

Management Considerations for Pediatric Oral Surgery and Oral Pathology

Latest Revision

2020

How to Cite: American Academy of Pediatric Dentistry. Management considerations for pediatric oral surgery and oral pathology. The Reference Manual of Pediatric Dentistry. Chicago, Ill.: American Academy of Pediatric Dentistry; 2023:527-36.

Abstract

This best practice defines, describes clinical presentation, and establishes criteria and therapeutic goals for common pediatric oral surgery procedures and oral pathological conditions. Pediatric oral surgery requires special considerations such as parental consent, knowledge of developing anatomy and dentition, potential for adverse effects on growth, behavior guidance, and peri- and postoperative management. Odontogenic infections usually are managed with pulp therapy, extraction, or incision and drainage. However, cases with systemic manifestations require antibiotic therapy. Extraction of erupted, unerupted, impacted, and supernumerary teeth are discussed with emphasis on a careful approach to avoid injury to adjacent teeth, permanent successors, and other hard and soft tissues. Considerations for surgical correction of frenulum attachments are reviewed. Guidance is provided for biopsies, a procedure which can establish a definitive diagnosis for most oral lesions. Common lesions in infants include Epstein pearls, Bohn nodules, and dental lamina cysts, and rare lesions include congenital epulis of the newborn and melanotic neuroectodermal tumor of infancy. Management of these lesions and natal and neonatal teeth is reviewed. Oral lesions in children and adolescents including eruption cysts, mucoceles, recurrent aphthous stomatitis, and pyogenic granuloma also are addressed. While most lesions are mucosal conditions, developmental anomalies, or inflammatory lesions, practitioners should be vigilant for neoplastic diseases.

This document was developed through a collaborative effort of the American Academy of Pediatric Dentistry Councils on Clinical Affairs and Scientific Affairs to offer updated information and guidance on management considerations for pediatric oral surgery and oral pathology.

KEYWORDS: DIAGNOSIS; ORAL; ORAL SURGICAL PROCEDURES; PATHOLOGY; TOOTH EXTRACTION

Purpose

The American Academy of Pediatric Dentistry (AAPD) intends this document to define, describe clinical presentation, and set forth general criteria and therapeutic goals for common pediatric oral surgery procedures and oral pathological conditions.

Methods

Recommendations on management considerations for pediatric oral surgery and oral pathology were developed by the Council on Clinical Affairs and adopted in 2005.¹ This document is a revision of the previous version, last revised in 2015.² It is based on a review of the current dental and medical literature related to pediatric oral surgery, including a search of the PubMed®/MEDLINE database using the terms: pediatric AND oral surgery, oral pathology, extraction, odontogenic infections, impacted canines, third molars, supernumerary teeth, mesiodens, mucocele, eruption cyst, eruption hematoma, gingival keratin cysts, Epstein pearls, Bohn's nodules, congenital epulis of newborn, dental lamina cysts, natal teeth, neonatal teeth, squamous papilloma, verruca vulgaris, irritation fibroma, recurrent aphthous stomatitis, localized juvenile spongiotic gingival hyperplasia, and pyogenic granuloma; fields: all; limits: within the last 10 years, humans, English, clinical trials. Papers for review were chosen from

the list of articles matching these criteria and from references with selected articles. When data did not appear sufficient or were inconclusive, recommendations were based upon expert and/or consensus opinion by experience researchers and clinicians. In addition, the manual *Parameters of Care: Clinical Practice Guidelines for Oral and Maxillofacial Surgery*,³ developed by the American Association of Oral and Maxillofacial Surgeons (AAOMS), was consulted.

General considerations

Surgery performed on pediatric patients involves special considerations unique to this population. Several critical issues deserve to be addressed.

Preoperative considerations

Informed consent

Before any surgical procedure, informed consent must be obtained from the parent or legal guardian. For more information, refer to AAPD's *Informed Consent*.⁴

ABBREVIATIONS

AAOMS: American Association of Oral and Maxillofacial Surgeons. **AAPD:** American Academy of Pediatric Dentistry. **HPV:** Human papilloma virus. **VKDB:** Vitamin K deficiency bleeding.

Medical evaluation

Important considerations in treating a pediatric patient include obtaining a thorough medical history, obtaining appropriate medical and dental consultations, anticipating and preventing emergency situations, and being prepared to treat emergency situations.⁵

Dental evaluation

It is important to perform a thorough clinical and radiographic preoperative evaluation of the dentition as well as a clinical examination of extraoral and intraoral soft tissues.⁵⁻⁷ Radiographs can include intraoral films and extraoral imaging if the area of interest extends beyond the dentoalveolar complex. Surgery involving the maxilla and mandible of young patients is complicated by the presence of developing tooth follicles. Knowledge of the anatomy of a child's developing maxilla and mandible and the avoidance of injury to the dental follicles can prevent complications.⁸ To minimize the negative effects of surgery on the developing dentition, careful planning using radiographs, tomography,⁹ cone beam computed tomography,¹⁰ and/or three-dimensional imaging techniques¹¹ is necessary to provide valuable information to assess the presence, absence, location, and/or quality of individual crown and root development.^{8,12,13}

Growth and development

The potential for adverse effects on growth from injuries and/or surgery in the oral and maxillofacial region markedly increases the potential for risks and complications in the pediatric population. Traumatic injuries involving the maxillofacial region can adversely affect growth, development, and function. Therefore, a thorough evaluation of the growing patient must be done before surgical interventions are performed to minimize the risk of damage to the growing facial complex.¹⁴

Behavioral evaluation

Behavioral guidance of children in the operative and perioperative periods presents a special challenge. Many children benefit from modalities beyond local anesthesia and nitrous oxide/oxygen inhalation to minimize their anxiety.^{4,14} Management of children under sedation or general anesthesia requires extensive training and expertise.^{15,16} Special attention should be given to the assessment of the social, emotional, and psychological status and cognitive level of the pediatric patient prior to surgery.¹⁴ Children have many unvoiced fears concerning the surgical experience, and their psychological management requires that the dentist be cognizant of their emotional status. Answering questions concerning the surgery is important and should be done in the presence of the parent.

Peri- and postoperative considerations

Metabolic management of children following surgery frequently is more complex than that of adults. Special consideration should be given to caloric intake, fluid and electrolyte

management, and blood replacement. Comprehensive management of the pediatric patient following extensive oral and maxillofacial surgery usually is best accomplished in a facility that has expertise and experience in the management of young patients (i.e., a children's hospital).¹⁴

Recommendations

Odontogenic infections

In children, odontogenic infections may involve more than one tooth and usually are due to caries lesions, periodontal problems, pathology (e.g., dens invaginatus), or a history of trauma.^{17,18} Untreated odontogenic infections can lead to pain, difficulty eating or drinking, abscess, cellulitis, septicemia, airway compromise, and life-threatening infections.¹⁹ Facial cellulitis results from unresolved abscess that has spread to cutaneous or subcutaneous soft tissue planes in the head and neck region.¹⁹ In these children, dehydration is a significant consideration; prompt treatment of the source of infection is imperative.

