

THE SPECTRUM OF CHRONIC POST-TREATMENT PAIN IN HEAD AND NECK CANCER: A DESCRIPTIVE PHENOTYPIC STUDY

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INTRODUCTION

- Chronic post-treatment pain (CPTP) is a major survivorship issue in head and neck cancer (HNC), with prevalence estimates as high as 72% in some studies ^{1,2}.
- Despite this, current frameworks treat it as a single entity, obscuring clinical variance and heterogeneity.
- This study prospectively characterized CPTP by adapting the International Classification of Orofacial Pain (ICOP) and International Classification of Headache Disorders, third edition (ICHD-3) criteria to the HNC context, defining five distinct phenotypes analogous to primary pain disorders of the head and neck to estimate **prevalence, overlap, and clinical associations**.

METHODS

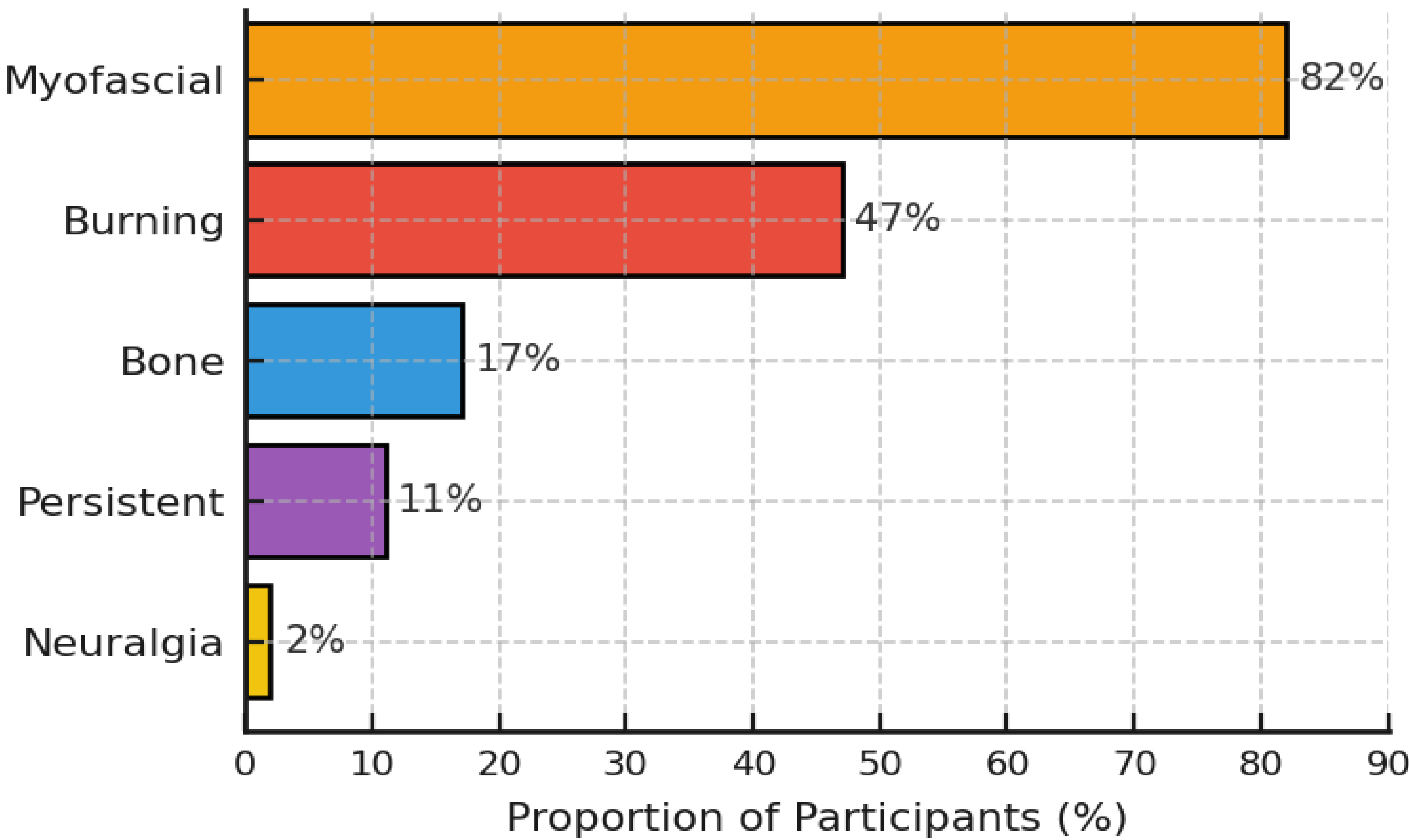
- This prospective observational study included 100 HNC survivors ≥3 months post-treatment with chronic pain.
- ICOP and ICHD-3 criteria were adapted to create 5 pain phenotypes: **orofacial bone pain, myofascial pain, neuralgia, persistent pain, and burning disorder**.
- A single examiner applied these criteria to characterize participants’ CPTP into phenotypes using their clinical history, examination, and radiographic findings.
- Pain intensity, distribution, and associated clinical features were recorded.

