

FLORIDA SOCIETY OF ORAL & MAXILLOFACIAL SURGEONS 2018 ANNUAL MEETING

EARLY ORAL CANCERS AND PRE-CANCERS & POTPOURRI OF ORAL PATHOLOGY

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Conflicts of Interests

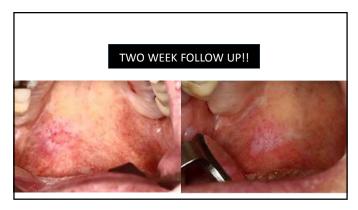
 Neither my immediate family nor I have any financial interests that would create a conflict of interest or restrict our independent judgment with regard to the content of this course.

Course Objectives

- Upon completion of this course, participants should be able to:
 - Recognize and formulate a differential diagnosis, understand the etiology and management of various oral and maxillofacial conditions.
 - Better recognize early mailignacies, improve diagnostic skills for oral soft and hard tissue lesions through practice sessions utilizing the audience response devices.







IDIOPATHIC LEUKOPLAKIA

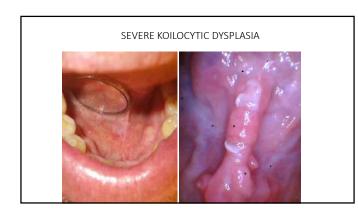
- FEATURES TO WORRY ABOUT
 - Occurrence in non-smoker
 - Thickened often corrugated appearance
 - Associated erythema
 - High risk location-horseshoe shaped area
 - ??pain
 - Multifocal or recurrent

NON-SMOKERS LEUKOPLAKIA

- 5-8 times INCREASED risk of oral cancer
- More frequent on tongue/floor of mouth(64 vs. 11%)
- More dysplasia(38 vs. 5 %)
- Younger patients
- Often very subtle lesions under tongue and on lingual frenum
- Likely high risk HPV related

TONSILLAR CRYPT EPITHELIUM

- Stratified squamous but basaloid so virus can invade epithelium <u>w/o surface ulceration</u>
- Well known localization for the replication of viruses
- Tonsillar crypt epithelium serves as reservoir for Epstein Barr virus and also HPV
- Majority of OPSCCA originate from tonsillar epithelium





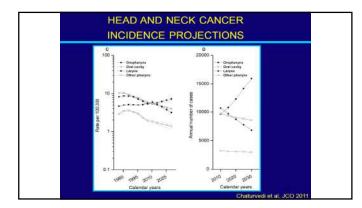


WHY TALK TO DENTISTS ABOUT HPV???

- Dentists may be next group of providers to participate in prevention of HPV and OPCancer
- Improve HPV knowledge and dentists communication skills with patients
- Highlight barriers to discuss HPV with patients
- Put HPV vaccination guestion on dentists patient health history
 - 47% DDS no discuss HPV with patients
 - 33% discuss with some patients(papilloma)
 - 19% Discuss with all patients
 - JADA 149(1) Jan 2018 pp 9-17

HPV & ORAL CANCER

- 70% of sexually active adults are HPV positive
- Prevalence of oral HPV in US population 7%
- Oro-pharyngeal cancer most common H & N cancer shortly

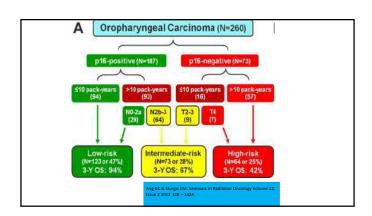


HPV & ORAL CANCER

- 10% male, 3.6% female
- Peak incidence: 30-34 y/o -7.37% 60-64 y/o -11.45%!!!
- HPV-16 incidence 1% (2.13 million people)
- But only 15,000 cases of oropharyngeal cancer/yr

HPV & ORAL CANCER

- Oropharyngeal cancers occur in younger age group.
- Especially troubling increase in non-smoking males with oropharyngeal ca
- Men 45 -60 2-3 x more likely to get oro-pharyngeal cancer
- Fortunately HPV associated cancers have a better prognosis
- HPV+HNSCC: Less chromosomal mutations (compared to smoking/ drinking associated tumors)









TESTING FOR HPV-TISSUE SAMPLES

- HPV-+ cases express high levels p16 tumor-suppressor protein
- Diffuse nuclear and cytoplasmic p16 protein-staining correlates strongly with presence of HPV by in-situ hybridization, and PCR.