With infections of the upper portion of the face, patients usually complain of facial pain, fever, and malaise.²⁰ Care must be taken to rule out sinusitis or non-odontogenic infections, as symptoms may mimic an odontogenic infection. Occasionally in upper face infections, it may be difficult to find the true cause.¹⁴ Infections of the lower face usually involve pain, swelling, and trismus.^{3,17} They frequently are associated with teeth, skin, local lymph nodes, and salivary glands.¹⁷ Most odontogenic infections occur in the upper face; however, infections in the mandibular region are more frequent in older children.²⁰

Most odontogenic infections can be managed with pulp therapy, extraction, or incision and drainage.⁵ Infections of odontogenic origin with systemic manifestations (e.g., elevated temperature [102 to 104 degrees Fahrenheit], facial cellulitis, difficulty in breathing or swallowing, fatigue, nausea) require antibiotic therapy.¹⁹ Severe but rare complications of odontogenic infections include cavernous sinus thrombosis and Ludwig's angina.^{17,19} These conditions can be life threatening and may require immediate hospitalization with intravenous antibiotics, incision and drainage, and referral/consultation with an oral and maxillofacial surgeon.^{17,19}

Extraction of erupted teeth

Maxillary and mandibular anterior teeth

Most primary and permanent maxillary and mandibular central incisors, lateral incisors, and canines have conical single roots. In most cases, extraction of anterior teeth is accomplished with a rotational movement due to their single root anatomy.⁵ However, there have been reported cases of accessory roots observed in primary canines.^{21,22} Radiographic examination is helpful to identify differences in root anatomy prior to extraction.²¹ Care should be taken to avoid placing any force on adjacent teeth that could become luxated or dislodged easily due to their root anatomy.

Maxillary and mandibular molars

Primary molars have roots that are smaller in diameter and more divergent than permanent molars. Root fracture in primary molars is not uncommon due to these characteristics as well as the potential weakening of the roots caused by the eruption of their permanent successors.⁵ Prior to extraction, the relationship of the primary roots to the developing succedaneous tooth should be assessed. To avoid inadvertent extraction or dislocation of or trauma to the permanent successor, pressure should be avoided in the furcation area or the tooth may need to be sectioned to protect the developing permanent tooth.

Molar extractions are accomplished by using slow continuous palatal/lingual and buccal force allowing for the expansion of the alveolar bone to accommodate the divergent roots and reduce the risk of root fracture.⁵ When extracting mandibular molars, care should be taken to support the mandible to protect the temporomandibular joints from injury.⁵

Fractured primary tooth roots

The presence of a root tip should not be regarded as a positive indication for its removal. The dilemma to consider when managing a retained primary tooth root is that removing the root tip may cause damage to the succedaneous tooth, while leaving the root tip may increase the chance for postoperative infection and delay eruption of the permanent successor.⁵ Radiographs can assist in the decision process. Expert opinion suggests that if the fractured root tip can be removed easily, it should be removed.⁵ If the root tip is very small, located deep in the socket, situated in close proximity to the permanent successor, or unable to be retrieved after several attempts, it is best left to be resorbed.⁵ The parent must be informed and a complete record of the discussion must be documented. The patient should be monitored at appropriate intervals to evaluate for potential adverse effects.

Management of unerupted and impacted teeth

There is a wide clinical spectrum of disorders of eruption in both primary and permanent teeth in children. These may be syndromic or nonsyndromic and include ankyloses,²⁷⁻²⁸ secondary retention,²⁸ tooth impaction, or primary failure of eruption.²⁹ Clinically, it may be difficult to differentiate between the various disruptions; however, there have been many reports^{30,31} to assist the clinician in making a diagnosis. Increasing evidence supports a genetic etiology for some eruption disruptions which may help in a definitive diagnosis.²⁹ Management of unerupted teeth will depend on whether the affected tooth is likely to respond to orthodontic forces. If not, surgical extraction is the preferred treatment option.²⁹

Impacted canines

Tooth impaction may occur due to a mechanical obstruction. Permanent maxillary canines are second to third molars in

frequency of impaction.³² Early detection of an ectopically erupting canine through visual inspection, palpation, and radiographic examination is important to maximize success of an intervention.³³ Routine evaluation of patients in mid-mixed dentition should involve identifying signs such as lack of canine bulges and asymmetry in pattern of exfoliation. Abnormal angulation or ectopic eruption of developing permanent cuspids can be assessed radiographically.³³ When the cusp tip of the permanent canine is just mesial to or overlaying the distal half of the long axis of the root of the permanent lateral incisor, canine palatal impaction usually occurs.³² Extraction of the primary canines is the treatment of choice to correct palatally displaced canines or to prevent resorption of adjacent teeth.³² One study showed that 78 percent of ectopically erupting permanent canines normalized within 12 months after removal of the primary canines; 64 percent normalized when the starting canine position overlapped the lateral incisor by more than half of the root; and 91 percent normalized when the starting canine position overlapped the lateral incisor by less than half of the root.³² If no improvement in canine position occurs in a year, surgical and/or orthodontic treatment were suggested.^{32,33} A Cochrane review³⁴ and a systematic review³⁵ reported no evidence to support extraction of primary canines to facilitate eruption of ectopic permanent maxillary canines. A prospective randomized clinical trial demonstrated that extraction of primary canines is an effective measure to correct palatally displaced maxillary canines and is more successful in children with an early diagnosis.³⁶ Consultation between the practitioner and an orthodontist may be useful in the final treatment decision.

Third molars

Panoramic or periapical radiographic examination is indicated in late adolescence to assess the presence, position, and development of third molars.⁷ The AAOMS recommends that a decision to remove or retain third molars should be made before the middle of the third decade.³ Evidence-based research supports the removal of third molars when pathology (e.g., cysts or tumors, caries, infection, pericoronitis, periodontal disease, detrimental changes of adjacent teeth or bone) is associated and/or the tooth is malpositioned or nonfunctional (i.e., an unopposed tooth).³⁷⁻³⁹ There is no evidence to support³⁷⁻⁴⁰ or refute³ the prophylactic removal of disease-free impacted third molars. Factors that increase the risk for surgical complications (e.g., coexisting systemic conditions, location of peripheral nerves, history of temporomandibular joint disease, presence of cysts or tumors)^{38,39} and position and inclination of the molar in question⁴¹ should be assessed. The age of the patient is only a secondary consideration.⁴¹ Referral to an oral and maxillofacial surgeon for consultation and subsequent treatment may be indicated. When a decision is made to retain impacted third molars, they should be monitored for change in position and/or development of pathology, which may necessitate later removal.

