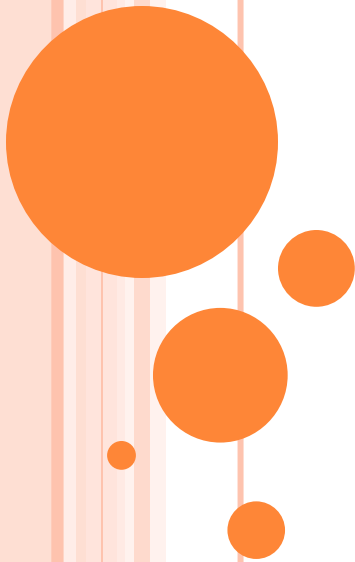


DEVELOPMENTAL DISTURBANCES CONTINUED...

**DEPTT OF ORAL PATHOLOGY &
MICROBIOLOGY**



PURPOSE STATEMENT

At the end of lecture the student should describe--

- ❑ Reactive lymphoid aggregate
- ❑ Lymphoid hamartoma
- ❑ Angiolymphoid hyperplasia with eosinophilia
- ❑ Lymphoepithelial cyst



LEARNING OBJECTIVES

S.N.	Learning Objectives	Domain	Level	Criteria	Condition
1	Enumerate clinical features	Cognitive	Must Know	All	
2	Write classification	Cognitive	Must Know	All	
3	Write pathogenesis	Cognitive	Must Know	All	
4	Write radiographic features	Cognitive	Must Know	All	
5	Write histologic features	Cognitive	Must Know	All	
6	Enumerate differential diagnosis	Cognitive	Nice to Know	All	
7	Write treatment & prognosis	Cognitive & Psychomotor	Must Know	All	

CONTENTS

- ❑ Reactive lymphoid aggregate
- ❑ Lymphoid hamartoma
- ❑ Angiolymphoid hyperplasia with eosinophilia
- ❑ Lymphoepithelial cyst



REACTIVE LYMPHOID AGGREGATE

- ✓ -Called as **Reactive lymphoid hyperplasia**.
- ✓ -Lingual tonsil located on the posterior portion of tongue on dorsolateral aspect.
- ✓ -One of the largest oral lymphoid.
- ✓ -It becomes inflamed & enlarged to become clinically evident.



- ❑ This enlargement is usually bilateral.
- ❑ If unilateral may be mistaken for early carcinoma.
- ❑ It is typically surrounded by crypt lined by stratified squamous epithelium.
- ❑ This reactive lingual tonsil called as “**foliate papillitis**”.
- ❑ It may occur in lymphoid aggregate in other locations. **Buccal mucosa** is very common.
- ❑ Clinically lymph node presents as firm, nodular submucosal mass with tenderness positive.
- ❑ Hyperplastic lymphoid polyps has also been described as **polypoid tissue**.
- ❑ It occurs on gingiva, buccal mucosa, tongue and floor of mouth.

LYMPHOID HAMARTOMA

❑ **-Other names**-Angiofollicular lymph node hyperplasia, Angiomatous lymphoid, Castleman tumor, Giant benign lymphoma, Hamartoma of lymphatics, Giant lymph node hyperplasia.

❑ **Types of castleman's disease:-**

❑ 1.Hyaline vascular type.

❑ 2.Plasma cell type.



❏ 1. Hyaline vascular type:-

Common site for occurrence:- chest, stomach & neck.

- If it so then called as localised disease of hyaline vascular type.
- Less common site for occurrence:- armpits, pelvis & pancreas.
- Growths represents abnormal enlargement of lymph nodes.
- It is generally asymptomatic.
- May develop as a non-cancerous growth of lymph node.

2. Plasma cell type-

- It is rare disorder characterised by noncancerous benign growth that may develop in lymphoid tissue throughout the body.
- May be associated with fever, weight loss, skin rash, hemolytic anemia, hypergammaglobulinemia.

3. Multicentric/generalised castleman's disease-

- Newly reported in medical literature.
- Affect multiple area of body.
- Exhibit as hepatosplenomegaly.

❏ Exact cause of Castleman's disease is not known.

❏ Some speculate that increased production of interleukin6 involves.

