Craniofacial Pain: A Case-Based Approach to Navigating Common Pitfalls

CareQuest Institute Continuing Education Webinar

January 11, 2024



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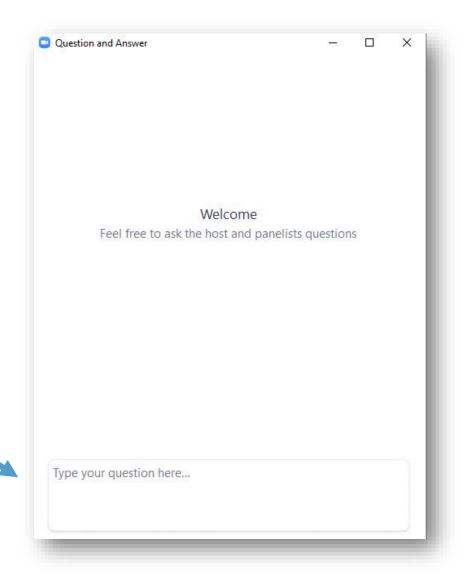
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Question & Answer Logistics

- Feel free to enter your questions into the Question & Answer box throughout the presentations.
- We will turn to your questions and comments toward the end of the hour.





Learning Objectives

At the end of this webinar, you'll be able to:

- Recognize the characteristics of temporomandibular disorders (TMD) and trigeminal neuralgia, including their symptoms, diagnostic criteria, and impact on patient quality of life.
- Recognize strategies for managing craniofacial pain, emphasizing the importance of a tailored, least invasive approach for each patient.
- Identify strategies to avoid common pitfalls in diagnosing and treating TMD and trigeminal neuralgia.



Craniofacial Pain: A Case-Based Approach to Navigating Common Pitfalls



WEBINAR | Thursday, January 11, 2024 | 7-8 p.m. ET | ADA CERP Credits: 1





Orofacial Pains

Sujay A. Mehta, DMD, MPH

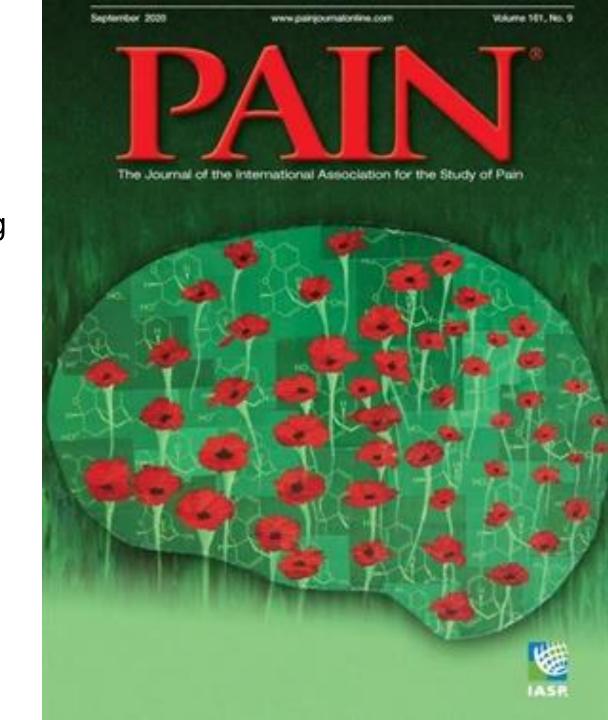
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Pain

- An unpleasant sensory and emotional experience associated with, or resembling that associated with, actual or potential tissue damage.
- A personal experience not strictly from sensory neurons.
- People learn pain concepts.
- Respect people's experience.
- Adverse effects on function.
- Non-verbal expression of pain.





Orofacial Pain

 The specialty of dentistry encompassing the diagnosis, management, and treatment of pain disorders to the jaw, mouth, face, head, and neck.

OFP recognized as a specialty on March 31, 2020.



What is the Scope of Orofacial Pain?

