 Feb;15(1):1–9.
59. El-Anwar MW, Nofal AAF, Khalifa M, Quriba AS. Use of autologous platelet-rich plasma in complete cleft palate repair. *Laryngoscope*. 2016 Jul;126(7):1524–8.
60. Kuzhalvaimozhi P, Ravindran V, Subhashini. Gingival health status in children with and without cleft palate: A case control study. *J Complement Med Res*. 2020;11(4):140.
61. Schreurs M, Maarten Suttorp C, Mutsaers HAM, Kuijpers-Jagtman AM, den Hoff JWV, Ongkosuwito EM, et al. Tissue engineering strategies combining molecular targets against inflammation and fibrosis, and umbilical cord blood stem cells to improve hampered muscle and skin regeneration following cleft repair. *Medicinal Research Reviews*. 2020 Jan 1;40(1):9–26.
62. Joshi VM, Kandaswamy E, Germain JS, Schiavo JH, Fm HS. Effect of hyaluronic acid on palatal wound healing: A systematic review. *Clin Oral Investig*. 2024 Oct 3;28(10):565.
63. Ballestas SA, Turner TC, Kamalakar A, Stephenson YC, Willett NJ, Goudy SL, et al. Improving hard palate wound healing using immune modulatory autotherapies. *Acta Biomater*. 2019 Jun;91:209–19.
64. Levrini L, Baldelli G, Castellani C, Ricci L, Dellavia CPB, Giannotta N, et al. Myofunctional Speech Therapy for Facial Rejuvenation and Orofacial Function Improvement: A Systematic Review. *J Funct Morphol Kinesiol [Internet]*. 2024 Jun 5;9(2). Available from: <http://dx.doi.org/10.3390/jfmk9020099>
65. The neuromotor effects of transverse friction massage. *Manual Therapy*. 2016 Dec 1;26:70–6.