ANGIOLYMPHOID HYPERPLASIA WITH EOSINOPHILIA

- ❑ -Other names-Epithelioid hemangioma, Histiocytoid hemangioma, Pseudopyogenic granuloma, Papular angioplasia, Inflammatory angiomatous nodules.
- ❑ -Uncommon idiopathic condition.
- ❑ -It presents as a isolated, grouped plaques or nodules in skin of head and neck.
- ❑ -Most patient presents lesions in peri-auricular region, forehead or scalp.

- ❑ -Sometimes it is marked distinctly by proliferation of blood vessel with enlarged endothelial cells.
- ❑ -Blood vessels are accompanied with inflammatory infiltrate that include eosinophils.
- ❑ -Lesion is benign, may be persistent and difficult to eradicate.
- ❑ -It may be benign or may be an unusual reaction to varied stimuli, including trauma.
- ❑ -It shows some similarities with **Kimura disease**.

❏ Clinical features-

❏ -Uncommon but not rare.

❏ -common in Japan.

❏ -Most common in Asians followed by Caucasians.

❏ -More common in female.

❏ -In age 20-50 years with mean onset 30-33 years.

❏ -Rare in elder patients and in non indian pediatric population.

- ❑ -Although it may be benign tumour, numerous factors suggest it as an unusual reactive process.
- ❑ -Condition may be multifocal.
- ❑ -May occur following various trauma or infection.
- ❑ -Hyperestrogenemic states like pregnancy, oral contraceptive use faster lesion growth.
- ❑ -Approximately 20% of patient have blood eosinophilia.



- ❑ -It may persist for years but do not show any malignant transformation.
- ❑ -Patients presents with expanding nodules or group of nodules, usually in vicinity of the ear.
- ❑ -Appear as dome shaped, smooth surface papule or nodules.
- ❑ -It may be associated with pain or pruritis.

- ❑ -Uncommon symptoms: pulsation and spontaneous bleeding.
- ❑ -Lesions range from erythematous to brown may be eroded or crusted.
- ❑ -Most lesions are 0.5-2 cm in diameter with range of 0.2-8 cm.
- ❑ **Differential diagnosis-**
- ❑ Granuloma faciale, Insect bites, Pyogenic granuloma (lobular capillary hemangioma), Angiosarcoma, Hemangioendothelioma, Hemangioma.

❏ Histologic Features-

❏ -Characteristic histological features include:-

❏ -Proliferation of small blood vessels.

❏ -Many of which is lined by endothelial cells with uniform ovoid nuclei and intracytoplasmic vacuoles.

❏ -Endothelial cells have cobblestone appearance.

- ❑ -Perivascular & interstitial infiltrate composed primarily of lymphocytes & eosinophils.
- ❑ -Eosinophils typically comprise of 5-15%.
- ❑ -Rarely account for as 50% of infiltrate.
- ❑ -Occasionally devoid of eosinophils.
- ❑ -Lymphoid aggregates with or without follicle formation.

❏ Treatment-

- ❏ -Not mandatory.
- ❏ -Intralesional corticosteroids & irradiation have been employed.
- ❏ -But not very effective.
- ❏ -Surgical removal of lesion demonstrate best results.
- ❏ -They may recur.

LYMPHOEPITHELIAL CYST

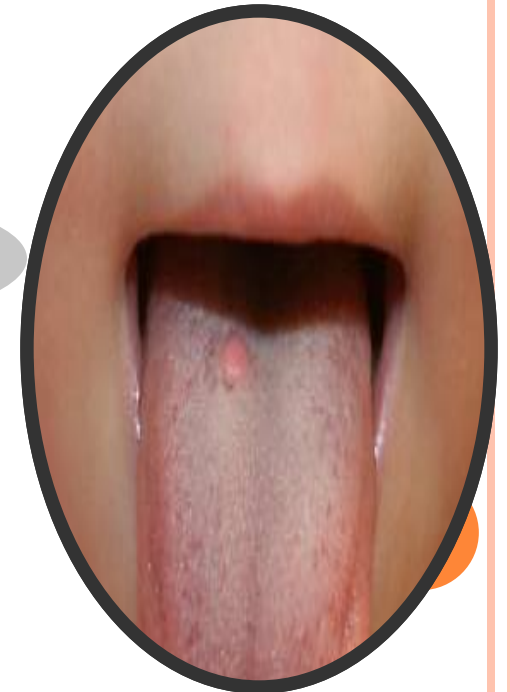
- ❑ -It develops within a benign lymphoid aggregate or accessory tonsil of oral or pharyngeal mucosa.
- ❑ It was first invented by Parmentier in 1857 as hydatid cyst.
- ❑ -Outside head & neck, this cyst is found in pancreas & testis.
- ❑ -Branchial cleft cyst, parotid cyst are similar cyst but they are much larger & parotid cyst does not contain surrounding lymphoid aggregate.