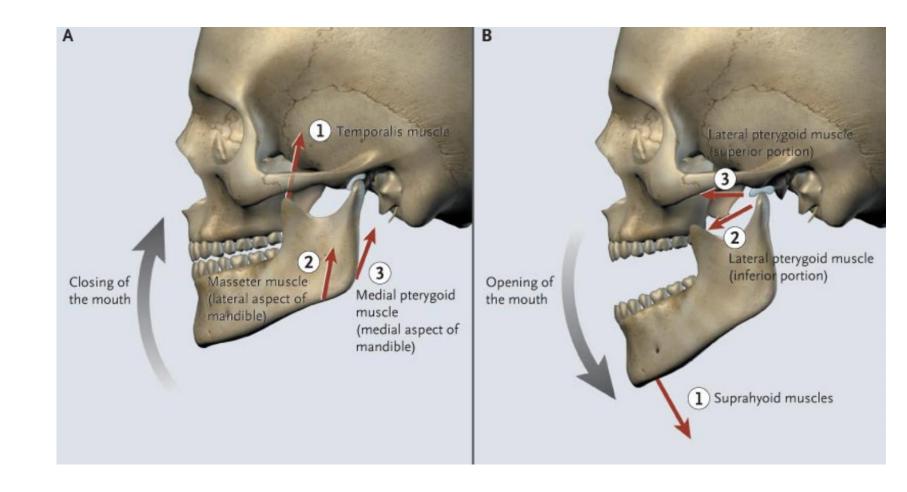
- Musculoskeletal
- Neuropathic
- Neuromuscular
- Psychogenic





Joint Disorders

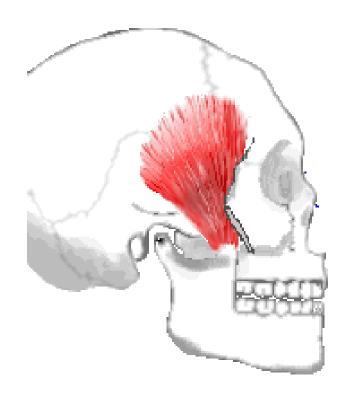
- Joint pain
- Hypomobility
- Hypermobility
- Joint diseases





Muscular Pains

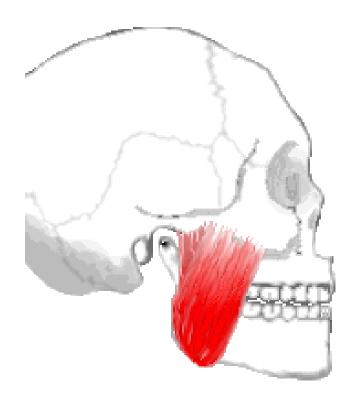
- Muscle pain
- Tendon
- Myositis
- Spasm





Muscular Pains

- Hypertrophy
- Neoplasm
- Movement disorders
- Central / Systemic





Management

- Self-care
- Biobehavioral
- Physical Modalities
- Oral appliances
- Occlusal treatments



Screening Questions

- Difficulty or pain with opening the mouth?
- Does your jaw lock, stick, or go out?
- Difficulty or pain with chewing, talking, or intimacy?
- Jaw stiffness, tightness, or fatigue?
- Jaw joint sounds or noises?
- Pain near the ears, temples, or cheeks?
- Frequent headaches, neck pains, or toothaches?
- Any recent injury to the head, neck, or jaw?
- Recent changes to your bite?
- Past treatments for facial pains or jaw pains?



History

- Chief complaint
- Medical history
- Dental history
- Psychosocial history



Exam Procedures

- Range of motion: opening and lateral excursions
- Palpate the TM joints
- Auscultate or palpate for joint sounds: clicking and crepitation
- Palpate masseters, temporlais, posterior, and lateral neck muscles
- Oral exam: excessive occlusal wear, tooth mobility, buccal mucosa ridging, tongue scalloping
- Symmetry and alignment of face, dental arches



Medications for TMD

Brian E. Cairns, PhD, DrMed, ACPR, RPh

January 11, 2024





Pharmacotherapy for Temporomandibular Disorders (TMD)



Temporomandibular Disorders Pain Mechanisms

Clin Oral Invest (2006) 10:261-268

- Peripheral (tissue specific nociceptor sensitization)
- Central mechanisms (central sensitization & diminished descending inhibition)
- Psychosocial (anxiety, depression, stress, maladaptive coping, parafunctional activities)

Psychosocial adjustment e.g. anxiety, depression Psychological stressors e.g. major or minor life events Neroendocrine function e.g. opioid peptides, Structural/functional substance P, dopamine, impairment serotonin, norepinephrine, TMJ'S, occlusion, muscles up or down regulation of Stress system-related receptors psychosocial variables e.g. personality, maladaptive coping, illness behaviour Physical adjustment e.g. muscular hyperactivity, disability, pain TMD SIGNS AND SYMPTOMS

Fig. 1 A range of psychosocial (central event) and physical variables (peripheral events) may modify or exacerbate the effects of stressors on disease-related outcomes