Supernumerary teeth

Supernumerary teeth and hyperdontia are terms to describe an excess in tooth number. Supernumerary teeth are thought to be related to disturbances in the initiation and proliferation stages of dental development.²¹ Although some supernumerary teeth may be syndrome-associated (e.g., cleidocranial dysplasia) or of familial inheritance pattern, most supernumerary teeth occur as isolated events.²¹

Supernumerary teeth can occur in either the primary or permanent dentition.^{21,42,43} In 33 percent of the cases, a supernumerary tooth in the primary dentition is followed by the supernumerary tooth complement in the permanent dentition.⁴⁴ Reports in incidence of supernumerary teeth can be as high as three percent, with the permanent dentition being affected five times more frequently than the primary dentition and males being affected twice as frequently as females.²¹

Supernumerary teeth will occur 10 times more often in the maxillary arch versus the mandibular arch.²¹ Approximately 90 percent of all single tooth supernumerary teeth are found in the maxillary arch, with a strong predilection to the anterior region.^{21,42} The maxillary anterior midline is the most common site, in which case the supernumerary tooth is known as a mesiodens; the second most common site is the maxillary molar area, with the tooth known as a paramolar.^{21,42} A mesiodens can be suspected if there is an asymmetric eruption pattern of the maxillary incisors, delayed eruption of the maxillary incisors with or without any overretained primary incisors, or ectopic eruption of a maxillary incisor.⁴⁵ The diagnosis of a mesiodens can be confirmed with radiographs, including occlusal, periapical, or panoramic films,⁴⁶ or computed tomography.^{9,10} Three-dimensional information needed to determine the location of the mesiodens or impacted tooth can be obtained by taking two periapical radiographs using either two projections taken at right angles to one another or the tube-shift technique (buccal object rule or Clark's rule)⁴⁷ or by cone beam computed tomography.^{10,12,13}

Complications of supernumerary teeth can include delayed and/or lack of eruption of the permanent tooth, crowding, resorption of adjacent teeth, dentigerous cyst formation, pericoronal space ossification, and crown resorption.^{42,48} Early diagnosis and appropriately timed treatment are important in the prevention and avoidance of these complications. Because only 25 percent of all mesiodens erupt spontaneously, surgical management often is necessary.^{44,49} A mesiodens that is conical in shape and is not inverted has a better chance for eruption than a mesiodens that is tubular in shape and is inverted.⁴⁸ The treatment objective for a nonerupting permanent mesiodens is to minimize eruption problems for the permanent incisors.⁴⁸ Surgical management will vary depending on the size, shape, and number of supernumeraries and the patient's dental development.⁴⁸ The treatment objective for a nonerupting primary mesiodens differs in that the removal of these teeth usually is not recommended, as the surgical intervention may disrupt or damage the underlying developing permanent teeth.⁵⁰ Erupted primary tooth mesiodens typically are left

to shed normally upon the eruption of the permanent dentition.⁵⁰

Extraction of an unerupted primary or permanent mesiodens is recommended during the mixed dentition to allow the normal eruptive force of the permanent incisor to bring itself into the oral cavity.⁴³ Waiting until the adjacent incisors have at least two-thirds root development will present less risk to the developing teeth but still allow spontaneous eruption of the incisors.³ In 75 percent of the cases, extraction of the mesiodens during the mixed dentition results in spontaneous eruption and alignment of the adjacent teeth.^{50,51} If the adjacent teeth do not erupt within six to 12 months, surgical exposure and orthodontic treatment may be necessary to aid their eruption.^{45,47}

Frenulum attachments

Frenulum attachments and their role in oral function increasingly have become topics of interest among a variety of health care specialists. Ankyloglossia (tongue-tie) and hypertrophic/restrictive maxillary frenula have been implicated in difficulties breastfeeding⁵³, incorrect speech articulation^{54,55}, caries formation^{56,57}, gingival recession⁵⁸, and aberrant skeletal growth⁵⁹. Studies have shown differences in treatment recommendations among pediatricians, otolaryngologists, lactation consultants, speech pathologists, surgeons, and dental specialists.^{54,60-66} Clear indications and timing of surgical treatment remain controversial due to lack of consensus regarding accepted anatomical and diagnostic criteria for degree of restriction and relative impact on growth, development, feeding, or oral motor function.^{54,60-66}

When indicated, frenuloplasty/frenotomy (various methods to release the frenulum and correct the anatomic situation) or frenectomy (simple cutting of the frenulum) may be a successful approach to alleviate the problem.^{54,60,65,67} Each of these procedures involves surgical incision, establishing hemostasis, and wound management.⁶⁸ Dressing placement or the use of antibiotics is not necessary.⁶⁸ Recommendations include maintaining a soft diet, regular oral hygiene, and analgesics as needed.⁶⁹ The use of electrosurgery or laser technology for frenectomies has demonstrated a shorter operative working time, a better ability to control bleeding, reduced intra- and postoperative pain and discomfort, fewer postoperative complications (e.g., swelling, infection), no need for suture removal, and increased patient acceptance.^{62,69,70} These procedures require extensive training as well as skillful technique and patient management.^{54,60,65,67,71-75}

Pediatric oral pathology

A wide spectrum of oral lesions occurs in children and adolescents, including soft and hard tissue lesions of the oral maxillofacial region. There is limited information on the prevalence of oral lesions in the pediatric population. The largest epidemiologic studies in the United States place the prevalence rate in children at four to 10 percent with the exclusion of infants.^{76,77} Although the vast majority of these lesions represent mucosal conditions, developmental anomalies, and reactive

or inflammatory lesions, it is imperative to be vigilant for neoplastic diseases.

Regardless of the age of the child, it is important to establish a working diagnosis for every lesion. This is based on obtaining a thorough history, assessing the risk factors and documenting the clinical signs and symptoms of the lesion. Based on these facts, a list of lesions with similar characteristics is rank ordered from most likely to least likely diagnosis. The entity that is judged to be the most likely disease becomes the working diagnosis and determines the initial management approach.

For most oral lesions, a definitive diagnosis is best made by performing a biopsy. By definition, a biopsy is the removal of a piece of tissue from a living body for diagnostic study and is considered the gold standard of diagnostic tests.⁷⁸ The two most common biopsies are the incisional and excisional types. Excisional biopsies usually are performed on small lesions, less than one centimeter in size, for the total removal of the affected tissue. An incisional biopsy is performed when a malignancy is suspected, the lesion is large in size or diffuse in nature, or a multifocal distribution is present. Multiple incisional biopsies may be indicated for diffuse lesions in order to obtain a representative tissue sample. Fine needle aspiration, the cytobrush technique, and exfoliative cytology may assist in making a diagnosis, but they are considered adjunctive tests because they do not establish a definitive diagnosis.^{79,80}

It is considered the standard of care that any tissue removed from the oral and maxillofacial region be submitted for histopathologic examination.⁸¹ Exceptions to this rule include carious teeth that do not have soft tissue attached, extirpated pulpal tissue, and clinically normal tissue, such as tissue from gingival recontouring.⁸¹ Gross description of all tissue that is removed should be entered into the patient record. In general, a soft tissue biopsy should be performed when a lesion persists for greater than two weeks despite removal of the suspected causative factor or empirical drug treatment. It is also imperative to submit hard or soft tissue for evaluation to a pathologist if the differential diagnosis includes at least one significant disease or neoplasm. Histopathologic examination not only furnishes a definitive diagnosis, but it provides information about the clinical behavior and prognosis and determines the need for additional treatment or follow-up. Another valuable outcome is that it allows the clinician to deliver evidence-based medical/dental care, increasing the likelihood for a positive result.⁷⁸ Furthermore, it presents important documentation about the lesion for the patient record, including the procedures taken for establishing a diagnosis.⁷⁸

