A systematic review of treatment for patients with burning mouth syndrome

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Abstract

Background: Burning mouth syndrome is a chronic idiopathic intractable intraoral dyseaesthesia that remains a challenge to clinicians due to its poorly understood pathogenesis and inconsistent response to various treatments.

Aim: This review aimed to study the short- (<3 months) and long-term (>3 months) effectiveness and sustainable benefit of different burning mouth syndrome treatment strategies and the associated side effects.

Materials and methods: Randomised controlled trials of burning mouth syndrome treatment compared with placebo or other interventions with a minimum follow up of 2 months were searched from the PubMed, Embase and Cochrane database (published to July 2020).

Results: Twenty-two studies were selected based on the inclusion and exclusion criteria and analysed. Nine categories of burning mouth syndrome treatment were identified: Anticonvulsant and antidepressant agents, phytomedicine and alpha lipoic acid supplements, low-level laser therapy, saliva substitute, transcranial magnetic stimulation, and cognitive behaviour therapy. Cognitive behaviour therapy, topical capsaicin and clonazepam, and laser therapy demonstrated favourable outcome in both short- and long-term assessment. Phytomedicines reported a short-term benefit in pain score reduction. The pooled effect of alpha lipoic acid (ALA) pain score improvement was low, but its positive effects increased in long term assessment.

Conclusion: A more significant volume in terms of sample size, multi-centres, and multi-arm comparison of therapeutic agents with placebo and longitudinal follow-up studies is recommended to establish a standardised burning mouth syndrome treatment protocol. Further studies are required to assess the analgesic benefits of topical clonazepam and capsaicin, alternative medicines with neurodegenerative prevention capability and psychology support in treating burning mouth syndrome and reducing systemic adverse drug reactions.

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Keywords

Burning mouth syndrome, glossodynia, treatment, systematic review

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Introduction

Burning mouth syndrome (BMS) is defined as idiopathic orofacial pain with intraoral burning or dyseaesthesia recurring daily for more than 2 hours per day and more than 3 months, without any identifiable causative lesions, with and without somatosensory changes in International Classification of Orofacial Pain, 2020 (1). BMS prevalence ranges from 0.1% to 3.9% and is primarily present in postmenopausal women aged between 50 and 70 (2,3). BMS commonly manifests as burning, prickling, tingling, itching or numbness.

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affecting the tongue, lip, palate, gums and other oral mucosae (4). The pain intensity increases throughout the day and peaks in the late evening (5). Patients often complain of dysgeusia, xerostomia, altered sensation in the oral mucosa, and psychological issues such as anxiety and depression. The pathogenesis of BMS has been hypothesised to be associated with psychological disorders (6) and peripheral and central neuropathy (7), but at present it is classified as idiopathic chronic pain (1). Diagnosing and managing patients with BMS remains a challenge to clinicians due to its poorly understood pathogenesis and inconsistent and limited response to various treatments. Besides, it has an exceptionally low spontaneous remission prevalence of 3–4% after 5–6 years of diagnosis (8). There are no global guidelines on BMS treatment, and published review articles included clinical studies with limited follow up periods (<2 months) (9–11). Based on the current universal ICOP criteria, the diversity of BMS patients’ underlying pain mechanism, and the difference in evidence on short- and long-term benefit of treatment in BMS (11), we sought to conduct a systematic review on different therapeutic strategies for patients presenting with BMS, with the question “which range of treatments have effective short (<3 months) and long-term (>3 months) outcomes in improving the pain symptoms in BMS patients?” Parallel with the aim of providing a personalised treatment for each patient, the sustainability of a treatment efficacy and patients’ compliance and response towards the therapy and its side effects should be considered.

Methodology

Search strategy

The study was carried out following the PRISMA guidelines (12). An electronic search on PubMed Medline (1946 to 1 July 2020), Embase Ovid (1980 to 1 July 2020), Cochrane Database of Systematic Reviews (1 July 2020) and Cochrane Central Register of Controlled Trials (CENTRAL) (1 July 2020) was conducted based on the combination of the following keywords: “burning mouth syndrome or glossalgia or stomatodynia AND treatment or therapy or therapeutic or management”. This review includes all randomised and controlled clinical trials with a placebo published in the English language. The included studies should state that the diagnosis of BMS is based on the absence of local and systemic pathological contributing factors and have a minimum follow up of treatment of 2 months. This systematic review was registered in PROSPERO (Protocol ID: CRD42020160892). We also performed a manual search on all included clinical trials in published systematic review articles for any potentially relevant studies.

Study selection

The search results were screened based on the relevant title and abstract by two independent authors. Where information from the abstract was inadequate to allow a decision, a full report was obtained. The full text was obtained for articles fulfilling the inclusion criteria. Any disagreements were resolved by discussion between the authors, and the review authors were not blinded to articles’ authorship. Studies meeting the inclusion criteria underwent data extraction and were evaluated for study risk of bias. The following data were obtained and recorded in a standardised pro-forma sheet on author and year of publication; study design or methodology; sample size and participant inclusion and/or exclusion criteria; types of intervention and follow-up time; the outcome and/or adverse effect from the intervention; statistical methods employed (Table 1).

Assessment of risk of bias

We used the Cochrane risk of bias assessment tool (13), which is based on seven main domains (Table 2). Each study was categorised based on the overall risk category and classified as low, unclear, or high risk. The quality of all included articles was assessed using the GRADE (14).

Outcome analysis

We analysed outcome data based on short term (≥2 month to ≤3 months) and long term (>3 months) changes in symptoms. The assessment method used in the included studies should be of equal measure. The standardised mean difference (SMD) in pain score (VAS) of treatment groups and placebo and their relative risk ratio (RR) for BMS pain improvement was recorded from the relevant studies with the 95% confidence interval (CI) where possible. Estimates of effect (and associated CI) were combined and pooled for studies reporting the same treatment.

Statistical analysis

Mean difference (MD) of the pre- to post-treatment VAS change scores were extracted from studies. For each study with comparisons between treatment and placebo at short term (≤3 months) and/or long term (>3 months), standardised mean differences (SMDs) of the VAS scores were calculated using pre-to-post-intervention change score (means) and post-intervention SDs (rather than change score SDs...
Table 1. Summary of included studies and quality of evidence.

| Author/Year | Intervention | Sample size: Mean age (years) (study/control) | Outcome assessment method | Finding summary | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|-------------|--------------|-----------------------------------------------|---------------------------|----------------|-----------------------|----------------|---------------------------|
| (20)        | Clonazepam (0.5 mg) Dosage: 0.5 mg Durations: 9 weeks Route: Oral | 10/10; 67.5/65.4 | - Numerical pain ratings (0–10) - BDI - ZMS - Taste test - Smell test - Salivary flow rate | - NPS: Significant difference between clonazepam (MD: 2.9) and placebo (MD: -1.5), (p = 0.011) - Taste and saliva: Clonazepam group show significant increase in taste score (p = 0.023) and salivary flow (p = 0.033) - No significant difference between clonazepam and placebo in taste (p = 0.83) and salivary flow (p = 0.060) - Depression and mood. | | No side effect on psychology | Moderate |
| (19)        | Clonazepam (0.5 mg) Dosage: 0.5–2.0 mg/day Duration: 6 months Route: Topical. Dissolved in mouth and spat out after 3 min | 33/33; 64.9/64.9 | - VAS | - Significant decrease in VAS for Clonazepam (MD: -4.7) - 23 study group improved more than 50% (p < 0.05) and three were totally asymptomatic - Not significant decrease in control group (MD: -3.2) - Reduced in tasted alteration and dryness in clonazepam group | | Test group: Five had sleepiness but did not require termination of treatment | Moderate |
| (39)        | Clonazepam (2 mg) Dosage: 2 mg/day Duration: 4 months Route: Oral Pregabalin (150 mg) Dosage: 150 mg/day Duration: 4 months | 25; 43; 25; 45; 25; 42 | - VAS | - Significant reduction in VAS score clonazepam (MD: -4.1, p < 0.001) - Significant reduction in pregabalin VAS score (MD: -4.7, p < 0.001) | | Four dizziness, two transient diarrhoea, two myalgia, Three increased appetite, one | Very low |