❏ Clinical features-

- ❏ -It presents as movable, painless submucosal nodule with yellow or yellow white discolouration.
- ❏ -Occasionally cyst are transparent.
- ❏ -Commonly found in oral cavity, at floor of mouth.
- ❏ -May occur on lateral, ventral tongue, soft palate, mucosa over pharyngeal tonsil.

- ❏ -Diagnose usually during teen years or third decade of life.
- ❏ -Generally diameter is less than 0.6 cm.
- ❏ -Surface of cyst indented with tonsillar crypts.
- ❏ -Superficial cyst may rupture to release foul tasting, cheesy, keratinaceous material.

- ❑ - This cyst has clinical appearance same as that of epidermoid, dermoid cyst of oral or pharyngeal mucosa.
- ❑ - But has less growth potential.
- ❑ - Never occur on alveolar mucosa hence can be easily distinguish from gingival cyst of adult or from
- ❑ - unruptured parulis or pus pockets.



❏ Histologic feature-

- ❏ -It is lined by atrophic and degenerated stratified squamous epithelium.
- ❏ -Usually lacks rete pegs.
- ❏ -Demonstrate a minimal granular cell layer.
- ❏ -Orthokeratin is seen to be sloughing from epithelial surface into cystic lumen.
- ❏ -Often sometimes fill lumen completely or sometimes shows dystrophic calcification.

- ❑ -Rarely mucus filled goblet cells seen within the superficial layers of epithelium.
- ❑ -Occasionally cyst shows epithelial lined communication with overlying mucosal surface.
- ❑ -Cyst is entrapped within well demarcated aggregate of mature lymphocytes.

- ❑ -Lymphoid aggregates may be hyperplastic.
- ❑ Combination of epithelium lined cyst with lymphoid aggregates is unique enough to make diagnosis. But it should not be confused with Warthins tumour.
- ❑ Warthin tumour is not lined by squamous epithelium but by bilayered cuboidal, columnar or oncocytic ductal epithelium.

❑ Treatment-

- ❑ -No treatment is required unless its location is such that it is constantly being traumatized.
- ❑ -Can be removed by conservative surgical excision.
- ❑ -No malignant transformation.
- ❑ -It can be involved with extranodal lymphoma

— Developmental disturbances associated with:

- Size of teeth
- Shape of teeth
- Number of teeth
- Growth of teeth



CONTENTS

- ❑ Microdontia
- ❑ Macrodontia
- ❑ Gemination
- ❑ Fusion
- ❑ Concrescence
- ❑ Dilaceration
- ❑ Talon's cusp
- ❑ Dens invaginatus
- ❑ Dens evaginatus
- ❑ Taurodontism
- ❑ Supernumerary roots
- ❑ Anodontia
- ❑ Supernumerary teeth



DEVELOPMENTAL DISORDER IN SIZE OF TEETH

❑ MICRODONTIA

❑ Smaller than normal teeth

❑ Three types

❑ True generalized microdontia
(Smaller size of teeth)

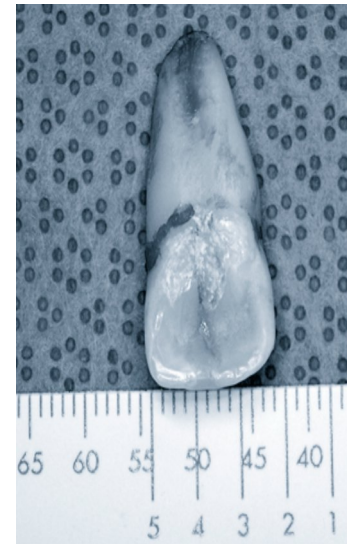
❑ Relative generalized microdontia
(Larger size of jaws)

❑ Microdontia involving single tooth
(Peg laterals, Third molars, premolars)