Many oral biopsies are within the scope of practice for a pediatric dentist to perform. If the tissue is excised, the following steps should be taken for optimum results:⁷⁸⁻⁸¹

1. select the most representative lesion site and not the area that is the most accessible.
2. remove an adequate amount of tissue. If the biopsy is too small or too superficial, a diagnosis may be compromised.
3. avoid crushing or distorting the tissue. Damage is most often observed from the forces of the tissue forceps, tearing the tissues or overheating the tissue from the use of electrosurgery or laser removal.
4. immediately place the tissue in a fixative, which for most samples is 10 percent formalin. It is critical not to dilute the fixative with water or other liquids because tissue autolysis will render the sample nondiagnostic.
5. proper identification of the specimen is essential. The formalin container should be labelled with the name of the patient and the location. Multiple tissue samples from different locations should not be placed in the same container, unless they are uniquely identified, such as tagged with a suture.
6. complete the surgical pathology form including patient demographics, the submitting dentist's name and address, and a brief but accurate history. It is important to have legible records so that the diagnosis is not delayed. Clinical photographs and radiographs often are very useful for correlating the microscopic findings.

Worldwide, the most frequently oral biopsied lesions in children include⁸²:

- mucocele;
- fibrous lesions;
- pyogenic granuloma;
- dental follicle;
- human papillomavirus (HPV) lesion;
- chronic inflammation;
- giant cell lesions (soft tissue);
- hyperkeratosis;
- peripheral ossifying fibroma;
- gingivitis;
- gingival hyperplasia;
- hemangioma;
- ulcer;
- lymphangioma;
- sialadenitis;
- Burkitt's lymphoma;
- melanotic macule;
- pleomorphic adenoma;
- nevus; and
- neurofibroma.

Lesions of the newborn

Palatal cysts of the newborn include Epstein pearls and Bohn nodules. These cysts are found in up to 85 percent of newborns.^{53,83-90} Epstein pearls occur in the median palatal raphe area^{53,83-85} as a result of trapped epithelial remnants along the line of fusion of the palatal halves.^{49,51} Bohn nodules are remnants of salivary gland epithelium and usually are found on the buccal and lingual aspects of the ridge, away from the midline.^{83,85} Gingival cysts of the newborn, or dental lamina cysts, are found on the crests of the dental ridges, and are most commonly are seen bilaterally in the region of the first primary

molars.⁸⁴ They result from remnants of the dental lamina. Palatal and gingival cysts of the newborn typically present as asymptomatic one to three millimeter nodules or papules. They are smooth, whitish in appearance, and filled with keratin.^{83,84} No treatment is required, as these cysts usually disappear during the first three months of life.^{53,83}

Congenital epulis of the newborn

Congenital epulis of the newborn, also known as granular cell tumor or Neumann's tumor, is a rare benign tumor seen only in newborns.⁹¹ This lesion is typically a protuberant mass arising from the gingival mucosa. It is most often found on the anterior maxillary ridge.^{86,87} Patients typically present with feeding and/or respiratory problems.⁸⁷ Congenital epulis has a marked predilection for females at 8:1 to 10:1.⁸⁶⁻⁸⁸ Treatment normally consists of surgical excision.⁸⁶⁻⁸⁸ The newborn usually heals well, and no future complications or treatment should be expected. Congenital epulis never recurs after excision.⁸⁸ There have been reports of spontaneous regression of untreated congenital epulis.^{88,91}

Melanotic neuroectodermal tumor of infancy

Melanotic neuroectodermal tumor of infancy is a rare occurrence that develops during the first year of life.⁸⁸ This lesion may be present at birth. It occurs in the anterior maxilla 70 percent of the time.⁸³ Less frequently, melanotic neuroectodermal tumor of infancy occurs in the skull, mandible, epididymis and testis, and brain.^{83,88} The classic presentation is a bluish or black rapidly expanding mass of the anterior maxilla. Radiographic findings include an ill-defined unilocular radiolucency with the displacement of tooth buds.⁸⁸ There can be a floating tooth appearance.⁸³ Surgical excision is required, and there is a 20 percent recurrence rate. Although this is a benign lesion, seven percent of reported cases have behaved malignantly resulting in metastasis and death.⁸⁸

Natal and neonatal teeth

Natal and neonatal teeth can present a challenge when deciding on appropriate treatment. Natal teeth have been defined as those teeth present at birth, and neonatal teeth are those that erupt during the first 30 days of life.^{92,93} The occurrence of natal and neonatal teeth is rare; the incidence varies from 1:1,000 to 1:30,000.^{92,93} The teeth most often affected are the mandibular primary incisors.⁹⁴ In most cases, anterior natal and neonatal teeth are part of the normal complement of the dentition.^{92,93} Natal or neonatal molars have been identified in the posterior region and may be associated with systemic conditions or syndromes (e.g., Pfeiffer syndrome, histiocytosis X).⁹⁴⁻⁹⁶ Although many theories exist as to why the teeth erupt prematurely, currently no studies confirm a causal relationship with any of the proposed theories. The superficial position of the tooth germ associated with a hereditary factor seems to be the most accepted possibility.⁹³

If the tooth is not excessively mobile or causing feeding problems, it should be preserved and maintained in a healthy

condition if possible.^{93,95,96} Close monitoring is indicated to ensure that the tooth remains stable and is not an aspiration risk to the infant.

Riga-Fede disease is a condition caused by the natal or neonatal tooth rubbing the ventral surface of the tongue during feeding, leading to ulceration.^{75,92} Failure to diagnose and properly treat this lesion can result in dehydration and inadequate nutritional intake for the infant.⁹⁶ Treatment should be conservative and focus on creating round, smooth incisal edges.⁹³⁻⁹⁶ If conservative treatment does not correct the condition, extraction is the treatment of choice.⁹³⁻⁹⁶

An important consideration when deciding to extract a natal or neonatal tooth is the potential for hemorrhage. Extraction is contraindicated in newborns due to risk of hemorrhage.⁹⁷ Unless the child is at least 10 days old, consultation with the pediatrician regarding adequate hemostasis may be indicated prior to extraction of the tooth. In particular, infants may be at risk for vitamin K deficiency bleeding (VKDB) if they did not receive a dose of vitamin K shortly after birth (within six hours of birth).⁹⁸ Infants can be at risk for VKDB until the age of six months if they do not receive a vitamin K injection.⁹⁸

Lesions occurring in children and adolescents

Eruption cyst (eruption hematoma)