(continued)
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary Short term (<3 months) | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|------------|--------------|-----------------------------------------------|---------------------------|----------------------------------------|----------------------|----------------|--------------------------|
| (21)       | Route: Oral  |ALA (600 mg)  
Dosage: 600 mg/day  
Duration: 4 months  
Route: Oral  
Trazodone (100 mg)  
Dosage 1: 100 mg  
Duration 1: 4 days  
Daily for 4 days  
Dosage: 200 mg Duration: 8 weeks  
Route: Oral | 11/17; 61.1/NA  
VAS  
MPQ  
BDI  
Global assessment | ● Eight in study group and 13 in placebo reported reduction in pain  
● Pain intensity was significantly decreased in (p < 0.01) in both trazodone (MD: −1.4) and placebo group (MD: −1.3)  
● No significant difference between both groups in treatment effect and treatment by time interaction for pain intensity  
● No significant differences between the groups MPQ for influence of pain on eating, speaking, sleeping or for the suffering caused by the pain  
● No significant difference between both group in the patient's global assessment of improvement or benefits of the treatment  
● Significant decreased in BDI for both groups in the depression score (p < 0.01)  
|  | ● No significant reduction in VAS score ALA (MD: −0.7)  
● Two mild nausea, one myalgia  
| Moderate |
| (30)       | Crocin (15 mg) | 26; 52.9 | VAS  
HAD  
DSM IV psychiatric diagnosis | No significant difference in VAS mean score between crocin and citalopram (p = 0.98) |  |  | Low |

(continued)
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary | Adverse Effect | Quality of Evidence (Grade) |
|------------|--------------|-----------------------------------------------|---------------------------|----------------|---------------|--------------------------|
|            | Dosage: 10 mg for first week followed by 20 mg daily Duration: 11 weeks Route: Oral | 30/30; 45/NA | BMS symptomatology change scale (worsening; unchanged; slight improvement; decided improvement; resolution) | Significant reduced in crocin VAS (MD: -7.8, \( p < 0.001 \)) Average recovery percentage of burning mouth score at the end of treatment for both crocin and citalopram was 87.45% No significant difference in depression and anxiety between crocin and citalopram (\( p = 0.76 \)) Average recovery percentage of depression score at the end of treatment for crocin was 30.57% and citalopram was 30.79% Average recovery percentage of anxiety score at the end of treatment for crocin was 15.44% and citalopram was 15.40% | No adverse effect reported | Low |
| (22) ALA (200 mg) Dosage: 600 mg/day Duration: 2 months Route: Oral Participant showing deterioration of symptoms within 4 months will be given another 1 month of supplement | 23/16; 67/59.3 | VAS | Statistically significant symptom improvement with ALA (97%) compares with control (40%) 87% ALA patients showed resolution or a decided improvement in symptom but none in control None of ALA group has worsening of BMS symptoms but 20% in control | 73% of ALA remains significantly stable (no changes) but 83% of control group shows significant deterioration | Test group: One patient has gastrointestinal upset. | Low |
| (24) ALA Dosage: 800 mg/day Duration: 8 weeks Route: Oral | | | No significant difference \( (p = 0.14) \) between ALA (MD: 2.2) or control (MD: 3.8) | | | Low |
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary Short term (<3 months) | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|------------|--------------|-----------------------------------------------|---------------------------|---------------------------------------|----------------------|---------------|---------------------------|
| (26)       | ALA (200 mg) | Dosage: 600 mg/day Duration: 2 months Route: Oral | 29/25; 62.13/62.13 | VAS | 64% in ALA and 27.6% in control group reported improvement of symptoms | No ALA patients and five control patients reported worsening of symptoms | Statistically significant differences between both groups ($p = 0.009$) | Low |
| (23)       | ALA (400 mg) and vitamin (C, PP, E, B6, 2, 1, 12, and folic acid) Dosage: 800 mg/day Duration: 8 weeks Route: Oral ALA: 400 mg Dosage: 800 mg/day Duration: 8 weeks Route: Oral GABA Dosage: 300 mg/day Duration: 60 days ALA: 20 GABA: 20 | 18/20; 67.3/NA | N/A | No adverse effect reported | Moderate |
| (25)       | ALA Dosage: 600 mg/day Duration: 60 days Route: Oral GABA Dosage: 300 mg/day Duration: 60 days | 20/60; 57.5/NA | Numerical category of burning scale: Category 1: negative changes (deterioration) Category 2: no changes | No significant difference between all three groups in MPQ | No significant difference between three groups in MPQ | Adverse effects appeared very mild. | Moderate |

(continued)
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|-------------|--------------|-----------------------------------------------|---------------------------|----------------|-----------------------|---------------|--------------------------|
| Route: Oral • ALA and GABA Dosage: 600 ALA and 300 GABA/day Duration: 60 days Route: Oral | Category 3: with positive changes (improvements) Category 4: with total recovery | Negative: 0%; no change: 50%; positive and total recovered: 50% GABA 5.7× higher than control group ALA + GABA: Negative: 0%; no change: 30%; positive and total recovered: 70% ALA + GABA 13.2× higher than control group Significant level of positive burning changes between group (p < 0.001) | | | | |
| (33) • ALA (400 mg) Dosage: 800 mg/day Duration: 8 weeks Route: Oral • Capsaicin (250mg chilli powder in 50 ml) Dosage: 750 mg/150 ml/day Duration: 8 weeks Route: Topical – oral rinse • Lysozyme lactoperoxidase (Biotene) Five times per day Duration: 8 weeks Route: Topical – oral rinse | Size Short term: ALA: 14 Capsaicin: 14 Biotene: 14 Control: 14 Long term: ALA: 9 Capsaicin:9 Biotene: 9 Age ALA: 64 Capsaicin: 62 Biotene: 62 | VAS | Significant improvement in VAS (p < 0.001) ALA: 57% improved (MD: −2.1). Capsaicin: 76% improved (MD: −3.2) Biotene: 57% remain unchanged (MD: −1.7) No statistically difference in VAS between groups ALA, capsaicin and Biotene No significant difference VAS improvement in control group All study groups, ALA, capsaicin and biotene, were statistically superior to control groups Significant decreased of spontaneous burning intensity between umPEA group (MD: −3.8) and control group (MD: −1.3) (p = 0.001) | | | |
| (31) Ultramicronised Palmitoylethanolamide (umPEA) (600 mg) Dosage: 1200 mg Duration: 60 days Route: Sublingual | 13/16 NA | NRS (scale 0 to 10) | Only capsaicin group shows significant reduction in VAS score (MD: −2.9) with 67% improved ALA (MD: −1.8) and Biotene (MD: −1.8) failed to show statistically significant of VAS score improvement with 55% remain unchanged for Biotene and 55% improved with ALA No difference in trend of VAS in control group | No statistically significant difference between umPEA (MD: −2.4) and control group (MD: −1.4). | | Low | |