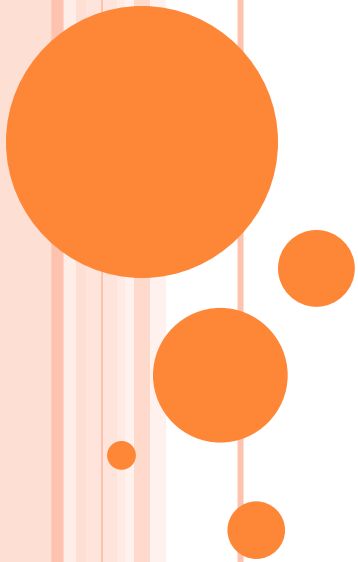
MACRODONTIA

- ❑ Larger than normal teeth
- ❑ Three types:
 - ❑ True generalized macrodontia
(Larger size of teeth)
 - ❑ Relative generalized macrodontia
(Smaller size of jaws)
 - ❑ Macrodontia involving single tooth
(Central Incisors)



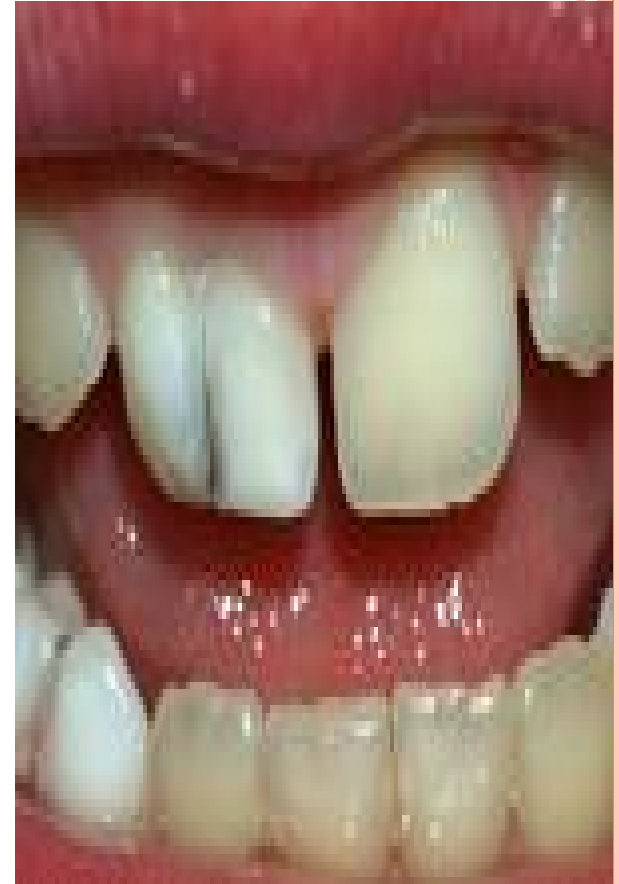
DEVELOPMENTAL DISTURBANCES

Shape of Teeth



GEMINATION (TWINNING)

- ❑ Attempt at division of single tooth germ by invagination, resulting in the formation of incomplete teeth.
- ❑ One structure with completely or incompletely separated crowns with single root and root canal
- ❑ Twinning refers to production of two teeth from single tooth bud resulting in formation of one normal and one supernumerary tooth.



FUSION

- ❑ Union of two normally separated teeth
- ❑ Complete or incomplete fusion
- ❑ Pressure or crowding of developing teeth
- ❑ Fusion may be between two normal teeth, a normal tooth and a supernumerary tooth.
- ❑ Clinical complications like, spacing, periodontal problems, esthetics.