The eruption cyst is a soft tissue cyst that results from a separation of the dental follicle from the crown of an erupting tooth.^{83,99} Fluid accumulation occurs within this created follicular space.^{85,89,100} Eruption cysts most commonly are found in the mandibular molar region.⁸⁹ Color of these lesions can range from normal to blue-black or brown, depending on the amount of blood in the cystic fluid.^{85,89,100} The blood is secondary to trauma. If trauma is intense, these blood-filled lesions sometimes are referred to as eruption hematomas.^{85,89,100} Because the tooth erupts through the lesion, no treatment is necessary.^{85,89,100} If the cyst does not rupture spontaneously or the lesion becomes infected, the roof of the cyst may be opened surgically.^{85,89}

Mucocele

The mucocele is a common lesion in children and adolescents resulting from the rupture of a minor salivary gland excretory duct, with subsequent leakage of mucin into the adjacent connective tissues that later may be surrounded in a fibrous capsule.^{83,85,99-101} Most mucoceles are well-circumscribed bluish translucent fluctuant swellings that are firm to palpation, although deeper and long-standing lesions may range from normal in color to having a whitish keratinized surface.^{85,99,100} Mucoceles most frequently are observed on the lower lip, usually lateral to the midline.⁸⁸ Mucoceles also can be found on the buccal mucosa, ventral surface of the tongue, retromolar region, and floor of the mouth (ranula).⁹⁹⁻¹⁰¹ Superficial mucoceles and some other mucoceles are short-lived lesions that burst spontaneously, leaving shallow ulcers that heal within a few days.^{85,100} Local mechanical trauma to the minor salivary gland is often the cause of rupture.^{50,53,86,87} Many lesions, however, require treatment to minimize the risk of recurrence.^{85,100}

Squamous papilloma

Squamous papilloma is a benign lesion caused by HPV types 1 and 6.⁸³ Squamous papilloma presents as soft painless, pink to white, pedunculated (stalked) lesions. The surface may display multiple fingerlike projections and may have a cauliflower like appearance.^{83,102,103} These lesions can occur anywhere in the oral cavity, but the tongue, lips, and soft palate are the most common sites.¹⁰³ Squamous papilloma generally occurs in adulthood, but 20 percent have been noted prior to age 20.⁹¹ Although they are viral in origin, the infectivity is low.^{83,102} Squamous papilloma do not have malignant potential.¹⁰³ Excision is the treatment of choice, and recurrence is uncommon.^{83,103}

Verruca vulgaris

Verruca vulgaris, or the common wart, is a lesion induced by HPV type 2 and generally found on the skin of the hand.¹⁰² Finger or thumb sucking can cause autoinoculation resulting in the development of intraoral lesions.¹⁰² Verruca vulgaris is similar in appearance to the squamous papilloma. This lesion can be sessile (broad based) or pedunculated and can display a rough bumpy surface.¹⁰³ Verruca vulgaris can be found on the lips, tip of tongue, and labial mucosa.⁸³ There is no risk of malignant transformation.⁸³ Excision of the entire lesion is recommended and recurrence is uncommon.¹⁰³

Irritation fibroma

The irritation fibroma is a reactive lesion occurring as a response to chronic trauma of the mucosa. The irritation fibroma presents as a firm nontender pink nodule and is composed of fibrous connective tissue.¹⁰³ The lesion does not exceed two millimeters in diameter.¹⁰⁴ The irritation fibroma can be found on buccal and labial mucosa, the tongue, and attached gingiva. Excisional biopsy is recommended. These can reoccur if the source of the irritation is not removed.^{103,104}

Recurrent aphthous stomatitis

Recurrent aphthous stomatitis is one of the most common oral lesions, occurring in 20-30 percent of children.⁸³ Recurrent aphthous stomatitis is caused by a T-cell mediated immunologic reaction to a triggering agent.¹⁰⁵ Three variants of aphthous ulcers are recognized:

1. Minor aphthous ulcerations. Minor aphthous ulcerations are the most common form, accounting for almost 80 percent of aphthous ulcers.¹⁰⁵ They have a yellowish-white membrane and are surrounded by an erythematous halo. These ulcers are three to 10 millimeters in diameter. Minor aphthous ulcers occur on nonkeratinized mucosa.¹⁰⁵ One to five ulcers often present during a single outbreak, and they heal in seven to 14 days without scarring.¹⁰⁶
2. Major aphthous ulcerations. Major aphthous ulcerations are larger and deeper and have a longer duration than the minor aphthous ulcer. These occur most commonly

on the labial mucosa, soft palate, and the tonsillar fauces.¹⁰⁵ The major aphthous ulcer can take up to six weeks to heal with potential scarring.¹⁰⁵

3. Herpetiform aphthous ulcerations. Herpetiform aphthous ulcerations can occur on any intraoral site.¹⁰⁶ As many as 100 small ulcerations can be present in a single occurrence.¹⁰⁵ The ulcerations may resemble primary herpetic stomatitis. These ulcerations may coalesce to form a larger ulceration.¹⁰⁵ Herpetiform aphthous ulcers heal within seven to 10 days, but recurrences are frequent.¹⁰⁶

Aphthous ulcers may be treated with topical anesthetics for relief of pain. Topical and systemic steroids, chlorhexidine rinses, and laser treatments can be used to manage these lesions.⁸³

Localized juvenile spongiotic gingival hyperplasia

Localized juvenile spongiotic gingival hyperplasia was originally known as puberty gingivitis.¹⁰⁷ It is thought to be an isolated patch of sulcular or junctional epithelium that is subjected to local factors such as mouth breathing or orthodontic appliances.⁸³ The lesion presents as an isolated bright red velvety patch or enlargement of anterior facial gingiva. This lesion bleeds easily and does not respond to oral hygiene measures. There is a female predilection.⁸³ Most lesions occur under the age of 20, with the median age at diagnosis being 12 years.¹⁰⁷ Excision is the treatment of choice, and up to 16 percent will recur.⁸³

Pyogenic granuloma

Pyogenic granuloma is a painless smooth or lobulated vascular lesion. The pyogenic granuloma is usually ulcerated and bleeds easily.^{83,107} This lesion can occur at any age but is most common in children and young adults. There is a female predilection,⁸³ and the pyogenic granuloma can occur in up to five percent of pregnancies.¹⁰⁸ The pyogenic granuloma is thought to be an exuberant tissue response to a local irritant or trauma.¹⁰⁸ Pyogenic granuloma most commonly occurs on maxillary anterior attached gingiva (75 percent) but can be found on tongue, lower lip, or buccal mucosa.¹⁰⁸ Treatment is complete excision with the removal of the source of irritant.^{83,108} This lesion can recur in three to 15 percent of cases.⁸³

References

1. American Academy of Pediatric Dentistry. Pediatric oral surgery. *Pediatr Dent* 2005;27(Suppl):158-64.
2. American Academy of Pediatric Dentistry. Management considerations for pediatric oral surgery and oral pathology. *Pediatr Dent* 2015;37(special issue):279-88.
3. American Association of Oral and Maxillofacial Surgeons. Dentoalveolar Surgery. In: Parameters of Care: Clinical practice guidelines for oral and maxillofacial surgery (AAOMS ParCare 2017 Ver 6). *J Oral Maxillofac Surg* 2017;75(8)Suppl 1:e50-73.