(continued)
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary Short term (< 3 months) | Long term (> 3 months) | Adverse Effect | Quality of Evidence (Grade) |
|------------|--------------|-----------------------------------------------|---------------------------|---------------------------------------|------------------------|----------------|-----------------------------|
| (29)       | Herbal catuama (310 mg) Dosage: 620 mg/day Duration: 8 weeks Route: Oral | 30/30; 63.6/61.5 | VNS (0–10) | At 8 weeks reduction in symptoms of test group was 52.4% and control group 24.2% | | Test group: One patient with somnolence and weight gain, one insomnia, three exacerbation of symptoms in first week of treatment | Moderate |
| (28)       | Hypericum perforatum (300 mg) Dosage: 900 mg/day Duration: 12 weeks Route: Oral | 19/20; 65.9/63.9 | VAS | No significant difference between study (MD: -1.8) and control group (MD: -1.1) in VAS (p = 0.222) | | Test group: One had severe headache in the fifth week of therapy | Moderate |
| (27)       | Lycopene-enriched extra virgin oil (300 ppm) Dosage: 900 ppm/day Duration: 12 weeks Routes: Topical spray and swallowed | 26/24; 61.7/64.9 | VAS* (grade 1 to 10) | Significant reduction in VAS score in both pain (MD: -3.0; p < 0.001) and burning (MD: -1.0; p = 0.003) symptoms | | No adverse effect reported | Low |

(continued)
Table 1. Continued.

| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|-------------|--------------|-----------------------------------------------|---------------------------|----------------|----------------------|----------------|--------------------------|
| (18)        | N-acetyl-5-methoxytryptamine. Melatonin (MLT) (3 mg) | 16/16; 64.4/64.4 | VAS, Number of sites, Patient global impression of pain changes, Symptom response categories (worse; no change; mild improvement; moderate improvement; strong improvement), MOS, HRS | No significant difference between MLT (MD: −0.6) and placebo (MD: −1.1) group in VAS score | Four MLT group and three control group reported improvement in pain changes | No adverse effect reported | Moderate |

Test group: 40% of patients dropped out because of side effects: Four self-reported heavy tremor, sexual disturbances, blurred vision, and severe heaviness; three lack of efficacy or pain improvement; one lost to follow-up.

| (34)        | Low level laser therapy: IR1W: 830 nm wavelength, 100 mW output power; continuous emissions, 3.57 W/cm², 5 J energy per point, 176 J/cm² radiant exposure, application time 50 sec per point. Duration: One session per week for 10 weeks. Total 10 sessions. IR3W: 830 nm wavelength, 100 mW output power, continuous emissions, 3.57 W/cm², 5 J energy per point, 176 J/cm² radiant exposure, | 20/19; 63.6/61.5 | VAS (0–100), VNS (0–10), OHIP 14 | VNS: Significant difference between IR1W laser (MD: −4.45) and control (MD: −2.53) at 11 weeks; p = 0.005 | VAS: Significant difference between IR1W laser (MD: −4.92) and control at 11 weeks; p = 0.004. VNS: Significant difference between IR3W laser (MD: −5.1) and control (MD: −2.53) at 11 weeks; p < 0.0001 | No adverse effect reported | Low |

(continued)
Table 1. Continued.

| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary | Short term (<3 months) | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|-------------|--------------|-----------------------------------------------|---------------------------|----------------|------------------------|------------------------|---------------|--------------------------|
|             | Low level laser therapy (LLLT) | 10/10; 60.3/67.6 | VAS, SF-36, OHIP14, EES, SCL 90-R, MPQ | VAS: | 90% LLLT and 20% control reported improvement | N/A | Low |
| (36)        | Dose: 810 nm wavelength, 12 J/cm² per session in a continuous mode | Duration: Twice a week session for 5 weeks consecutively Total 10 sessions |             | VAS: | Pain decreased significantly (p = 0.005) in the study group (MD: -2.9) versus control group (MD: 0.5) |             |             |
|             | Red laser: 685 nm wavelength, 35 mW output power, continuous emissions, 1.25 W/cm², 2 J energy per point, 72 J/cm² radiant exposure, application time 58 sec per point. Duration: Three sessions per week for 3 weeks. Total 9 sessions |  | VAS: | No significant difference between IR1W, IR3W and red laser group |             |             |
|             | VNS: | No significant difference between red laser (MD: -3.74) and control (MD: -2.53) at 11 weeks; p = 0.12 |             | VAS: | No significant difference between red laser and control at 11 weeks, p = 0.13 |             |             |
|             | VNS: | No significant difference between IR3W laser (MD: -53.0) and control at 11 weeks; p < 0.0001 |             | VNS: | No significant difference between red laser and control at 11 weeks, p = 0.13 |             |             |
|             | VNS: | No significant difference between IR1W, IR3W and red laser group |             | VNS: | No significant difference between red laser and control at 11 weeks, p = 0.13 |             |             |
|             | VNS: | No significant difference between IR1W, IR3W and red laser group |             | VNS: | No significant difference between red laser and control at 11 weeks, p = 0.13 |             |             |
|             | VNS: | No significant difference between IR1W, IR3W and red laser group |             | VNS: | No significant difference between red laser and control at 11 weeks, p = 0.13 |             |             |
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary Short term (<3 months) | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|-------------|--------------|---------------------------------------------|--------------------------|--------------------------------------|------------------------|----------------|--------------------------|
|            |              |                                             |                          | EES:                                 | control group in all categories                      |
|            |              |                                             |                          |                                     | EES: No significant difference in LLLT (MD: −0.1, p = 0.83) and between control (p = 0.32) |
| (32)       | Urea 10%     | 12/13; 66.3/58.4                           | EDOF-HC                  | Seven in study group and eight in control group have reduction in pain |
|            | 3–4 times daily |                                             | Xerostomia questionnaire  | No difference between study and control group pain intensity (p = 0.88); salivary flow (xerostomia) (p = 0.32); gustation (sweet p = 0.38, salty p = 0.69, sour p = 0.69, bitter p = 0.69); olfaction (p = 0.96); corneal reflex right (p = 0.20) and left (p > 0.99) |
|            | Duration: 3 months |                                             | QST                      | No significant differences in the somatosensory test between group (p > 0.05) |
|            | Route: Topical |                                             |                          | N/A                                  | Low |
| (35)       | Transcranial magnetic stimulation (rTMS) | 12/8; 63.4/64.4 | VAS                      | Significant reduction in pain with rTMS group (MD: −3.1, p = 0.002) |
|            | Dosage: total of 30,000 pulses |                                             | BPI                      | 75% reported > 50% decrease in BMS pain intensity |
|            | Duration: 10 sessions |                                             | SF-MPQ                  | Significant difference between rTMS and sham group (MD: −2.8, |
|            | Route: Transcranial |                                             | PHQ-9                   | | |
|            |              |                                             | PGIC                    | | |
|            |              |                                             | CGI-I                   | | |
|            |              |                                             |                         | Seven in rTMS group and five in sham group had headache at the beginning treatment but very mild and tolerated and disappeared in one or two days |
|            |              |                                             |                         | | |

(continued)
Table 1. Continued.

| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|-------------|--------------|-----------------------------------------------|---------------------------|----------------|-----------------------|---------------|-----------------------------|
|             | Tongue protector | 25/25; 61.0/61.4 | VAS, HAD, OHIP-49, SF-36 | VAS: Significant difference ($p < 0.001$) between active (MD: $-3.6$) and control (MD: $-1.4$) group | | No adverse effect observed | Very Low |
|             | Dosage: 15 min 3 times/daily | | | | | | |
|             | Duration: 2 months | | | | | | |
|             | Route: Oral appliance | | | | | | |