265

95% of Recommendations for TMD Treatment Involve Analgesic Medications

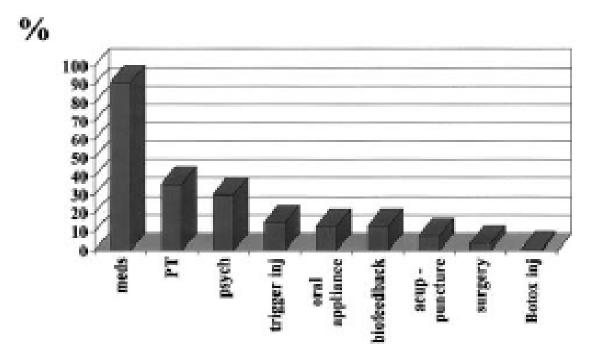
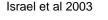


FIGURE 3. Treatment recommendations by the center for patients with chronic orofacial pain, meds, medication; PT, physical therapy; psych, psychiatric management; inj, injections; Botox, botulinum toxin.





Non-Steroidal Anti-Inflammatory Drugs

- Reduction of prostaglandins (PGE₂) through inhibition of cyclooxygenase
- Full benefit may not be seen for 3 weeks or longer, a two-week trial suggested
- Treatments have very low evidence, with short duration studies and low numbers of subjects studied.

Ibuprofen	600mg po TID	OTC, max dose 1800-2400 mg/day
Naproxen	500mg po BID	OTC, max dose 1500 mg/day
Diclofenac	50mg po BID	Increased cardiovascular risk
Diclofenac 2.32%	2g BID to affected area	Facial irritation, may take weeks to
		achieve effectiveness
Piroxicam	20 mg po daily	risk of GI issues high***



Non-Steroidal Anti-Inflammatory Drugs

Chronic oral NSAID use may result in:

- Decrease renal function (avoid in kidney dysfunction)
- Gastrointestinal ulceration (can use with proton pump inhibitor, COX2 selective agent celecoxib not an effective alternative)
- Increased blood pressure in susceptible individuals (uncontrolled HT, ACE inhibitors)
- Anti-platelet actions (avoid in patients treated with anticoagulants).
- Acetaminophen (paracetamol) may be considered if NSAIDs contraindicated, but no evidence of efficacy in TMDs



Skeletal Muscle Relaxants

- Skeletal muscle relaxants act in the central nervous system to reduce motor neuron excitability leading to decreased muscle tone.
- Used for the treatment of myofascial TMD pain, often in combination with an NSAID (e.g. diazepam and ibuprofen).
- Evidence for short-term use only (2-3 weeks).

cyclobenzaprine	10 mg po HS	significant sedation, impaired coordination,
		interacts with antidepressants (MAOIs,
		tricyclics, SNRIs), trazodone
diazepam	5 mg po QID	significant sedation, amnestic properties,
		impaired coordination



Antidepressants

- Enhance endogenous descending inhibition of pain
- Slow onset to full efficacy (several weeks required for effect)
- Tricyclic antidepressants (TCAs), serotonin selective reuptake inhibitors (SSRIs) and serotonin noradrenaline reuptake inhibitors (SNRIs) are commonly used, but few have been subjected to clinical trials in TMD

amitriptylline 25 mg po HS

sedation, dizziness, blurred vision, constipation, dry mouth



Gabapentinoids

- Bind to the $\alpha_2\delta$ subunit of voltage gated calcium channels
- Over time, may decrease transport and synthesis of $\alpha_2\delta$ subunits resulting in decreased calcium influx

dizziness, mental clouding, highly sedating

- May take 8 weeks or longer to see effect
- Some efficacy for myofascial TMD pain

gabapentin 300 mg po daily (HS),

mg

increase by 300 mg q72h until pain controlled or maximum dose of 4200



Botulinum Neurotoxin

- Destroys SNAP 25, a protein critical to vesicular release of neurotransmitters
- Decreased release of acetylcholine from motor neuron axon terminals at the neuromuscular junction lowers muscle tone
- Decreased release of glutamate and neuropeptides from the endings sensory afferent fibers decreases neurogenic inflammation and raises mechanical pain threshold
- Onset of analgesia may be within hours, lasting months between injections

botulinum 25-50 units depending unintended muscle weakness, allergic neurotoxin on muscle q3-4 months reaction