References continued on the next page.

4. American Academy of Pediatric Dentistry. Informed consent. *The Reference Manual of Pediatric Dentistry*. Chicago, Ill.: American Academy of Pediatric Dentistry; 2020:470-3.
5. Adewumi AO. Oral surgery in children. In: Nowak AJ, Christensen JR, Mabry TR, Townsend JA, Wells MH, eds. *Pediatric Dentistry Infancy through Adolescence*. 6th ed. St Louis, Mo.: Elsevier; 2019:399-409.
6. Kaban L, Troulis M. Preoperative assessment of the pediatric patient. In: *Pediatric Oral and Maxillofacial Surgery*. Philadelphia, Pa.: Saunders; 2004:3-19.
7. American Academy of Pediatric Dentistry. Prescribing dental radiographs for infants, children, adolescents, and persons with special health care needs. *The Reference Manual of Pediatric Dentistry*. Chicago, Ill.: American Academy of Pediatric Dentistry; 2020:248-51.
8. Murray DJ, Chong DK, Sandor GK, Forrest CR. Denti-gerous cyst after distraction osteogenesis of the mandible. *J Craniofac Surg* 2007;18(16):1349-52.
9. Ramesh A. Panoramic imaging. In: Mallya SM, Lam WN, eds. *Oral Radiology: Principles and Interpretation*. 8th ed. St. Louis, Mo.: Elsevier; 2019:132-50.
10. Scarfe WC, Farman AG. Cone-beam computed tomography volume preparation. In: Mallya SM, Lam WN, eds. *White and Pharoah's Oral Radiology: Principles and Interpretation*. 8th ed. St. Louis, Mo.: Elsevier; 2019:165-80.
11. Mallya SM. Other imaging modalities. In: Mallya SM, Lam WN, eds. *White and Pharoah's Oral Radiology: Principles and Interpretation*. 8th ed. St. Louis, Mo.: Elsevier; 2019:2218-38.
12. Katheria BC, Kau CH, Tate R, Chen JW, English J, Bouquet J. Effectiveness of impacted and supernumerary tooth diagnosis from traditional radiography versus cone beam computed tomography. *Pediatr Dent* 2010;32(4):304-9.
13. Serrant PS, McIntyre GT, Thomson DJ. Localization of ectopic maxillary canines—Is CBCT more accurate than conventional horizontal or vertical parallax? *J Orthod* 2014; 41(1):13-8.
14. Ferneini EM, Bennett JD. Oral surgery for the pediatric patient. In: Dean JA, ed. *McDonald and Avery's Dentistry for the Child and Adolescent*, 10th ed. St Louis, Mo.: Elsevier; 2016:627-44.
15. American Academy of Pediatric Dentistry. Use of anesthesia providers in the administration of office-based deep sedation/general anesthesia to the pediatric dental patient. *The Reference Manual of Pediatric Dentistry*. Chicago, Ill.: American Academy of Pediatric Dentistry; 2019:327-30.
16. Kaban L, Troulis M. Deep sedation for pediatric patients. In: *Pediatric Oral and Maxillofacial Surgery*. Philadelphia, Pa.: Saunders; 2004:86-99.
17. Kaban L, Troulis M. Infections of the maxillofacial region. In: *Pediatric Oral and Maxillofacial Surgery*. Philadelphia, Pa.: Saunders; 2004:171-86.
18. Seow W. Diagnosis and management of unusual dental abscesses in children. *Aust Dent J* 2003;43(3):156-68.
19. Baker S, Parico L. Pathologic paediatric conditions associated with a compromised airway. *Int J Paediatr Dent* 2010; 20(2):102-11.
20. Thikkurissy S, Rawlins JT, Kumar A, et al. Rapid treatment reduces hospitalization for pediatric patients with odontogenic-based cellulitis. *Am J Emerg Med* 2010;28 (6):668-72.
21. Regezi J, Sciubba J, Jordan R. Abnormalities of teeth. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:373-88.
22. Mochizuki K, Ohtawa Y, Kubo S, Machida Y, Yakushiji M. Bifurcation, bi-rooted primary canines: A case report. *Int J Paediatr Dent* 2001;11(5):380-5.
23. Andersson L, Blomlöf L, Lindskog S, Feiglin B, Hammarström L. Tooth ankylosis. Clinical, radiographic and histological assessments. *Int J Oral Surg* 1984;13(5): 423-31.
24. American Academy of Pediatric Dentistry. Management of developing dentition and occlusion in pediatric dentistry. *The Reference Manual of Pediatric Dentistry*. Chicago, Ill.: American Academy of Pediatric Dentistry; 2020: 393-409.
25. Tieu LD, Walker SL, Major MP, Flores-Mir C. Management of ankylosed primary molars with premolar successors: A systematic review. *J Am Dent Assoc* 2013;144(6):602-11.
26. O'Connell AC, Torske KR. Primary failure of tooth eruption: A unique case. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 1999;87(6):714-20.
27. Frazier-Bowers SA, Koehler KE, Ackerman JL, Proffit WR. Primary failure of eruption: Further characterization of a rare eruption disorder. *Am J Orthod Dentofacial Orthop* 2007;131(5):578, e1-11.
28. Raghoebar GM, Boering G, Vissink A. Clinical, radiographic and histological characteristics of secondary retention of permanent molars. *J Dent* 1991;19(3):164-70.
29. Frazier-Bowers SA, Puranik CP, Mahaney MC. The etiology of eruption disorders – Further evidence of a 'genetic paradigm'. *Semin Orthod* 2010;16(3):180-5.
30. Rhoads SG, Hendricks HM, Frazier-Bowers SA. Establishing the diagnostic criteria for eruption disorders based on genetic and clinical data. *Am J Orthod Dentofacial Orthop* 2013;114(2):194-202.
31. Sharif MO, Parker K, Lyne A, Chia MSY. The orthodontic-oral surgery interface part two: Diagnosis and management of anomalies in eruption and transpositions. *Brit Dent J* 2018;225(6):491-6.
32. Ericson S, Kurol J. Early treatment of palatally erupting maxillary canines by extraction of the primary canines. *Eur J Orthod* 1988;10(4):283-95.
33. Richardson G, Russel K. A review of impacted permanent maxillary cuspids – Diagnosis and prevention. *J Can Dent Assoc* 2000;66(9):497-501.
34. Parkin N, Benson P, Shah A, et al. Extraction of primary (baby) teeth for unerupted palatally displaced permanent canine teeth in children. *Cochrane Database Syst Rev* 2009;15(2):CD004621.
35. Naoumova J, Kurol J, Kjellberg H. A systematic review of the interceptive treatment of palatally displaced maxillary canines. *Eur J Orthod* 2011;33(2):143-9.
36. Bazargani F1, Magnuson A, Lennartsson B. Effect of interceptive extraction of deciduous canine on palatally displaced maxillary canine: A prospective randomized controlled study. *Angle Orthod* 2014;84(1):3-10.
37. Song F, O'Meara S, Wilson P, Goldner S, Kleijnen J. The effectiveness and cost-effectiveness of prophylactic removal of wisdom teeth. *Health Technol Assess* 2000;4(1):1-55.
38. Haug R, Perrott D, Gonzalez M, Talwar R. The American Association of Oral and Maxillofacial Surgeons age-related third molar study. *J Oral Maxillofac Surg* 2005;63(8): 1106-14.