- BPI: Significant improvement for rTMS group (MD: $-2.1$, $p = 0.003$) and not in control group
- SFMPQ: Non-significant difference in affective score and present pain intensity in rTMS (MD: $-1.2$) and sham (MD: $-0.8$) group
- PHQ-9: No significant difference in rTMS (MD: $-5.6$) and sham group (MD: $-1.0$)
- PGIC: Significant difference in rTMS (MD: $3.3$, $p < 0.01$) but not in sham group (MD: $1.4$)
- CGI-I: Significant improvement in rTMS (MD: $-2.3$, $p < 0.01$) but not in sham group (MD: $-0.62$)
- HAD: Depression non-significant ($p = 0.205$) between active (MD: $-1.0$) and control groups (MD: $-0.04$)
- Anxiety non-significant ($p = 0.69$) between active (MD: $-0.1$) and control groups (MD: $-0.2$)
| Author/Year | Intervention | Sample size; Mean age (years) (study/control) | Outcome assessment method | Finding summary | Long term (>3 months) | Adverse Effect | Quality of Evidence (Grade) |
|------------|--------------|---------------------------------------------|---------------------------|----------------|-----------------------|---------------|--------------------------|
|            | Cognitive therapy (CT) | 15/15                                       | VAS (1–7)                 | • OHIP-49: Significant difference ($p=0.008$) between active (MD: -18.4) and control (MD: -1.9) |                     | N/A           | Low                      |
|            | One hour once a week A total of 12–15 sessions |                               |                           | • SF-36: Significant difference ($p<0.05$) between active and control group in physical role, bodily pain, general health, emotional role |                     |               |                          |
|            |              |                                             |                           | • Tongue protector group has better oral health (OHIP) and quality of life (SF-36) |                     |               |                          |
| (37)       |              |                                             |                           | • VAS (1–7) | Significant reduction in pain symptoms (VAS) in CT group (MD: -2.8, $p<0.001$) than control group |                     |               |                          |
|            |              |                                             |                           | • Significant reduction in CT group pain symptoms (VAS) (MD: -3.6, $p<0.001$) than control group |                     |               |                          |

VAS: Visual analogue scale; VNS: visual numerical scale; NRS: Numeric rating scale; MPQ: McGill Pain Questionnaire; BPI: Brief Pain Inventory; EDOF-HC: Orofacial Pain Clinic Questionnaire; SF-36: Short Form Health Survey (SF-36); OHIP 14: Oral Health on Quality of Life; HAD: Hospital Anxiety and Depression Scale; BDI: Beck Depression Inventory; ZMS: Zerssen Mood Scale; PGIC: Patient Global Impression of Change; CGI-I: Clinical Global Impression for global Improvement Scale; EES: Epworth Sleepiness Scale; STQ: Patient Health Questionnaires-9; HRS: Hamilton Rating Scale; SCL-90-R: Symptom Checklist-90-R; MOS: Medical Outcomes Survey of Sleep Scale; QST: Quantitative somatosensory testing; MD: mean difference from baseline; ALA: Alpha lipoic acid; GABA: Gabapentin; N/A: Not available.
### Table 2. Risk of biased analysis of included studies.

| Treatment                          | Author                                                                 | Random sequence generation | Allocation concealment | Blinding of participants | Blinding of outcome assessors | Incomplete outcome data | Selective reporting | Other bias |
|------------------------------------|------------------------------------------------------------------------|-----------------------------|------------------------|---------------------------|-------------------------------|-------------------------|---------------------|------------|
| Clonazepam Systemic (oral)         | Heckmann et al. 2012                                                  | +                           | +                      | +                         | +                             | –                       | –                   | –          |
|                                    | Çınar et al. 2018                                                    | ?                           | –                      | –                         | –                             | –                       | +                   | +          |
| Clonazepam Topical (rinse)         | Rodriguez de Rivera-Campillo et al. 2010                              | +                           | ?                      | +                         | +                             | –                       | –                   | +          |
| Pregabalin                         | Çınar et al. 2018                                                    | !                           | –                      | –                         | +                             | +                       | –                   | !          |
| GABA                               | Lopez-D'alessandro et al. 2011                                       | +                           | +                      | +                         | –                             | –                       | –                   | –          |
| Trazodone                          | Tammiala-Salonen et al. 1999                                         | +                           | +                      | +                         | +                             | +                       | –                   | –          |
| Citalopram                         | Pakfetrat et al. 2019                                                | ?                           | –                      | +                         | +                             | +                       | +                   | –          |
| Crocin                             | Pakfetrat et al. 2019                                                | !                           | –                      | +                         | +                             | +                       | +                   | –          |
| ALA                                | Femiano et al. 2002                                                  | ?                           | –                      | +                         | +                             | –                       | –                   | –          |
|                                    | Lopez-Jornet et al. 2009                                             | +                           | +                      | +                         | +                             | –                       | –                   | –          |
|                                    | Palacios-Sanchez et al. 2015                                         | ?                           | ?                      | +                         | –                             | –                       | –                   | –          |
|                                    | Carbone et al. 2009                                                  | +                           | –                      | +                         | +                             | –                       | –                   | –          |
|                                    | Lopez-D'alessandro et al. 2011                                       | +                           | +                      | +                         | +                             | –                       | –                   | –          |
|                                    | Marino et al. 2010                                                   | +                           | –                      | –                         | +                             | +                       | –                   | –          |
|                                    | Çınar et al. 2018                                                    | !                           | –                      | –                         | +                             | +                       | +                   | !          |
| ALA + Vitamin                      | Carbone et al. 2009                                                  | +                           | –                      | +                         | +                             | !                       | +                   | +          |
| ALA + GABA                         | Lopez-D'alessandro et al. 2011                                       | +                           | +                      | ?                         | +                             | +                       | –                   | –          |
| Capsaicin Topical (Rinse)          | Marino et al. 2010                                                   | +                           | –                      | ?                         | +                             | +                       | –                   | –          |
| Ultramicronised palmitoylethanolamide | Ottaviani et al. 2019                                           | +                           | –                      | +                         | +                             | –                       | +                   | –          |
| Herbal catuama                     | Spanemberg et al. 2012                                               | +                           | +                      | +                         | –                             | –                       | +                   | –          |
| Hypericum perforatum               | Sardella et al. 2008                                                 | +                           | +                      | +                         | +                             | –                       | –                   | +          |
| Lycopene-enriched extra virgin oil | Cano-Carrillo et al. 2014                                             | +                           | +                      | +                         | +                             | –                       | –                   | –          |
| Melatonin                          | Varoni et al. 2018, 2019                                             | +                           | +                      | +                         | +                             | –                       | –                   | –          |
| Low level laser therapy            | Spanemberg et al. 2015                                               | ?                           | ?                      | ?                         | ?                             | +                       | +                   | ?          |
|                                    | de Pedro et al. 2020                                                 | ?                           | –                      | +                         | +                             | –                       | –                   | –          |
|                                    | da Silva et al. 2014                                                 | ?                           | –                      | +                         | +                             | –                       | –                   | –          |
| Urea Topical (Rinse)               | Lysozyme lactoperoxidase Topical (Rinse)                              | +                           | –                      | ?                         | +                             | –                       | –                   | –          |
|                                    | Transcranial magnetic stimulation                                    | +                           | ?                      | ?                         | ?                             | –                       | –                   | –          |
| Tongue protector                   | Lopez-Jornet et al. 2011                                             | +                           | ?                      | –                         | +                             | +                       | –                   | +          |
| Cognitive therapy                  | Bergdahl et al. 1995                                                 | ?                           | –                      | –                         | +                             | +                       | –                   | +          |

ALA: alpha lipoic acid; GABA: gabapentin; ‘?': unclear risk; ‘+': low risk; ‘- ': high risk
which were not provided in several studies). Means and/or standard deviations for baseline and post-treatment pain intensity were calculated for two studies based on the length of error bars in graphs and a ruler and two other studies using raw data (provided in papers). Continuous data were pooled using the Hedges g statistic as a formulation for the SMD under the fixed effects model. For categorical (dichotomous) outcomes (e.g. n ≥ vs. n < 50% decrease in VAS pain intensity, or number of patients demonstrating improvement from baseline versus the number showing no change/worsened score), relative risks (RRs) and associated 95% CI were calculated to express the estimate of treatment effect (15). Where zeros caused problems with the computation of the RR or its CIs, 0.5 was added to frequency cells (16,17). Where appropriate, RR data were pooled (under a fixed effect model). Formal meta-analyses were not performed in this review due to the heterogeneity of the included studies’ methods and outcome data such as varying assessment times within short- and long-term testing periods, differences in treatment regime (e.g. timing or dosage of medication administration), different outcome assessments of burning or general pain improvement, and incomplete data (e.g. variance not reported).