IDIOPATHIC LEUKOPLAKIA

- FEATURES TO WORRY ABOUT
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VERRUCO-PAPILLARY HYPERKERATOSIS





(ERYTHRO-) Leukoplakia

- 82% of transformed leukoplakias
- 4 times risk of oral cancer (23.4%)

ERYTHROPLAKIA

ERYTHROPLAKIA

HISTOLOGIC SPECTRUM:

Benign keratosis – 0%
Mild dysplasia - 10%
Severe dysplasia or ca-in-situ - 40%
Invasive carcinoma - 50%

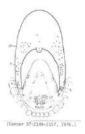
ERYTHROPLAKIA

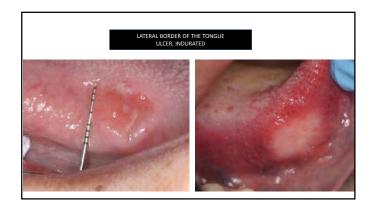
- Red lesions should be viewed with suspicion, biopsied.
- Suspicious red lesion may be observed for 10-14 days, if persists biopsy
- Recurrence and multifocal involvement are common
- Long-term follow-up is needed

HIGH RISK LOCATIONS ORAL CANCER

- 90% LOCATED IN:
 - LATERAL OR VENTRAL TONGUE
 - FLOOR OF THE MOUTH
 - LINGUAL FRENUM
 - •SOFT PALATE ANTERIOR PILLAR COMPLEX
 - FREE AND MARGINAL GINGIVA- ring around the collar

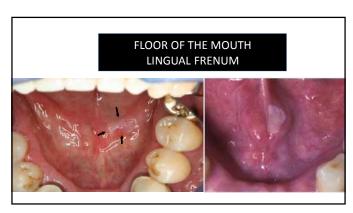
HIGH RISK LOCATIONS

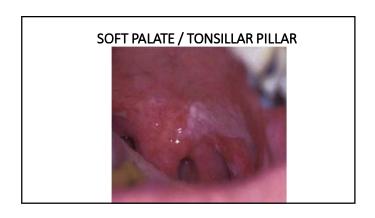














Proliferative Verrucous Leukoplakia (PVL)

- Middle aged females(4:1F>M)
- Mean age 63.9(over 62)
- Little relation to smoking
- Little known of etiopathogenesis

$TABLE\ 1: \ {\it Diagnostic Criteria for Proliferative Verrucous Leukoplakia.}^{19}$

The diagnostic criteria are met if an individual has three major criteria (Including E) or two major criteria (Including E) and two minor criteria

Major Criteria:

- A. A white lesion involving two different oral sites, most frequently found in the gingiva, alveolar process and palate

 B. Presence of a verrucous area

- C. The lesions are evolving and have spread
 D. Presence of recurrence in a previously treated area
 E. Histologically, the tissue sample may display hyperkeratosis, verrucous hyperplasia, verrucous carcinoma or squamous cell carcinoma

Minor Criteria:

- Million Criteria.

 A. A white lesion that measures at least 3 cm when including all the affected sites in the oral cavity

 B. Female

 C. Nonsmoker

- C. Nonsmoker D. Disease progression longer than five years

Proliferative Verrucous Leukoplakia (PVL)

- Recurrent(av.71%)/persistent
- Progresses to multiple sites
- ??HPV 16,18 positive (0-80%)
- High (40-100% av.64%) risk for transformation
- Time to transformation 4.7-11.6 years mean 6 years
- Verrucous carcinoma or squamous
- 39% of 277 patients died of disease within 7 year F/U

Otolaryngol Head Neck Surg 2015 Oct;153(4):504-11

PROLIFERATIVE VERRUCOUS CARCINOMA









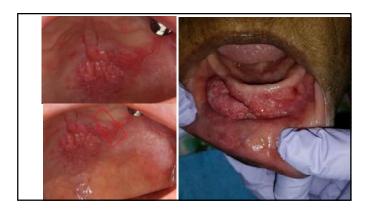


Proliferative Verrucous Leukoplakia (PVL)

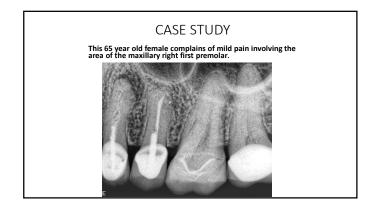
- No specific treatment modality has proven effective
- Laser ablation rapid & high rate of recurrence
- 2 not so recent papers no association with HPV
- But surgery alone 18/25 recurrences within 6 months
- Surgery plus antiviral (anti-HPV immunomodulatory agent) isoprinosine or methisoprinol--2/25
- 18 months post op 2 additional recurrences in anti viral group(4/25) none in surgery group(18/25)