39. Pogrel M, Dodson T, Swift J, et al. White paper on third molar data. American Association of Oral and Maxillofacial Surgeons. March 2007. Available at: "https://www.aaoms.org/docs/govt_affairs/advocacy_white_papers/white_paper_third_molar_data.pdf". Accessed July 24, 2020.
40. Friedman JW. The prophylactic extraction of third molars: A public health hazard. *Am J Public Health* 2007;97(9):1554-9.
41. Almendros-Marques N, Alaejos-Algarra E, Quinteros-Borgarello M, Berini-Aytes L, Gay-Escoda C. Factors influencing the prophylactic removal of asymptomatic impacted lower third molars. *Int J Oral Maxillofac Surg* 2008;37(1):29-35.
42. Neville BW, Damm DD, Allen CM, Chi AC. Abnormalities of the teeth. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:49-110.
43. Dean JA. Managing the developing occlusion. In: McDonald and Avery's *Dentistry for the Child and Adolescent*. 10th ed. St. Louis, Mo.: Elsevier; 2016:415-78.
44. Taylor GS. Characteristics of supernumerary teeth in the primary and permanent dentition. *Trans Br Soc Study Orthod* 1970-71;57:123-8.
45. Primosch R. Anterior supernumerary teeth—Assessment and surgical intervention in children. *Pediatr Dent* 1981;3(2):204-15.
46. Tadinada A, Potluri. Dental anomalies. In: Mallya SM, Lam EW, eds. *White and Pharoah's Oral Radiology: Principles and Interpretation*. 8th ed. St. Louis, Mo.: Elsevier; 2019:335-63.
47. Mallya SM, White S, Pharoah M. Projection geometry. In: Mallya SM, Lam MN, eds. *White and Pharoah's Oral Radiology: Principles and Interpretation*. 8th ed. St. Louis, Mo.: Mosby Elsevier; 2019:81-8.
48. Christensen JR, Fields HW Jr., Sheats RD. Treatment planning and management of orthodontic problems. In: Nowak AJ, Christensen JR, Mabry TR, Townsend JA, Wells MH, eds. *Pediatric Dentistry: Infancy through Adolescence*. 6th ed. Philadelphia, Pa.: Elsevier; 2019:512-53.
49. Neville BW, Damm DD, White DK. Pathology of the teeth. In: *Color Atlas of Clinical Oral Pathology*. 2nd ed. Baltimore, Md.: Williams & Wilkins; 2003:58-60.
50. Russell K, Folwaczna M. Mesiodens: Diagnosis and management of a common supernumerary tooth. *J Can Dent Assoc* 2003;69(6):362-6.
51. Howard R. The unerupted incisor. A study of the post-operative eruptive history of incisors delayed in their eruption by supernumerary teeth. *Dent Pract Dent Rec* 1967;17(9):332-41.
52. Giancotti A, Grazzini F, De Dominicis F, Romanini G, Arcuri C. Multidisciplinary evaluation and clinical management of mesiodens. *J Clin Pediatr Dent* 2002;26(3):233-7.
53. Neville BW, Damm DD, Allen CM, Chi AC. Developmental defects of the oral and maxillofacial region. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:1-48.
54. Suter VG, Bornstein MM. Ankyloglossia: Facts and myths in diagnosis and treatment. *J Periodontol* 2009;80(8):1204-19.
55. Webb AN, Hao W, Hong P. The effect of tongue-tie division on breastfeeding and speech articulation: A systematic review. *Int J Pediatr Otorhinolaryngol* 2013;77(5):635-46.
56. Coryllos E, Genna CW, Salloum A. Congenital tongue-tie and its impact on breastfeeding. *Breastfeeding: Best for baby and mother. Am Acad Pedia (newsletter)* 2004; Summer:1-7.
57. Kotlow L. The influence of the maxillary frenum on the development and pattern of dental caries on anterior teeth in breastfeeding infants: Prevention, diagnosis, and treatment. *J Hum Lact* 2010;26(3):304-8.
58. John J, Weddell JA, Shin DE, Jones JJ. Gingivitis and periodontal disease. In: JA Dean, ed. *McDonald and Avery's Dentistry for the Child and Adolescent*, 10th ed. St. Louis, Mo.; Elsevier; 2016:243-73.
59. Geddes D, Langton D, Gollow I, Jacobs L, Hartmann P, Simmer K. Frenulotomy for breastfeeding infants with ankyloglossia: Effect on milk removal and sucking mechanism as imaged by ultrasound. *Pediatrics* 2008;122(1):e188-e194.
60. Segal L, Stephenson R, Dawes M, Feldman P. Prevalence, diagnosis, and treatment of ankyloglossia. *Can Fam Physician* 2007;53(6):1027-33.
61. Boutsis EZ, Tatakis DN. Maxillary labial frenum attachment in children. *Int J Paediatr Dent* 2011;21(4):284-8.
62. Kotlow L. Diagnosing and understanding the maxillary lip-tie (superior labial, the maxillary labial frenum) as it relates to breastfeeding. *J Hum Lact* 2013;29(4):458-64.
63. O'Callahan C, Macary S, Clemente S. The effects of office-based frenotomy for anterior and posterior ankyloglossia on breastfeeding. *Int J Pediatr Otorhinolaryngol* 2013;77(5):827-32.
64. Finigan V, Long T. The effectiveness of frenulotomy on infant-feeding outcomes: A systemic literature review. *Evid Based Midwifery* 2013;11(2):40-5.
65. Webb AN, Hao W, Hong P. The effect of tongue-tie division on breastfeeding and speech articulation: A systematic review. *Int J Pediatr Otorhinolaryngol* 2013;77(5):635-46.
66. Delli K, Livas C, Sculean A, Katsaros C, Bornstein MM. Facts and myths regarding the maxillary midline frenum and its treatment: A systematic review of the literature. *Quintessence Int* 2013;44(2):177-87.
67. Devishree G, Gujjari SK, Shubhashini PV. Frenectomy: A review with the reports of surgical techniques. *J Clin Dent Res* 2012;6(9):1587-92.
68. Kaban L, Troulis M. Intraoral soft tissue abnormalities. In: *Pediatric Oral and Maxillofacial Surgery*. Philadelphia, Pa.: Saunders; 2004:147-53.
69. Shetty K, Trajtenberg C, Patel C, Streckfus C. Maxillary frenectomy using a carbon dioxide laser in a pediatric patient: A case report. *Gen Dent* 2008;56(1):60-3.
70. Olivi G, Chaumanet G, Genovese MD, Beneduce C, Andreana S. Er,Cr:YSGG laser labial frenectomy: A clinical retrospective evaluation of 156 consecutive cases. *Gen Dent* 2010;58(3):e126-33.
71. Kupietzky A, Botzer E. Ankyloglossia in the infant and young child: Clinical suggestions for diagnosis and management. *Pediatr Dent* 2005;27(1):40-6.
72. Hogan M, Wescott C, Griffiths M. Randomized, controlled trial of division of tongue-tie in infants with feeding problems. *J Paediatr Child Health* 2005;41(5-6):246-50.
73. Díaz-Pizán M, Lagravère M, Villena R. Midline diastema and frenum morphology in the primary dentition. *J Dent* 2006;26(1):11-14.