Results
A total of 95 full text published articles were reviewed; 22 were included in this review (Table 1), and 73 were excluded (Table 3). Figure 1 shows the study selection flow process.

Characteristics of studies
All 22 included studies were randomised controlled clinical trials with one triple-blinded study (participant, caretaker, and assessor) (18), 14 double-blinded studies (19–32), four single-blinded studies (participants) (33–36), and three non-blinded studies (37–39). Three of the four single-blinded studies have a common concern with assessor blinding as they involved patient-reported outcomes (33,34,36). Fourteen (64%) studies described the method employed in generating the randomised sequence; online website or computer software, and randomisation tables, balls, or blocks (18–21,23–25,27–29,31,33,35,38). Eight studies reported on examiners’ allocation concealment (18,20,21,24,25,27–29). Five studies (22%) have a high risk of attrition bias (24,26,29,32,35), and eight studies (36%) have a high risk of reporting bias (20,22,24–27,31,32). In the reviewers’ opinion, none of the studies was graded high, with two very low (38,39), 12 low (22,24,26,27,30–37) and eight moderate (18–21,23,25,28,29).

Twenty studies were randomised controlled trials (RCT) with placebo parallel-group comparison (18–29,31–38), and two studies were a comparison between different parallel cohort treatment groups (30,39). The 20 placebo-controlled randomised trials consisted of 16 trials with two-arm (18–22,24,26–29,31,32,35–38) (14 intervention vs. placebo and two non-intervention vs. intervention), one trial with three-arm (23), and three trials with four arms (25,33,34) comparison between intervention and placebo. The remaining two non-placebo RCT were two-arm (30) and three-arm (39) trials investigating several different treatment interventions. Thirteen studies with a follow-up period between 2 and 3 months were categorised as short-term assessment (18,21–26,29,31–33,35,37). Seven studies were reporting long term assessments (> 3 months), ranging between 4 and 12 months (19,22,23,31,33,36,37).

The total pool of treated participants was 623, with a wide age range from 43 to 89 years. All BMS participants were appropriately defined as having chronic pain for more than 3 months, with normal oral mucosa and absence of contributing local or systemic factors, except De Rivera Campillo et al. (19) (duration of BMS was less than 6 months), Cinar et al. (39) (average duration of BMS was 17 days), Ottaviani et al. (31) (duration of pain was 1 month), and Bergdahl et al. (37) (no description of BMS duration).

The visual analogue scale (VAS) or visual numerical scale (VNS) of either 0–10 or 0–100 scores were the primary assessment tools in measuring post-therapy pain improvement (18,20,21,23,24,27–29,31,33–35,38) except Bergdahl et al. (37) with a VAS scale of 1–7. Six studies used categorical changes in pain improvement as their assessment tool (22,23,25,26,32,33). Supplementary assessment tools such as the McGill Pain Questionnaire (21,23,35,36), faces scales (29), Orofacial Pain Clinic Questionnaire (EDOF-HC) (32) and Brief pain Inventory (BPI) (35) were used to evaluate pain intensity and associated characteristics further. Face scales classified patients’ expression of happiness based on a pictured face scale of 0–5 (lower is better). Secondary outcome assessment of participants’ quality of health, anxiety and depression, and quality of sleep were evaluated using patient-reported questionnaires, such as 36-Short Form Health Survey (SF-36), Oral Health on Quality of Life (OHIP 14), Patient Health Questionnaires-9 (PHQ-9), Patient Global Impression of Change (PGIC), Clinical Global Impression for global Improvement Scale (CGC-Z), Hospital Anxiety and Depression Scale (HADS), Beck Depression Inventory (BDI), Zerssen Mood Scale (ZMS), Hamilton Rating Scale (HRS), Psychometric
### Table 3. Reasons for studies’ exclusion.

| Author                        | Reason for exclusion                                                                 |
|-------------------------------|---------------------------------------------------------------------------------------|
| 1. Okayasu et al. 2020        | Non randomisation. No control. Follow up at 4 weeks                                   |
| 2. Paudel et al. 2020         | Non randomisation. Retrospective study. No control                                     |
| 3. Diep et al. 2019           | Non randomisation. Case series. No control                                            |
| 4. Bris et al. 2019           | Non randomisation. Case series                                                        |
| 5. Adamo et al. 2020          | Non randomisation. Unavailable post treatment result for control                      |
| 6. Jeong 2019                 | Follow up at 2 weeks                                                                  |
| 7. Iris et al. 2017           | Follow up at 4 weeks                                                                  |
| 8. Ilankizhai et al. 2016     | Review paper                                                                          |
| 9. Aravindhan et al. 2014     | Review paper                                                                          |
| 10. Miziara et al. 2015       | Review paper                                                                          |
| 11. Van Heerden WFP et al., 2011 | Review paper                                                                      |
| 12. Garg et al. 2017          | Non randomisation. No control. Case series                                            |
| 13. Jimson et al. 2015        | Review paper                                                                          |
| 14. Skrinjar et al. 2020      | Follow up at 2 weeks                                                                  |
| 15. Suga et al. 2019          | Non randomisation. No control                                                         |
| 16. Pereira et al. 2020       | Review paper                                                                          |
| 17. Nakase et al. 2004        | Non randomisation. Unavailable inclusion criteria on glossodynia. Follow up at 4 weeks |
| 18. Bessho et al. 1998        | Unclear definition on glossodynia. May included second burning mouth syndrome        |
| 19. Grechko et al. 1996       | Non randomisation. Study included second burning mouth syndrome                      |
| 20. Bardellini et al. 2019    | Follow up at 4 and 5 weeks                                                            |
| 21. Ritchie et al. 2018       | Review paper                                                                          |
| 22. Barbosa et al. 2018       | Follow up at 4 weeks                                                                  |
| 23. Sikora et al. 2018        | Follow up at 2 weeks                                                                  |
| 24. De Souza et al. 2018      | Systematic review paper                                                               |
| 25. Liu et al. 2018           | Systematic review paper                                                               |
| 26. Fenelon M et al., 2017    | Non randomisation. Retrospective study                                                |
| 27. Haggman-Henrikson et al. 2017 | Systematic review paper                                                            |
| 28. Kuten-Shorrer et al. 2017 | Non randomisation. No control                                                         |
| 29. Restivo et al. 2017       | Non randomisation. Case series. No control                                            |
| 30. Al-Maweri et al. 2017     | Systematic review paper                                                               |
| 31. Valenzuela et al. 2017    | Follow up at 2 and 4 weeks                                                            |
| 32. McMillan et al. 2016      | Systematic review paper                                                               |
| 33. Sugaya et al. 2016        | Follow up at 2 weeks                                                                  |
| 34. Cui et al. 2016           | Systematic review paper                                                               |
| 35. Valenzuela et al. 2016    | Follow up at 30 days                                                                  |
| 36. Kisely et al. 2016        | Systematic review paper                                                               |
| 37. Arduino et al. 2016       | Follow up at 21 days and 5 weeks                                                      |
| 38. Treldal et al. 2016       | Follow up at 2 weeks                                                                  |
| 39. Zakrzewska et al. 2016    | Systematic review paper                                                               |
| 40. Jurisic Kveisic et al. 2015 | Follow up at 4 weeks                                                               |
| 41. Lopez-Jornet et al. 2013  | Control arm included active study treatment                                           |
| 42. Komiyama et al. 2013      | No control group                                                                     |
| 43. Ko et al. 2012            | Non randomisation. No control. Follow up at 4 weeks                                   |
| 44. De Moraes et al. 2012     | Review paper                                                                          |
| 45. Silvestre et al. 2012     | Follow up at 1 week                                                                   |
| 46. Buchanan et al. 2010      | Review paper                                                                          |
| 47. Scardina et al. 2010      | Non randomisation                                                                     |
| 48. Kho et al. 2010           | Non randomisation. Follow up at 4 weeks                                               |
| 49. Lopez-Jornet et al. 2010  | Review paper                                                                          |
| 50. Gremeau-Richard et al. 2010 | Non comparable follow up time. Clonazepam follow up at 3 weeks and local anaesthesia at 15 min |
| 51. Barker et al. 2009        | Non control. Same group of drugs in comparison                                        |
Symptom Checklist-90-R (SCL-90-R), Medical Outcomes Survey (MOS) of Sleep Scale and Epworth Sleepiness Scale (ESS).