Intraarticular (TMJ) Injection

- Indicated for limited mouth opening, joint pain, and inflammation due to their antiinflammatory actions
- Corticosteroids have a short duration of effect, should be used no more than three times per year – response declines with repeated use

triamcinolone

10-20mg IA q3-4 months

(betamethasone, hydrocortisone, methylprednisolone)

post injection flare, pain, exacerbation of degenerative joint changes (osteoporosis with long term use, risk for osteonecrosis?)

hyaluronic acid

many different protocols

pain, bruising, redness, itching, and swelling



Opioid Analgesics for TMD

- Reserved for moderate to severe chronic TMD pain that has not responded adequately to all other analgesic treatments
- No clinical trials demonstrating the effectiveness of oral opioid analgesics in TMD



Case-Based Learning

Sujay A. Mehta, DMD, MPH January 11, 2024



Case

40-year-old female presents with toothache & consults her local dental office



Dental Office

Routine endodontic procedure with obturation

Rx NSAID, a "dental rinse", and steroid

Returns to Dental Office

- Pt c/o 2 types of pains
- "Nasty"
- No sign of infection but there is limited opening
- Rx physio, Endo, & Oral Med consult

Endodontist

- No obvious dental / endo pathology
- Rx NSAIDS
- Endodontist agrees to Oral Med consult

Oral Med

At this point pt has significant limited opening

Rx Physio and TP injection

 Pt elects to try physio - not keen to an injection to their face

Family MD

History noted for "dental procedure"

Rx antibiotics

Recommended a colleague from oral surgery

Physiotherapist

 History reviewed a change to pain quality after dental procedure with trial to NSAIDS, ABX, and concerns to limited opening

 After a few visits, it seems clear pain is unilateral, but they have an exaggerated pain reaction to rehab

Oral & Maxillofacial Surgeon

- Hx noted for prior Endo
- Current pain and limited opening
- Rx
 - Return to dentist for oral appliance
 - NSAID & muscle relaxant
 - MRI of TMJs

Emergency Department

- Attends ED with complaints of severe pain
- Unilateral pains following dental procedure
- No benefit from NSAIDs and muscle relaxants
- Rx Gabapentin 100 mg bid

Current

- Carbamazepine 400 mg bid
- MRI confirmed vascular loop contacting trigeminal nerve at base of brain
- Pending neurosurgical consult

Medications for Trigeminal Neuralgia

Brian E. Cairns, PhD, DrMed, ACPR, RPh

January 11, 2024





Pharmacotherapy for Trigeminal Neuralgia



Pharmacological Treatment of Trigeminal Neuralgia

- The similarity of signs and symptoms of trigeminal neuralgia and other orofacial pain conditions often complicate the diagnosis.
- First line management of this condition is pharmacological suppression of aberrant afferent discharge.
- There are no proven abortive treatments for paroxysms, so therapy is focused on prophylaxis of attacks and reduction of ongoing pain.
- The effectiveness of pharmacological therapies commonly decrease over time.



Carbamazepine

- Voltage gated sodium channel blocker, anticholinergic
- Electrocardiogram should be undertaken to confirm no atrioventricular conduction abnormalities (contraindication to use)
- 100 mg po daily or BID, increase by 100 mg q2d up to 1600mg maximum dose to relieve pain
- Rare Stevens-Johnson syndrome (SJS) and toxic epidermal necrolysis (TEN) associated with HLA-B*1502 allele
- Regular blood tests to monitor cell counts, electrolytes and organ function required
- Mutagenic, teratogenic
- Common fatigue, drowsiness, cognitive difficulties, disturbed sleep, gait imbalance
- Many drug interactions due to metabolism by CYP3A4



Oxcarbazepine

- Derivative of carbamazepine (200 mg carbamazepine=300 mg oxcarbazepine) with similar mechanism
- Electrocardiogram should be undertaken to confirm no atrioventricular conduction abnormalities (contraindication to use)
- 300-2700 mg po daily (BID to QID), increase dose 300mg q3d
- Common fatigue, drowsiness, cognitive difficulties, disturbed sleep, gait imbalance



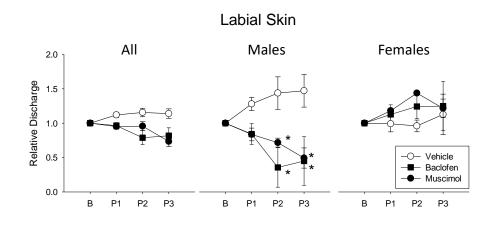
Baclofen

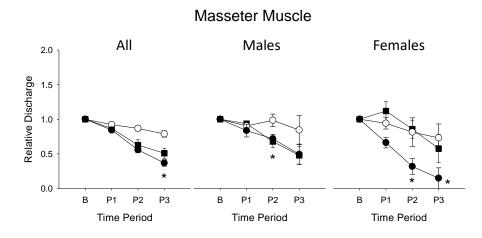
GABA_B receptor agonist

Activation of GABA_B receptors has been shown to inhibit sensory transmission through the trigeminal ganglion