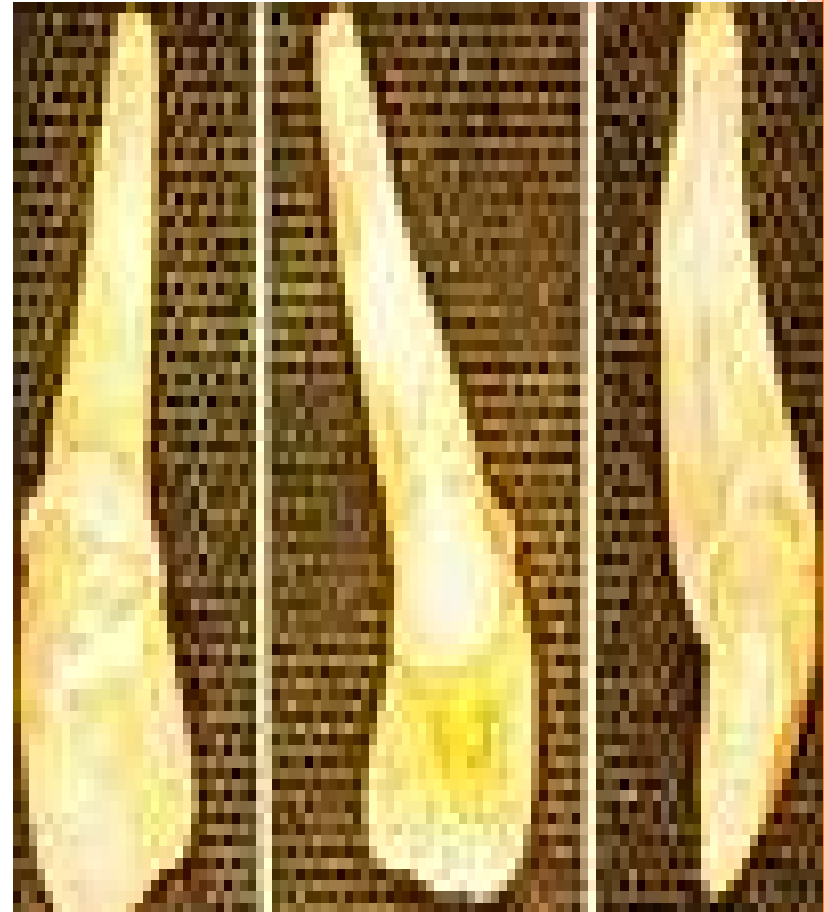
CONCRESCENCE

- ❑ Form of fusion which occurs after root formation
- ❑ Teeth united by cementum
- ❑ Due to traumatic injury, crowding leads to loss of interdental bone
- ❑ May occur before or after tooth eruption
- ❑ Difficult extraction



DILACERATION

- ❑ Refers to abnormal bend or curvature in crown or root.
- ❑ Due to trauma during tooth formation
- ❑ Can occur anywhere in the tooth
- ❑ Complications like extraction, root canal instrumentation.



TALONS CUSP

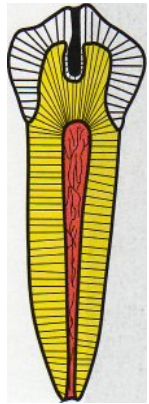
- ❑ Anomalous structure resembling an eagles talon on lingual aspect of incisors.
- ❑ Consists of normal enamel, dentin and pulp.
- ❑ Complications like dental caries, esthetics, malocclusion etc.
- ❑ Rubinstein taybi syndrome
 - ☛ Developmental retardation
 - ☛ Broad thumbs and great toes
 - ☛ Delayed or incomplete descent of testis in males
 - ☛ Bone age below fiftieth percentile
 - ☛ Talons cusp



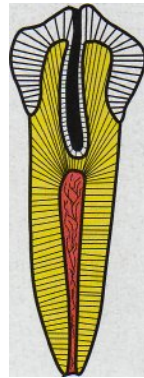
DENS IN DENTE

- ❑ Invagination in the surface of crown before calcification has occurred.
- ❑ Etiology
 - 🦷 Increased localized external pressure
 - 🦷 Focal growth retardation
 - 🦷 Focal growth stimulation in tooth bud
- ❑ Maxillary lateral incisor most commonly involved
- ❑ Radiographically
 - 🦷 Pear shaped invagination of enamel