The substantial heterogeneity in the treatment methodology and regime, the follow-up time and inadequately reported statistical data precluded formal meta-analysis on the efficacy of a treatment in this review. However, combined SMD VAS scores or RR of studies with similar interventions were pooled with 95% CI. Two studies without comparison with placebo (30,39) and another, which described outcomes using median values (27), were qualitatively analysed.

Effects of treatment

The effectiveness of various treatments and pooled efficacy for similar treatments for BMS between short- and long-term outcomes are shown in Figures 2 to 5, respectively.

Anticonvulsants

Clonazepam. The efficacy of clonazepam in reducing BMS pain symptoms was reported in two studies with oral (20,39) and one with topical administration (19).

Short term (2 months). Treating BMS pain symptoms with daily oral systemic clonazepam 0.5 mg has shown favourable results of pain score reduction but was not statistically significant in the SMD analysis (SMD −0.63, 95% CI −1.56 to 0.29) (20). Despite the improvement in the taste, odour, and salivary flow rate, there were no statistically significant differences in improvement between clonazepam and placebo groups in taste (p = 0.83) and salivary flow (p = 0.03). Clonazepam did not improve patients’ ZMS mood and BDI depression scores.

Long term (4 months and 6 months). Administration of 2 mg clonazepam has been reported to reduce VAS score significantly at 4 months (MD −4.1, p < 0.001) (39). Eight of the 25 participants developed side-effects such as dizziness (n = 4), transient diarrhoea (n = 2) and myalgia (n = 2) with the use of clonazepam. Within the clonazepam group, 70% of patients described an improvement in pain intensity, and three participants were completely asymptomatic after 6 months of daily rinsing with 0.5 to 2.0 mg clonazepam (19). The application of topical clonazepam significantly decreased patients’ VAS score (MD −4.7) (SMD −1.06, 95% CI −1.58 to −0.54) in comparison to placebo than oral ingestion clonazepam (20) (MD −3.2) (SMD −0.63, 95% CI −1.56 to 0.29) and no significant difference in the total number of tablets dissolved in the mouth as a topical application between both clonazepam and placebo groups. Six months of clonazepam rinse statistically significantly reduced pain scores by 13-fold (RR 13.0, 95% CI 3.35–50.39).

Table 3. Continued.

| Author                        | Reason for exclusion                                      |
|-------------------------------|----------------------------------------------------------|
| Miziara et al. 2009           | Non comparative outcome assessment                        |
| Cavalcanti et al. 2009        | Follow up at 30 days                                      |
| Toida et al. 2009             | Included secondary burning mouth syndrome patients        |
| Buchanan et al. 2008          | Review paper                                              |
| Mingeuez Serra et al. 2007    | Review paper                                              |
| Patton et al. 2007            | Systematic review paper                                   |
| Buchanan et al. 2005          | Systematic review paper                                   |
| Zakrzewska et al. 2005        | Follow up at 2 weeks                                      |
| Gremeau-Richard et al. 2004   | Non randomisation. Follow up at 30 days                   |
| Petruzzi et al. 2004          | Non randomisation                                         |
| Femiano et al. 2004           | Systematic review paper                                   |
| Zakrzewska et al. 2003        | Review paper                                              |
| Scala et al. 2003             |                                                           |
| Femiano 2002                  | Unsure overlapping of recruited patient pools in Femiano 2000 trial or Femano and Scully 2002 trial |
| Maina et al. 2002             | Included secondary burning mouth syndrome patients        |
| Zakrzewska et al. 2001        | Systematic review paper                                   |
| Femiano et al. 2000           | Follow up at 1 month                                      |
| Sardella et al. 1999          | Follow up at 4 weeks                                      |
| Formaker et al. 1998          | Non randomisation. No definition on burning mouth syndrome |
| Grushka et al. 1998           | Non randomisation. No control                             |
| Dym et al. 2020               | Review paper                                              |
Five clonazepam participants reported sleepiness as adverse effect, but they were not suspended from the trial.

**Gabapentin**

*Short term (2 months).* Patients receiving 300 mg gabapentin has shown a similar result to alpha lipoic acid (ALA), with half of the total number of patients evidencing improvement in pain or total pain recovery (25). A more than three-fold likelihood of positive change relative to placebo were reported with the use of gabapentin in the short-term assessment of 20 BMS patients (25) (RR 3.33, 95% CI 1.58–7.02). It is associated with approximately a five-fold likelihood of decrease in pain levels compared with placebo if combined with ALA (RR 4.67, 95% CI 2.40–9.09) (25).

**Pregabalin**

*Long term (4 months).* At 4 months of assessment, 150 mg pregabalin showed a significant reduction in VAS scores (MD $-4.7, p < 0.001$) (39). Six of the 25 participants had side effects such as increase in appetite ($n = 3$), vertigo ($n = 1$), mild nausea ($n = 1$) and diarrhoea ($n = 1$).

**Antidepressants**

**Trazadone**

*Short term (2 months).* Administration of 100 mg trazodone daily for the first 4 days followed by 200 mg for 8 weeks significantly decreased patients’ VNS pain intensity against baseline (MD $-13.9, p < 0.01$), but there was no significant difference with the placebo group (SMD $-0.06, 95\%\ CI -0.72$ to 0.59; RR 0.95,
Figure 2. Forest plot showing standardised mean differences (SMD) and 95% confidence intervals for short-term outcomes (≥ 2 months and ≤ 3 months) of RCTs comparing an intervention with placebo for the treatment of BMS (with separate pooled effects for ALA).

Figure 3. Forest plot showing standardised mean differences (SMD) and 95% confidence intervals for long-term outcomes (≥ 3 months) of RCTs comparing an intervention with placebo for the treatment of BMS (with separate pooled effects for ALA).

Figure 4. Forest plot showing relative risks (RRs) and 95% confidence intervals for short-term outcomes (improvement on VAS at ≤ 3 months) of RCTs comparing an intervention with placebo for the treatment of BMS (with pooled effect for ALA).
95% CI 0.61–1.49) (21). If the assessment was based on the “Patients’ Global Assessment of Improvement” evaluation, trazodone and placebo groups reported improvements in pain intensity of 73% and 76%, respectively, and were not significant (p > 0.05). One patient in the trazodone group reported a worsening of symptoms. Both the trazodone and placebo groups significantly improved their BDI depression scores (p < 0.01). The most common side effects were dizziness and drowsiness, with seven patients dropping out due to dizziness. Other side effects included abdominal pains, headache, palpitation, tremor, xerostomia, and urinary incontinence.

**Citalopram**

**Short term (11 weeks).** The use of citalopram 10 mg daily followed by an increment to 20 mg after 1 week showed an improvement of VAS score of 87.45% (MD: −7.8, p < 0.001) (30). However, comparison with crocin reported no significant difference between their post treatment VAS scores (p = 0.98). The Hamilton questionnaires analysis revealed a significant reduction of depression and anxiety scores, with an average recovery percentage of improvement of 30.57% (SD 15.81) and 15.44% (SD 11.86), respectively. There was no significant difference in comparison between both groups in depression (citalopram: 19.4, SD 4.65; crocin: 19.0, SD 3.97, p = 0.76) or anxiety (citalopram: 18.6, SD 5.11; crocin: 18.0, SD 4.38, p = 0.76).