10 mg po daily up to 100 mg, increase by 5 mg q3d, tid with higher doses

Nausea, somnolence, tiredness, and gastrointestinal symptoms

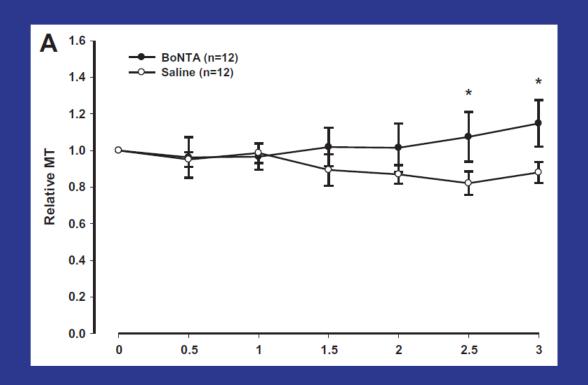


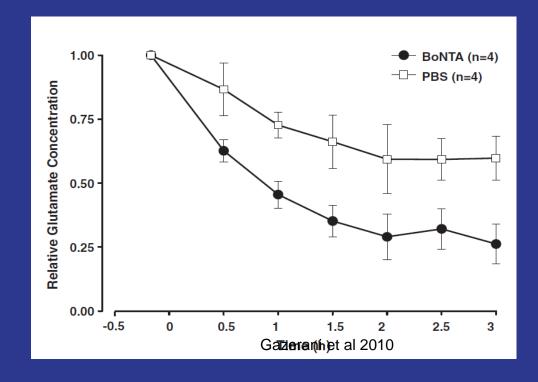




Ranjbar-Ekbatan and Cairns 2020 45

Botulinum Neurotoxin





- Dose 25-100 units into trigger zone q3months
- Transient facial paralysis, facial asymmetry or edema at the injection site



Other Analgesics

- Lamotrigine, gabapentin, pregabalin, phenytoin, tizanidine, pimozide may be used clinically, but relatively little or no evidence supports their use.
- The use of opioids for pain in this condition is not supported by evidence.



Summary

- For TMD, NSAIDs plus or minus skeletal muscle relaxants is a reasonable initial choice for pain.
- Longer term management of TMD could include botulinum neurotoxin (myofascial), or corticosteroids/hyaluronic acid (joint).
- For TN, the drug of choice remains carbamazepine (oxcarbazepine).
- Alternative or add-on therapy includes baclofen, botulinum neurotoxin, or other neuroleptics.
- No evidence for the use of opioid analgesics in either condition.



Practical Applications — Key Takeaways

- Orofacial pain conditions can be complex
- Do what you can do eliminate or rule out oral and dental causes to pain
- Diagnosis requires a solid history and a team approach
- Recognize and refer early



CONSENSUS STUDY REPORT

TEMPOROMANDIBULAR DISORDERS

Priorities for Research and Care

NAM Report 2020

- Urgent need to transform dental education to include OFP and TMD
- Improve inter professional education opportunities
- Centers of Excellence to provide care, coordinate research, and provide telehealth



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Questions & Answers

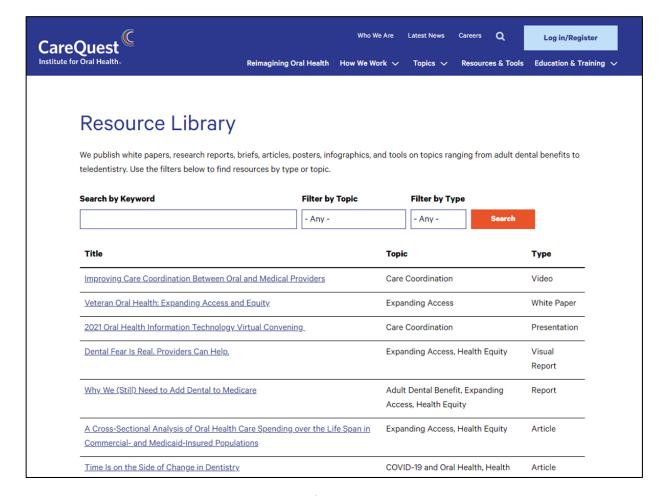




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