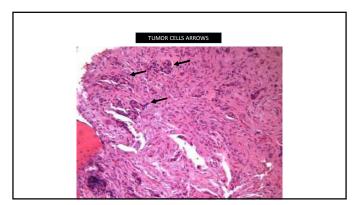


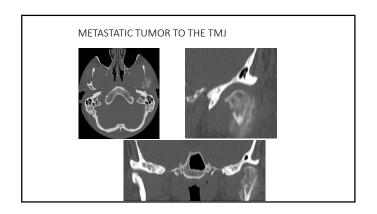


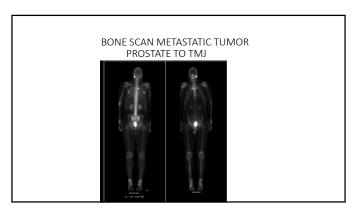












METASTATIC TUMORS TO THE JAWS-SYMPTOMS

- Pain
- Swelling
- Loosening of teeth
- Presence of a mass
- Paresthesia

METASTATIC TUMORS TO THE JAWS

| • | Breast | 23% | |
|---|--------------|-----|-----|
| • | Lung | | 15% |
| • | GI | | 8% |
| • | Male Repro | | 7% |
| • | Female Repro | | 3% |
| • | Renal | | 3% |
| • | Thyroid | | 3% |
| | | | |

METASTATIC TUMORS TO THE JAWS

- Usually present as radiolucent defects.
- Defect well circumscribed or ill defined ("moth-eaten" appearance).
- Some carcinomas (prostate and breast), osteoblastic resulting in radiopaque or mixed radiolucent and radiopaque lesions.

CASE STUDY

- a. One of the 3 P's
- b. Gingival manifestation of a systemic disorder
- c. Peripheral odontogenic neoplasm
- d. Malignancy





GRANULOMATOSIS WITH POLYANGIITIS

(Formerly Wegener Granulon

- Unknown cause
- Wide age range
- No gender predilection
- 90% of cases in Caucasians
- Can involve almost every organ system in the body
- Early stage (before renal involvement) = Strawberry gingivitis and enlargement of one or more major salivary glands
- Late stage (after renal involvement) = Oral ulcerations
- Diagnosis
 - PR3-ANCA (previously c-ANCA): Seen in 90-95% of generalized Wegener granulomatosis and 60% of early or localized cases
 - ELISA test for antibodies against PR3
- Mean survival for untreated patients with disseminated disease = 5 months
- 80% of patients die within 1 year and 90% die within 2 years
- Treatment: Oral prednisone and cyclophosphamide
- With appropriate therapy, 75% of patients have prolonged remission

3 FLAVORS OF WEGENER'S GRANULOMATOSIS

- Generalized Wegener's Granulomatosis
 - Initial upper & lower respiratory tract and rapid renal involvement

Limited Wegener's Granulomatosis

Respiratory tract w/o rapid kidney involvement

Superficial Wegener's Granulomatosis

Lesions primarily of skin and mucosa Systemic involvement develops slowly

HYPERTROPHIC LICHEN PLANUS???

PRETEST LP OR CANCER? 52 y/o female with irregular white and red lesion on right buccal mucosa for 3 years 5/10/2011



HISTOLOGIC OVERLAP

• 29% OF DYSPALSTIC AND MALIGNANT ORAL LESIONS HAD 3 OR MORE OF THE FIVE HISTOLOGIC FEATURES OF LICHEN PLANUS

PRETEST LP OR CANCER?

I have a patient I have been seeing for some time. Originally presented with well localized erythematous lesion of palate. A biopsy prior to my evaluation Dx: lichen planus. Did another biopsy 2009 and immunoflourescence confirmed diagnosis. Treated her with clobetasol ointment and diflucan to no avail. I'm not sure I'm actually dealing with I'm of the mill' lichen planus and would like you to take a look. I've attached a few pictures from September.



Epidemiology Lichen planus

- Common mucocutaneous disorder- 2% of population.
- Increasing frequency
- Disease of middle aged and elderly.
- More common in females.

Oral Lichen Planus has distinctive clinical features and classical distribution... or does it??



LICHEN PLANUS

- 90% of patients have bilateral reticular lesions in the posterior buccal mucosa.
- The tongue is the next most commonly involved followed by the gingiva and alveolar ridge.

Reticular Lichen Planus



EROSIVE LICHEN PLANUS



Factors in the Etiopathogenesis of LichenOID MUCOSITIS

- Drug induced
- Foreign body (gingiva) prophy jet
- Dental materials

TRIGGERS of LichenOID MUCOSITIS/Lichen Planus

- ? Stress
- Trauma
- Aspirin(actually NSAID's /Amalgam
- Yeast
- Idiopathic (true lichen planus)

<u>S</u>TAY

- Reduce stress
- Exercise 5x week for 30 minutes
- Must sweat to be effective

TRAUMA

• Triggers the condition in susceptible individuals

KOEBNER PHENOMENON

 Development of isomorphic pathologic lesions in the traumatized uninvolved skin of patients who have cutaneous disease