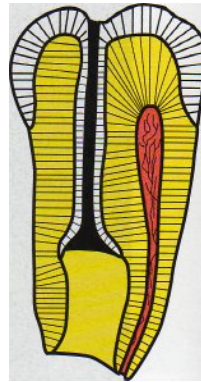




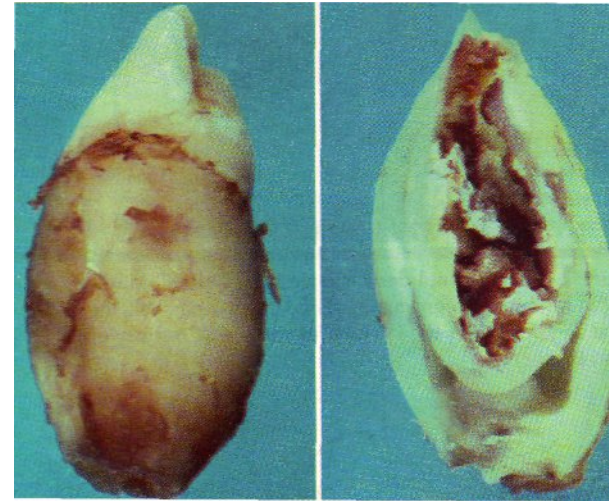
Type I



Type II



Type III



DENS EVAGINATUS

- ❑ Accessory cusp or globule of enamel on the occlusal aspect between buccal and lingual cusp of premolars
- ❑ Proliferation and evagination of IEE and odontogenic mesenchyme.
- ❑ May lead to disturbance in occlusion, displacement of teeth, pulp exposure, infection



TAURODONTISM

- ❑ Bull like teeth/ Resembling teeth of cud chewing animals
- ❑ Classified as
 - 🦷 Hypotaurodont
 - 🦷 Hypertaurodont
 - 🦷 Mesotaurodont
- ❑ Results due to failure of invagination of HERS
- ❑ Deciduous & Permanent teeth
- ❑ Molars are affected, Unilateral but can be bilateral.



❏ R/F

- 🦷 Rectangular shape
- 🦷 Large pulp chamber
- 🦷 Bifurcation near the apex



Figure 1: Apically displaced fracture and short roots in mandibular premolars



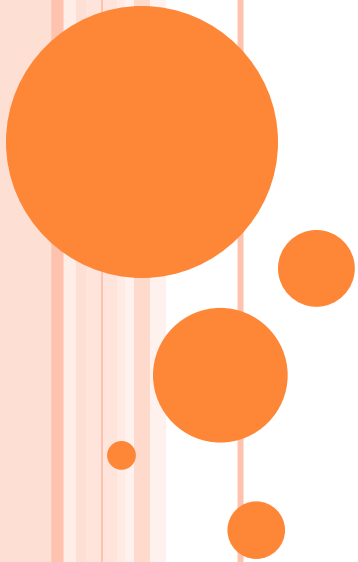
SUPERNUMERARY ROOTS

- ❑ Extra root
- ❑ Usually cuspids and bicuspid
- ❑ Molars may also have an extra root
- ❑ Problem during extraction, Root canal treatment



DEVELOPMENTAL DISTURBANCES

Number of Teeth



ANODONTIA

❑ True

- ☞ Total/ Complete
- ☞ Partial/ Incomplete

❑ False

- ☞ Total/ Complete
- ☞ Partial/ Incomplete



SUPERNUMERARY TEETH

- ▣ Incisors
- ▣ Premolars
- ▣ Paramolars
- ▣ Distomolars
- ▣ **Gardner Syndrome**
 - ☛ Multiple Osteoma
 - ☛ Multiple Supernumerary teeth
 - ☛ Compound Odontoma
 - ☛ Unerupted teeth
 - ☛ Hypercementosis





- PREDECIDUOUS DENTITION

- HORNIFIED STRUCTURE ON GINGIVA RESEMBLING TEETH

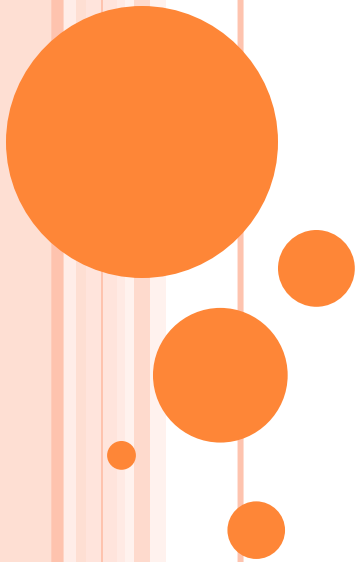
- ▣ Postpermanent dentition

- 🦷 Third set of dentition



DEVELOPMENTAL DISTURBANCES

Growth (Eruption) of Teeth



- ❑ Premature eruption
- ❑ Delayed eruption
- ❑ Eruption squelstrum
- ❑ Multiple unerupted teeth
- ❑ Embedded teeth
- ❑ Impacted teeth
- ❑ Ankylosed teeth (Submerged teeth)



Thank You!!!