References continued on the next page.

74. Gontijo I, Navarro R, Haypek P, Ciamponi A, Hadda A. The applications of diode and Er:YAG lasers in labial frenectomy in infant patients. *J Dent Child* 2005;72(1):10-5.
75. Kara C. Evaluation of patient perceptions of frenectomy: A comparison of Nd:YAG laser and conventional techniques. *Photomed Laser Surg* 2008;26(2):147-52.
76. Kleinman DV, Swango PA, Pindborg JJ. Epidemiology of oral mucosal lesions in United States school children: 1986-87. *Community Dent Oral Epidemiol* 1994;22(4):243-53.
77. Shulman JD. Prevalence of oral mucosal lesions in children and youths in USA. *Int J Pediatr Dent* 2005;15(2):89-97.
78. Melrose RJ, Handlers JP, Kerpel S, Summerlin DJ, Tomich CJ. The use of biopsy in dental practice. The position of the American Academy of Oral and Maxillofacial Pathology. *Gen Dent* 2007;55(5):457-61.
79. Rethman M, Carpenter W, Cohen E, et al. Evidence-based clinical recommendations on screening for oral squamous cell carcinomas. *J Am Dent Assoc* 2010;141(5):509-20.
80. Kazanowska K, Halon A, Radwan-Oczko M. The role and application of exfoliative cytology in the diagnosis of oral mucosa pathology – Contemporary knowledge with review of the literature. *Adv Clin Exp Med* 2014;23(2):299-305.
81. American Academy of Oral and Maxillofacial Pathology. Submission policy on excised tissue. Available at: "http://www.aaomp.org/wp-content/uploads/2016/12/Policy_on_Excised_Tissue-Final-11-9-2013.pdf". Accessed July 25, 2020.
82. Hong C, Dean D, Hull K, et al. World workshop on oral medicine: VII: Relative frequency of oral mucosal lesions in children, a scoping review. *Oral Diseases* 2019;25(Suppl. 1):193-203.
83. Flaitz CM. Differential diagnosis of oral lesions and developmental anomalies. In: Nowak AJ, Christensen JR, Mabry TR, Townsend JA, Wells MH, eds. *Pediatric Dentistry: Infancy through Adolescence*. 6th ed. Philadelphia, Pa.: Elsevier; 2019:8-49.
84. Hays P. Hamartomas, eruption cysts, natal tooth, and Epstein pearls in a newborn. *ASDC J Dent Child* 2000;67(5):365-8.
85. Aldred MJ, Cameron AC, Georgiou A. Pediatric oral medicine and pathology and radiology. In: Cameron AC, Widmer RP, eds. *Handbook of Pediatric Dentistry*. 4th ed. Philadelphia, Pa.: Mosby Elsevier; 2013:209-68.
86. Lapid O, Shaco-Levey R, Krieger Y, Kachko L, Sagi A. Congenital epulis. *Pediatrics* 2001;107(2):E22.
87. Marakoglu I, Gursoy U, Marakoglu K. Congenital epulis: Report of a case. *ASDC J Dent Child* 2002;69(2):191-2.
88. Neville BW, Damm DD, Allen CM, Chi AC. Soft tissue tumors. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:473-515.
89. Neville BW, Damm DD, Allen CM, Chi AC. Odontogenic cysts and tumors. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:632-89.
90. Regezi JA, Sciubba JJ, Jordan RC. Cysts of the jaws and neck. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:245-68.
91. McDonald JS. Tumors of the oral soft tissues and cysts and tumors of bone. In: Dean JA, ed. *McDonald and Avery's Dentistry for the Child and Adolescent*. 10th ed. St. Louis, Mo.: Elsevier; 2016:603-26.
92. Cunha RF, Boer FA, Torriani DD, Frossard WT. Natal and neonatal teeth: Review of the literature. *Pediatr Dent* 2001;23(2):158-62.
93. Leung A, Robson W. Natal teeth: A review. *J Natl Med Assoc* 2006;98(2):226-8.
94. Galassi MS, Santos-Pinto L, Ramalho T. Natal maxillary primary molars: Case report. *J Clin Pediatr Dent* 2004;29(1):41-44.
95. Stein S, Paller A, Haut P, Mancini A. Langerhans cell histiocytosis presenting in the neonatal period: A retrospective case series. *Arch Pediatr Adolesc Med* 2001;155(7):778-83.
96. Slayton RL. Treatment alternatives for sublingual traumatic ulceration (Riga-Fede disease). *Pediatr Dent* 2000;22(5):413-4.
97. Rushmah M. Natal and neonatal teeth: A clinical and histological study. *J Clin Pediatr Dent* 1991;15(4):251-3.
98. Centers for Disease Control and Prevention. What is vitamin K deficiency bleeding? Available at: "<https://www.cdc.gov/ncbddd/vitamink/facts.html>". Accessed July 25, 2020.
99. Flaitz CM, Haberland C. Oral pathology and associated syndromes. In: Nowak AJ, Casamassimo PS, eds. *The Handbook: Pediatric Dentistry*. 5th ed. Chicago, Ill.: American Academy of Pediatric Dentistry; 2018:46-100.
100. Regezi J, Sciubba J, Jordan R. Salivary gland diseases. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:185-224.
101. Neville BW, Damm DD, Allen CM, Chi AC. Salivary gland pathology. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:422-72.
102. Regezi J, Sciubba J, Jordan R. Verrucal-papillary lesions. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:148-60.
103. Neville BW, Damm DD, Allen CM, Chi AC. Epithelial pathology In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:331-421.
104. Regezi J, Sciubba J, Jordan R. Connective tissue lesions. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:161-84.
105. Neville BW, Damm DD, Allen CM, Chi AC. Allergies and immunologic diseases. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:303-30.
106. Regezi J, Sciubba J, Jordan R. Ulcerative conditions. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:23-79.
107. Neville BW, Damm DD, Allen CM, Chi AC. Periodontal disease. In: *Oral and Maxillofacial Pathology*. 4th ed. St. Louis, Mo.: Elsevier; 2016:140-63.
108. Regezi J, Sciubba J, Jordan R. Red-blue lesions. In: *Oral Pathology: Clinical-Pathologic Correlations*. 7th ed. St. Louis, Mo.: Elsevier; 2017:114-33.