TRAUMATIC LICHENOID MUCOSITIS





Α

- Aspirin (actual NSAIDS)
- Amalgam

Drugs Implicated in Oral Lichenoid Reactions Category Drugs

| Antimicrobials | Dapsone Ketoconazole Para-aminosalicylic acid Sodium aminosalicylate Streptomyrin Sparloacin Suffamethoazole, Sulfasalazine Tetracycline |
|-------------------|---|
| Antiparasitics | Antimony compounds (tibophen, tibocaptate) Organic arsenicals Chloroquine Pyrimethamine Quinacrine ACE inhibitors -Captopril Chlorothiazide, Hydrochlorothiazide |
| Antihypertensives | Labetaid, Praction, Propanoiol Lasix Mercurial diuretics Methyldopa |

Drugs Implicated in Oral Lichenoid Reactions Category Drugs – Cont.

| Antiarthritics | Aurothioglucose Colloidal gold (Europe only) Gold sodium thiomalate and thiosulfate |
|--|---|
| Anxiolytics | Fenclofenac Ibuprofen |
| Non-Steroidal Anti-inflammatory Agents | Naproxen Phenylbutazone |
| | Chlorpropamide Tolazamide Tolbutamide |
| Hypoglycemic agents | Allopurinol |
| Uricosurics | |

Drugs Implicated in Oral Lichenoid Reactions Category Drugs —

Cont.

| Amitripyline | Copidingel (Plavis) | C

Drug Induced

- Often erosive
- May involve the lips
- Often asymmetric

LICHENOID DRUG ERUPTIONS



Lichenoid Drug Eruption

- Takes an average 12 months to develop.
- Can develop after 10 years on same drug
- Ebbs and flows while on drug
- May take up to 24-months to clear.
- Usually improves in 2-8 weeks..

YEAST

- Red, fuzzy lesion
- Exacerbate the condition





Treatment

- Not entirely satisfactory
- No cure available
- Immune suppression is the key
- Only 1 out of 15 resolve spontaneously.

Topical steroids are the mainstay of therapy.

Lichen Planus Treatment

- Clobetasol gel 0.05%
- Dsp. 15 or 30 gm.
- Sig. apply sparingly <u>b.i.d.</u>
- Very strong use for Tx initiation

TREATMENT 2 WEEKS TOPICAL CLOBETASOL • BEFORE • AFTER



Lichen Planus Treatment

- Lidex gel 0.05%
- Dsp. 15 or 30 gm.
- Sig. Apply sparingly 3-4 times a day.
- Good for long term treatment

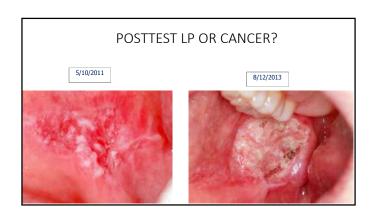
Lichen Planus Treatment

 Stringent oral hygiene and prophylaxis procedures Q 3-6 months.



Follow-up

- Patients must be seen 1-2 times a year.
- Return if condition worsens.
- Erosive / atrophic types much more risky.
- Biopsy IF lesions change clinical appearance or resist Tx.



POSTTEST LP OR CANCER?

I have a patient I have been seeing for some time. Originally presented with well localized erythematous lesion of palate. A biopsy prior to my evaluation Dx: lichen planus. Did another biopsy March of 2009 and immunoflourescence confirmed diagnosis. Treated her with clobetasol ointment and diffucan to no avail. I'm not sure I'm actually dealing with 'run of the mill' lichen planus and would like you to take a look. I've attached a few pictures from September.





EB Virus Positive B-cell Lymphoproliferative Disorder (E-LPD)

- E-LPD) spectrum of lymphoid expansion entities caused by immunosuppression or immunosenescence and associated with Epstein-Barr virus.
- Immunosenescence seen in 65+ y/o
- Immunosuppression can be post transplant or with any use of immunosuppressive drugs
- First sign is lymphadenopathy then destructive lesions(recalcitrant oral ulcers), ranging up to florid neoplasia with aggressive behavior

POST TRANSPLANT LYMPHOPROLIFERATIVE DISEASE

- Can be self limited with decreased immunosuppression or go on to be full blown monoclonal lymphoma.
- Therefore early diagnosis is critical
- 1/3 involve the H & N
- Oral cavity is often the first place lesions appear.