**Phytomedicine**

**Topical capsaicin**

**Short term (2 months).** Rinsing with 250 mg of chilli powder emulsified in 50 ml water with a dose concentration of 3.54 mg/ml capsaicin has been reported to induce a significant reduction in VAS score (MD −3.2, p < 0.01) with 76% of participants reporting an improvement in symptoms, but one patient-reporting a worsening (33). Capsaicin provides an immediate short term pain relief (SMD −1.49, 95% CI −2.35 to −0.63) and is statistically significant with 21 times better than placebo (RR 21.00, 95% CI 1.35 to 326.97). Topical capsaicin has shown a better clinical pain management outcome than oral ALA and lysozyme-lactoperoxidase, despite no statistically significant VAS difference in intergroup comparison.

**Long term (4 months).** Capsaicin showed superiority in maintaining VAS score reduction in long term (MD −2.9, p = 0.03) compared to lysozyme-lactoperoxidase, boric acid rinse and ALA (33). It also demonstrates sustainable benefit in long term administration (SMD −1.09, 95% CI −2.11 to −0.06) (33). It is 13 times better than placebo but not statistically significant (RR 13.00, 95% CI 0.84–201.27). An improvement in pain intensity was reported by 67% of participants, while one patient remained the same, reported worsening of pain. No adverse effect was noted during the trial.

**Ultramicronised palmitoylethanolamide (umPEA)**

**Short term (2 months) and long term (4 months).** Ottaviani et al. revealed a short-term (60 days) benefit with 1200 mg/day umPEA in BMS patients (SMD −0.70, 95% CI −1.39 to −0.01) but declining pain relief at 4 months (SMD −0.26, 95% CI −0.94 to 0.41) compared to placebo group (31). There were no side effects observed in patients treated with umPEA.

**Herbal catuama**

**Short term (3 months).** Catuama shows promising VNS (0–10) score reduction results compared to placebo with a minimal adverse effect of sleep alteration observed in the study (SMD −0.68, 95% CI −1.21 to −0.16) (29). Catuama shows a greater alleviation of patient symptoms with a lower faces scale score at both 8 and 12 weeks than placebo (p ≤ 0.001). The mean reduction of the face score were 1.6 and 1.5 for
8 and 12 weeks, respectively, while there were no changes in participants’ happiness in the control group with a similar mean reduction faces scale scores of 0.6 at 8 and 12 weeks. The majority of patients tolerated the treatment well, with none of the patients in the test group reporting xerostomia. The side effects reported by patients that took Catuama included somnolence and weight gain (n = 1), insomnia (n = 1), and exacerbation of the pain symptoms intensity in the first week of treatment (n = 2). A drop-out of eight (21.1%) participants in the treatment group, and four (11.8%) in the placebo group were reported.

Hypericum perforatum

Short term (3 months). At the end of 12 weeks of therapy, there was a reduction in the number of oral mucosa burning sites and improved ability to cope with the burning pain, there was no statistically significant difference with the placebo group (SMD −0.23, 95% CI −0.87 to 0.41) (28). The HAD questionnaires showed that approximately 50% of patients in both treatment and placebo groups evidenced better coping ability on their pain symptoms at the end of the trial. One participant developed a severe headache in the fifth week of active therapy (28).

Crocin

Short term (11 weeks). Crocin showed a significant reduction in VAS score (MD −7.8, p < 0.001) and has a similar improvement 87.5% of burning mouth score as citalopram (30). A significant improvement in depression and anxiety scores by 30.79% (SD 13.24) and 15.40% (SD 13.98), respectively, were reported. Crocin displayed similar effects as citalopram in treating burning pain, depression, and anxiety.

Lycopene enriched extra virgin oil (LVO)

Short term (3 months). A combination of topical spray and swallowing of 900 ppm LVO daily for 12 weeks led to a significant reduction in the median pain score (MD −3.0, p < 0.001) and burning (MD −1.0, p = 0.003) compared to baseline, but there was no significant difference (p = 0.99) when compared with the placebo group (27). Evaluation of SP-36 and OHIP-14 questionnaire scores showed no difference in changes to quality of life between treatment and placebo groups. HAD anxiety scores did not differ between treatment and placebo groups or significantly change throughout the trial period. The cholesterol and triglycerides levels were not remarkably raised after 12 weeks of LVO administration.

Alpha lipoic acid (ALA)

Short term (2 months). Four ALA trials (22,25,26,33) showed promising pain reduction in comparison to placebo during short term assessment (Femiano and Scully: RR 2.42, 95% CI 1.55 to 3.77; Lopez D’alessandro: RR 3.67, 95% CI 1.78 to 7.54; Palacios-Sanchez: RR 2.32, 95% CI 1.20 to 4.48; Marino: RR 17.0, 95% CI 1.08 to 268.86) while two did not (Carbone: SMD −0.06, 95% CI −0.75 to 0.64; RR 0.95, 95% CI 0.33 to 2.76; Lopez Jornet: SMD 0.56, 95% CI −0.10 to 1.22) (23,24). The pooled ALA suggested a more than double increase in likelihood of pain improvement (RR 2.44, 95% CI 1.57 to 3.78, p < 0.001) compared to placebo (22,23,25,26,33). However, there were no significant changes in the pooled ALA VAS scores (SMD −0.17, 95% CI −1.08 to 0.75, t −0.36, p = 0.72), reflecting the heterogeneity across studies (23,24,33). One patient had to discontinue treatment during the trial due to gastrointestinal upset such as nausea, dyspepsia and pyrosis (24).

Long term (4 months and 12 months). Two studies (23,33) assessed the persistence of the observed improvement for 2 months after discontinuation of therapy and described a stable decrease of VAS score (Carbone: MD −1.8, SD 3.19, p = 0.01; Marino: MD −1.8, p > 0.05). Long-term use of ALA did not result in any statistically significant improvement over placebo, suggested by the pooled VAS mean score changes (SMD −0.40, 95% CI −0.95 to 0.15, p = 0.15) (23,33) and the likelihood of improvement (RR 3.66, 95% CI 0.55–24.45, p = 0.18) (22,23,33). A study comparing ALA 600 mg with two other drugs (clonazepam and pregabalin) showed no significant improvement at 4 months of assessment (MD −0.72, p > 0.05). Three out of 25 patients reported side effects, including mild nausea (n = 2) and myalgia (n = 1) (39). A 1-year follow-up showed a sustained effect on pain intensity in 73% of patients. In this study, patients with signs of improvement within the first 4 months of treatment were given an extended treatment of 1 month ALA 600 mg (22).

ALA and gabapentin

Short term (2 months). A combination of 600 mg ALA and 300 mg gabapentin in a randomised, double-blind clinical trial described a notable pain reduction, with 70% of patients demonstrating a partial or complete improvement in pain intensity compared to 15% in the placebo group (25). The combined use of ALA and gabapentin gave a five-fold likelihood (RR 4.67, 95% CI 2.40–9.09) (p < 0.001) of decrease pain intensity while ALA only
has four times the likelihood of beneficial effect (RR 3.67, 95% CI 1.78 to 7.54).

**ALA and vitamins**

*Short term (2 months) and long term (4 months).* Combining vitamins such as vitamin C, PP, E, B6, 2,1, 12 and folic acid with 800 mg ALA did significantly improve VAS score (MD −1.0, SD 1.83, p = 0.047) and a further reduction in VAS score was noted 2 months after termination of treatment (MD −1.8, SD 3.19, p = 0.047) (23). However, there was no significant difference between ALA and vitamins (SMD 0.21, 95% CI 0.44–0.85) (SMD −0.15, 95% CI −0.79 to 0.50) compared to ALA monotherapy (SMD −0.06, 95% CI −0.75 to 0.64) (SMD −0.23, 95% CI −0.93 to 0.47) or placebo in both short (p = 0.60) and long-term assessment (0.79). ALA as a monotherapy led to a higher reduction in VAS score at 2 months (MD −1.6, p = 0.013) but no statistically significant difference compared to placebo (p = 0.60) compared to baseline, but there was no significant difference between the ALA (monotherapy), ALA and vitamin (combination) and placebo groups. No adverse effects were reported in the study (23).