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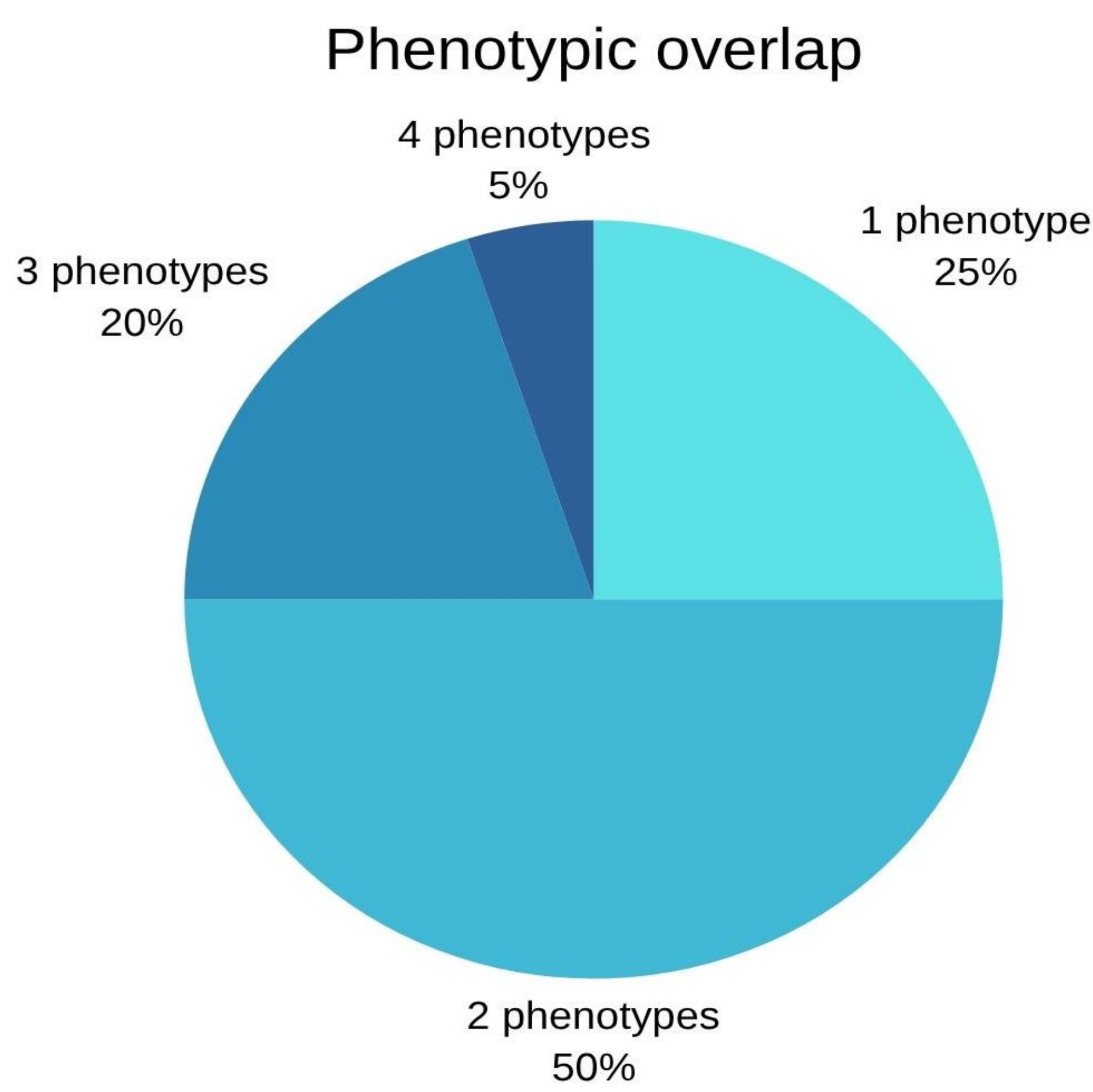
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RESULTS

- Among 100 HNC survivors (mean age 49, 54% male), 81% had squamous cell carcinoma and 52% had stage IV disease.
- Most underwent multimodal therapy: surgery 82%, radiation 93%, chemotherapy 20%.
- Pain was constant in **85%**, mean intensity **5.3 ± 1.8** (out of 10 on the numeric verbal pain rating scale) with highest intensity **7.7 ± 1.3**, affecting **5.8 ± 3.3** sites, mostly lateral cervical (61%) and masseteric (60%). The most common descriptors were dull ache and burning, predominant phenotypes were **myofascial** (82%), and **burning** disorder (47%).



- Patients exhibited one or more concurrent phenotypes, with a mean **2.1 ± 0.9** concurrent phenotypes per patient.



- Significant associations were identified between specific clinical features and certain pain phenotypes:

Sign/symptom	Significantly Associated Phenotype
Xerostomia	Burning pain (p<0.01)
Sensory alteration	Burning pain (p<0.05)
Tightness	Myofascial pain (p<0.001)
Trismus	Persistent and myofascial pain (p<0.01)

CONCLUSION

- CPTP in HNC survivors is not a single entity but an amalgamation of overlapping phenotypes **resembling primary pain disorders**.
- Recognizing these patterns highlights shared mechanisms and supports the need for **phenotype-based, personalized management strategies** to improve patients’ quality of life.