POST TRANSPLANT LYMPHOPROLIFERATIVE DISEASE

- E-LPD second most common form of neoplasia in solid organ transplant patients
- Important early cause of cancer related death and graft loss.
- Distinct clinicopathologic entity first described in the oral cavity as recalcitrant destructive EBV+ oral ulcers in 2010

CASE STUDY

70y/o male painful gingival ulcer. Present 3 months and is not healing despite root planning and excision. History of kidney transplant in 2104

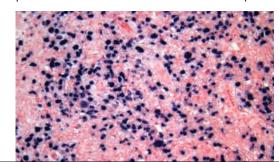




PTLD

- 1/3 of cases in H & N
- Second most common form of neoplasia in solid organ transplant patients
- Early cause of cancer related death and graft loss
- First sign lymphadenopathy then destructive lesions

Epstein-Barr virus encoded small RNA (EBER)



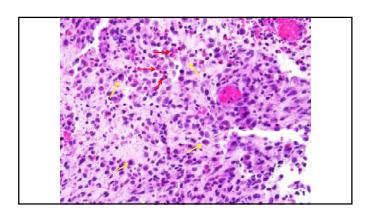
CASE STUDY

20's y/o male C.C. pain and drainage lower jaw. Seen in several emergency departments for jaw swelling and drainage. Treated 10 years ago with both chemo and radiation therapy for bilateral cancer of his hips









LANGERHANS CELL DISEASE (Histiocytosis X; Eosinophilic Granuloma

- Histiocytosis X: spectrum of clinicopathologic disorders characterized by proliferation of histiocyte-like cells and eosinophils.
- Now considered clonal neoplastic process involving myeloid dendritic histiocytic cells
- This disorder is characterized by single or multiple osteolytic bone lesions

CLINICO-PATHOLOGIC SPECTRUM

- Eosinophilic granuloma of bone- one or more lesions w/o visceral involvement
- Hand-Schüller-Christian disease- chronic disseminated bone, skin, viscera
- Letterer-Siwe disease- acute disseminated with prominent cutaneous, visceral, and bone involvement, in infants

LANGERHANS CELL DISEASE

- Wide age range with patients being reported even in the 6th and 7th decades.
- 50%+ seen under age 10.
- 55% single system disorder with bone lesions solitary or multiple, most common clinical presentation.

LANGERHANS CELL DISEASE

- Skull, ribs, vertebrae, mandible most frequent sites.
- Jaws affected in 30% skull 21% of adults.
- Most common lesions in oral cavity are teeth floating in air, gingivitis/periodontitis, unexplained mass or mucosal ulcers

RESEMBLES AGGRESSIVE PERIODONTAL DISEASE

LANGERHANS CELL DISEASE

- Clinical course quite varied depending on age and stage of the disease
- Pts with multi-focal disease respond initially to treatment but often recur same or distant site
- The bilateral cancer in this patient's hips was actually LCH which has recurred 10 years after initial remission

TREATMENT LANGERHANS CELL DISEASE

- Single agent treatment such as prednisone intralesional and or systemic, prednisone combined with vinblastine and/ or curettage of readily accessible bone lesions.
- Low dose radiation for less accessible lesions.





MY ASSESSMENT

- Judging from wear pattern in teeth something opened up vertical.
- Assume condyles are normal and her thyroid function.
- I think acquired macroglossia from vascular malformation/hemorrhagic diathesis, amyloid accumulation or clotting disorder
- If tall evaluate for Beckwith-Weidman syndrome.
- Other signs of bruising or clotting problems on the skin??

AMYLOIDOSIS

- Heterogeneous group of conditions characterized by deposition of extracellular material called amyloid.
- Amyloid associated with: multiple myeloma, rheumatoid arthritis or chronic infections including tuberculosis.

SYSTEMIC AMYLOIDOSIS

- Occurs in several forms:
 - primary
 - myeloma-associated
 - secondary
 - hemodialysis-associated
 - heredofamilial

Primary & Myeloma-Associated Amyloidosis

- Affect older adults (av. 65)
- Mucocutaneous lesions and macroglossia from amyloid deposits
- **Macroglossia** 12-40%, diffuse or nodular enlargement of tongue.

MYELOMA TREATMENT

- Multiple drugs
 - Dexamethasone
 - I.V. Bisphosphonates
 - Proteases inhibitors(Velcade)
 - Monoclonal antibodies(attack CD-38)
 - Melphalan-nitrogen mustard alkylating agent
 - Immune modulating drugs
 - Lenalidomide(revlimid)

TREATMENT SIDE EFFECTS

- 81% infections
- Blood clots
- Thrombocytopenia
- MRONJ
- Low blood counts
- Mouth sores

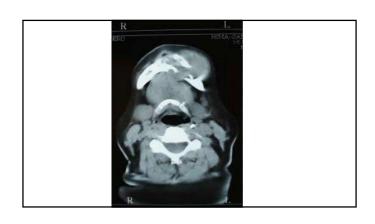
Patient with history of myeloma now with soft tissue mass











CASE STUDY

- 57 y/o Cauc male 5 cm x 3 cm spongy mass hard palate. Lesion first noted by hygienist.
- Patient reluctant to have biopsy because asymptomatic and had multiple similar "lipomas" on his back.
- Lesion did not blanch with pressure. Swelling did not communicate with teeth

HARD PALATE

DIAGNOSIS: MANTLE CELL LYMPHOMA

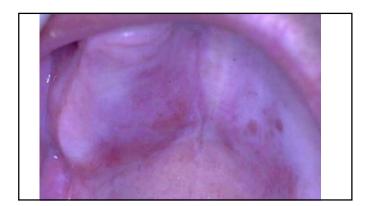
- One of four types of small B-cell lymphomas
- Distinct subtype due to short survival and aggressive course
- Predilection for elderly males, hard palate
- Treatment: chemotherapy(RCHOP), radiation and possible BMT

NON-HODGKIN'S LYMPHOMA

- Occurs primarily in adults
- Makes up 5% of oral malignancies
- Second most common malignancy of oral cavity
- Nontender slowly enlarging mass involves cervical, axillary or inguinal nodes

ORAL NON-HODGKIN'S LYMPHOMA

- Nontender, diffuse soft tissue swellings of buccal vestibule, gingiva, or posterior hard palate.
- \bullet Appear $\underline{\text{\it erythematous or purplish}}$ and have "boggy" consistency.



CASE HISTORY

This 30's female lump in palate for 2 yrs. Asked her DDS about it. He referred her
to ENT specialist who diagnosed normal anatomy. Developed headaches and
given therapy for TMD. Recently returned to her dentist who noticed the lump
and referred her to me for a second opinion.





POLYMORPHOUS ADENOCARCINOMA

- Recently recognized -1983
- Almost exclusively in the minor SG
- 60% hard or soft palate,16% BM, 12% upper lip
- 2/3rds in females

MINOR SALIVARY GLAND TUMORS

- Almost 50% malignant
- "Smaller the gland, the greater the chances of malignancy"
- Palate most frequent site (42 to 54%)
- Posterior lateral hard or soft palate
- Lips- second most common

POLYMORPHOUS ADENOCARCINOMA

- Tumor cells have uniform appearance
- Different growth patterns hence, "polymorphous"
- Perineural invasion common
- Wide surgical excision
- Overall prognosis relatively good, 80% cure rate

CASE STUDY

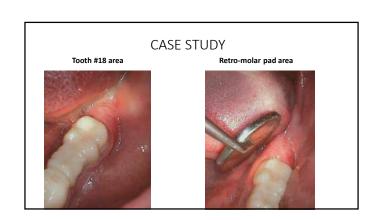
33 y/o male pain in area of tooth #18 and paresthesia lower lip 4 weeks duration. Root canal therapy and antibiotic did not relieve pain/paresthesia. Referred to a periodontist who did RP & S. Then referred to Dr Cohen for consult. Dr Cohen recommended oral surgeon do a biopsy. Diagnosis???







CASE STUDY CECT obtained Pt. placed on Augmentin 875mg for 10 days 11/13/14 Referral to Oral Surgeon



LOW GRADE CHONDROBLASTIC OSTEOSARCOMA

- Important early radiographic change symmetric widening of periodontal ligament space.
- When combined with pain/discomfort and other radiographic changes important in early diagnosis
- Lip paresthesia is key diagnostic feature
- See paresthesia only with malignancy and osteomyelitis
- Radical surgery preceded by chemo and radiation most common Tx.

OSTEOSARCOMA (Osteogenic Sarcoma)

- Most common primary malignant tumor of bone.
- Distal femur and proximal tibia are most frequent sites
- Only 7% occur in jaws.
- Mean age for jaw 33, (10-15 years older than for long bones)

OSTEOSARCOMA

- Swelling(100%) and pain(5%) most common symptoms.
- Peripheral border ill defined and indistinct.
- Spiking resorption of roots of teeth
- "Classic" sunburst or sun-ray 25% of jaw osteosarcomas

OSTEOSARCOMA

- Local uncontrolled disease cause of death>> than distant metastases.
- Jaw osteosarcomas lesser tendency to metastasize
- Metastases most often involve lungs and brain.
- 80% survival rate for jaw lesions with radical surgery alone(3cm margins).
- Long bone lesions chemo, radiation and surgery

CASE STUDY

Middle aged female 3-4 week history of swollen gums. DDS sent her to 2 MDs who Rx antibiotics w/o improvement. She is very tired all the time. DDS sent her to a periodontist who does biopsy





ACUTE MYELOGENOUS LEUKEMIA

- Begins with malignant transformation of stem cell, proliferates in bone marrow and eventually overflows into peripheral blood
- Leukemic cells "crowd out" normal defense cells and erythrocyte precursors.

LEUKEMIA

- Classified according to their histogenesis and clinical behavior
- Myeloid(granulocyte/monocytes)or lymphocytic (histogenetic origin)
- Broad category would be acute or chronic (clinical course)

LEUKEMIA

- Acute leukemias- untreated, aggressive course and result in death within a few months (survival is < six months).
- Chronic leukemias more indolent course, but end result is same (chronic - survival over one year).

LEUKEMIA

- Certain types of leukemia show specific chromosomal abnormalities
- Chronic myeloid leukemia (CML)- genetic alteration, Philadelphia chromosome, translocation of chromosomal material between long arms of chromosomes 22 and 9.

LEUKEMIA

- Environmental agents increased risk- benzene, ionizing radiation & HTLV-1
- Myeloid leukemias affect adults
- Acute myeloid leukemia (AML) broader age range including children.
- Chronic myeloid leukemia (CML) third and fourth decades.

LEUKEMIA

- Chronic lymphocytic leukemia (CLL) most common type in adults, affects elderly adults median age 60.
 - 24% of all leukemias
- Acute lymphoblastic leukemia (ALL) occurs in children(2-3), most common childhood malignancy.

LEUKEMIA

- Acute leukemia-abrupt onset
- Clinical signs and symptoms- accumulation of neoplastic cells in marrow causes marked reduction in normal white and red cells
- Causes infections and anemia respectively

LEUKEMIA

- Complain of easy bruising and bleeding.
- Petechial hemorrhages posterior hard and soft palate due to thrombocytopenia

LEUKEMIA

- Spontaneous gingival hemorrhage, result of decreased platelet counts,<10-20,000mm3 (thrombocytopenia).
- Fever associated with infection may be initial sign



LEUKEMIA

- Oral ulcers often present- due to impaired ability to combat normal flora.
- Gingival tissues most severely affected because of abundant plaque around teeth.

LEUKEMIA

- Leukemic cells infiltrate soft tissues- most frequent with myelo-monocytic types
- Results in diffuse gingival enlargement.



LEUKEMIA

- Diagnosis is established by finding poorly differentiated leukemic cells in peripheral blood and bone marrow.
- Treatment consists of various forms of chemotherapy.

LEUKEMIA PROGNOSIS

- Depends on type and age of patient.
- In children with **ALL** 95% complete remission, 75-85%% cured
- Adults similar 80% response rate but much lower cure rate(35-40%)

CASE STUDY 26 y/o female with sore throat and swollen glands





PRIMARY HERPES

- In adults can present with primarily pharyngeal involvement
- Clusters of vesicles form and coalesce into irregular shallow ulcers, Herpes=creep
- Coated tongue is constant for all primary HSV patients



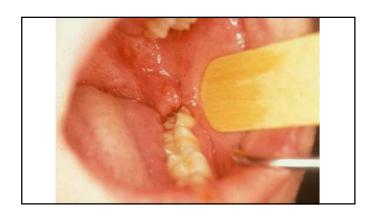


PRIMARY HERPES

- Vesicles occur on all mucosal surfaces and sometimes skin
- Always (almost) ulcerates marginal gingiva
- Fever, lymphadenopathy & swallowing difficulties

PRIMARY HERPES

- Usually affects young children
- \bullet Lesions heal completely if not traumatized
- Initial lesions prefer inflamed tissues i.e. erupting thirds and palatal of max. centrals
- Often cause gingival hypertrophy (opposite of ANUG) and lip ulcers

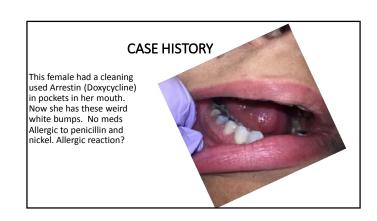


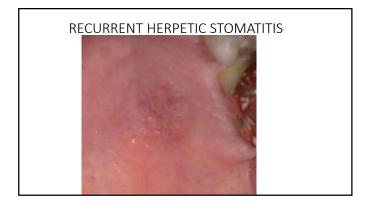
TREATMENT PRIMARY HERPES CHILDREN

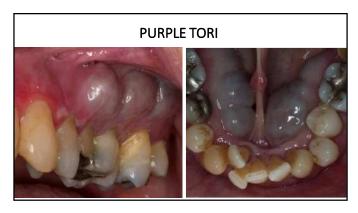
- Acyclovir elixir 200mg/5cc (banana flavored)
- 1tsp rinse 30 seconds and swallow 5x/day for 7 days

TREATMENT PRIMARY HERPES ADULTS

- Valcyclovir 1 Gram BID for 10 days
- Significantly reduces incidence of recurrences







Enrolling as Ordering/Referring Providers in Medicare Part B

Dentists.

(a) historically, have not participated in Medicare from inception,

(b) and have been falsely led to believe that by not participating in Medicare, they have Opted-Out,

(c) also, dentists have been incorrectly informed by the ADA that the Part D exclusion in October 2017 amendment of ACA for prescribing would include ordering and referring.

Affordable Care Act, Section 6405 (March 2010)

However, the enactment of the Affordable Care Act, Section 6405 (March 2010) stated that, "Physicians Who Order Items or Services are required to be Medicare Enrolled Physicians or Eligible Professionals," requires physicians or other eligible professionals to be enrolled in the Medicare Program (CMS-4159) to order or refer items or services for Medicare beneficiaries.

Only physicians and certain types of non-physician practitioners are eligible to order or refer items or services for Medicare beneficiaries.

- (1)Physicians-
- (a) doctor of medicine or osteopathy,
- (b) doctor of dental medicine,
- (c) doctor of dental surgery,
- (d) doctor of podiatric medicine
- (2)Non-physicians practitioners
- (a) doctor of optometry- (optometrists may only order and refer DMEPOS products/services
- (b) laboratory tests
- (c) X-Ray services

CMS will be fully implementing CMS-4159 (effective date of January 2019)

Over the last 4 years several implementation dates for CMS-4159 have been changed- but now has a final date of January 2019

CMS-4159- makes it mandatory for all dentists to enroll as ordering/referring providers by validating with their personal NPI.

(note: the NPI of an office practice cannot be used to validate as an ordering and referring provider)

CMS emphasizes that Medicare will only reimburse

- (a) for specific items or services when those items or services are ordered or referred by providers authorized by Medicare statute and regulation
- (b) claims that a billing provider submits in which the ordering/referring provider is not authorized by statute and regulation will be denied as a non-covered service.
- (c) The denial will be based on the fact that neither statute nor regulation allows coverage of certain services when ordered or referred by the identified supplier or provider specialty

CMS, further emphasizes that

- (a) the Medicare beneficiary **cannot be Balanced Billed** (ie..patient cannot be billed for any balances) when the claim is denied,
- (b) the patient cannot sign an ABN(Advance Beneficiary Notice) in anticipation of the denied claim in order to recover the liability,
- (c) Dentists who enroll as ordering and referring providers or have opted out will not be participating in Medicare-(they will not be able to bill claims on behalf on their Medicare patients)

Loss of Revenue to Pathology

In the last 4 years several dentists who refer biopsies have not validated or have chosen not to validate. This has resulted in

- (a) gross revenue losses in excess of \$600,000 in denied Medicare claims for biopsies
- (b) Combined with an additional \$150,000 in lost revenue from secondary insurances and copays on these denied biopsies

Three ways to Enroll in Medicare

- (1) Medicare Participating Provider- Dentists (usually an oral surgeon) enrolls and participates fully by filing medical claims on behalf of Medicare beneficiaries,
- (2) Ordering and Referring Provider- this enrollment is for dentists who refer and order services such as biopsies on behalf of their Medicare patients-(these dentists do not file medical claims on behalf of their patients-since patients pay for services out of pocket)
- (3) Opted-Out Provider- these are providers who are exempted from Medicare by submitting a signed affidavit the Medicare Administrative Contractor (MAC) for Florida

How to Enroll as Participating Medicare Provider

There are two ways to become a participating provider with Medicare.
(1)Enroll online through the PECOS portalhttps://nppes.cms.hhs.gov/IAWeb/worning.do?fwdurl=/

(2)Fill out a **CMS-855i** form and send it to First Coast Service Options Inc, who is the Medicare Administrative Contractor (MAC) for Florida.

Address: First Coast Service Options INC
Medicare Provider Enrollment
P.O. Box 44021
Jacksonville, FL 32231

How to Enroll as an Ordering and Referring Provider

There are two ways to become an **Ordering and Referring provider with Medicare**. (1)Enroll online through the **Part D Prescribers Eligible for Easy Enrollment**https://data.cms.gov/8550EasyEnroll

(2)Fill out a **CMS-8550** form and send it to First Coast Service Options Inc, who is the Medicare Administrative Contractor (MAC) for Florida.

Address: First Coast Service Options INC
Medicare Provider Enrollment
P.O. Box 44021
Jacksonville, FL 32231

How to Opt-Out of Medicare

When Opting Out of Medicare :

(1) Complete and submit a Medicare Opt-Out Affidavit to:

Medicare Provider Enrollment

P.O. Box 3409

Mechanicsburg, PA 17055-1849

(2) Have a **Private Contract** between Provider and Medicare Part B beneficiary **signed and attached to each patient new file** before medical services are performed

(3) Give each Medicare beneficiary a signed copy of the Private Contract

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