**Melatonin**

*Short term (2 months).* A cross-over clinical trial involving intervention with a high melatonin dosage (12 mg/day) did not provide pain relief (SMD 0.24, 95% CI −0.39 to 0.87; RR 1.18, 95% CI 0.31–4.43) and sleep score improvement compared to placebo (18). Ten participants reported no changes in symptoms, and one participant reported worsening of symptoms. The value of VAS score and serum plasma melatonin concentration was negatively associated, but it was not statistically significant (p > 0.05). Two patients in the melatonin group demonstrated a positive correlation between decreased VAS scores and increased sleep hours. The Hamilton rating scale for anxiety (HAM) assessment scores was always higher in the melatonin group than placebo, with a statistically significant decrease in the melatonin group’s anxiety score (p < 0.05). An approximate two-fold of patients reported sleep impairment using melatonin (n = 10, 62.5%) compared to placebo (n = 6, 37.5%). Mild daytime sleepiness was seen in melatonin and placebo groups, with high ESS scores but not significant between them (p > 0.05). The main adverse effect of melatonin that leads to the discontinuation of treatment on four patients were heavy tremor, sexual disturbances, blurred vision, and severe heavy headedness. Four patients were dropped from the study due to lack of efficacy, pain improvement, and follow-up loss.

**Low-level laser therapy (LLLT)**

*Short term (11 weeks).* A significant reduction in pain score by three to five units was observed in the study using the red (p = 0.13) and infrared laser (IR1W p = 0.004 and IR3W p < 0.001) (34). The red laser group (SMD −0.47, 95% CI −1.13 to 0.18) did not demonstrate a significant difference from the control group, but both IR1W (SMD −0.80, 95% CI −1.46 to −0.14) and IR3W (SMD −1.14, 95% CI −1.83 to −0.45) showed a statistically significant difference from the control group (34). No side effects were noted from the laser therapy.

*Long term (4 months).* A recent trial has suggested the advantage of photobiomodulation in treating orofacial neuropathic pain, including BMS with a significant 4.5-fold likelihood of pain reduction in comparison to placebo (RR 4.50, 95% CI 1.28–15.81) and a more than 1-point decrease in VAS (SMD −1.12, 95% CI −2.10 to −0.15) (36), but no improvement in patients’ psychology and quality of life. There was no significant improvement in McGill Pain scores, patient oral health quality scores (OHP), physical and emotional scores (SF-36) and sleepiness (ESS). However, there was a significant decrease in SCL-90-R interpersonal sensitivity, somatisation, and anxiety between the photobiomodulation group and the placebo group (p = 0.04). No adverse effects were reported.

**Saliva substitutes**

*Topical lysozyme lactoperoxidase (Biotene)*

*Short term (2 months) and long term (4 months).* Lysozyme lactoperoxidase (Biotene) rinse was prescribed to BMS patients diagnosed with xerostomia (33) and reported a decrease in pain score of 1.7 units during short-term assessment (SMD −0.93, 95% CI −1.72 to −0.13) but no advantage over placebo was seen in long-term assessment (SMD −0.73, 95% CI −1.72 to 0.26). A 13-fold (RR 13.00, 95% CI 0.80–210.82) and nine-fold (RR 9.00, 95% CI 0.55–146.12) likelihood of pain reduction compared with placebo was observed in both short- and long-term analyses (33).

The lubricating rinse lysozyme lactoperoxidase significantly reduced the VAS score (MD −1.7, p = 0.01), but there was no significant difference between lysozyme lactoperoxidase with capsaicin rinse and oral ALA, respectively (33). The pain score remained unchanged in 57% and 55% of patients in both short and long-term assessment.

*Topical urea*

*Short term (3 months).* Statistical analysis showed no statistically significant difference between the
application of 10% urea for 3 months and the placebo group ($p = 0.34$) (RR $0.95$, 95% CI $0.50$–$1.80$) (32). There is no difference in pain intensity after treatment ($p = 0.88$), although clinically 58.3% of patients demonstrated a reduction in pain intensity.

**Transcranial magnetic stimulation (rTMS)**

**Short term (2 months).** Ten days of 30,000 pulses of rTMS therapy over the left GDLPMC significant reduced VAS score (MD: $-3.1$, $p = 0.002$) with 75% of patients reporting a decrease in pain intensity of more than 50% compared to baseline (35). There was a significant difference compared with placebo (MD: $-2.8$, $p = 0.005$) (SMD $-0.33$, 95% CI $-1.25$ to $0.60$). There was a significant improvement in sensory SFMPQ in the rTMS group (MD $-4.84$, $p = 0.002$) but no difference in the SFMPQ affective scores and present pain intensity. PGIC and CGO-I assessments described positive changes from the patient in the rTMS group. There were no significant changes in patient mood based on PHQ-9 (MD $5.59$, $p = 1.00$).

**Tongue protector**

**Short term (2 months).** The hypothesis of wearing the tongue protector to prevent continuous irritation of tongue on teeth or denture has a statistically significant difference in improvement in VAS score between wearer (MD $-3.6$) and non-wearer with habitual avoidance reminder (MD $-1.4$, $p < 0.001$; SMD $-1.15$, 95% CI $-1.76$ to $-0.54$) (38). Participants did not show any improvement in the depression and anxiety score. There was a significant improvement in patient quality of life based on OHIP-49 and SF36 assessments.

**Cognitive therapy**

**Short term (12–15 weeks) and long term (6 months).** At the end of weekly behavioural therapy for 12–15 weeks, patients reported a significant improvement in their pain score for both short- (SMD $-2.16$, 95% CI $-3.09$ to $-1.24$) and the long-term effects were sustained over 6 months post-treatment: (SMD $-3.38$, 95% CI $-4.53$ to $-2.23$) (37). There were statistically significant changes between the therapy and the placebo group ($p < 0.001$).

**Discussion**

At present, there is no definitive curable treatment for BMS. Its aetiology remains uncertain with various suggested pathogenesis such as peripheral and central neuropathy disorders, psychological disorders, changes in gonadal, adrenal and neurosteroid levels, a dopamine D2 receptor (DRD2) $957C>T$ genotype and the association between BMS and other neurological diseases such as Parkinson’s disease (40–43). BMS treatment primarily aims at eliminating the painful burning dysaesthesia. Phenotyping BMS patients’ aetiology could achieve this based on their clinical histories and responses toward various treatments. In this review, we discuss nine BMS therapies: Anticonvulsants (19,20,25,39), antidepressants (21,30), phytomedicines and food supplements (18,22–29,31–33), lower-level laser therapy (34,36), saliva substitute (32,33), transcranial magnetic stimulation (35), oral appliances (38) and cognitive behavioural therapy (37).

Preceding systematic reviews included clinical trials of 2 weeks follow-up assessment results. It is crucial to have a more extended review period of patients’ responses towards the therapy, the sustainability of the treatment effects and the possible side effects before considering that a treatment has been effective. Hence, to ensure sufficient, sustainable benefits of the treatments, this review includes studies with a minimum follow up of 2 months and divided them into short term ($\leq 3$ months) and long term ($> 3$ months) treatments (11).

The majority of the included studies had small sample sizes. The diversified BMS patients’ characteristics such as presence or absence of psychological disorders, taste disturbance, and xerostomia make recruitment for a larger homogenous sample group difficult in a clinical trial. The concurrent use of psychotherapeutic drugs or therapies and anti-inflammatory analgesic medications in patients may influence the presentation of the BMS population trials due to the ambiguity of whether these psychological disorders preceded BMS (21,26,32,35,38).

**Anticonvulsant**

**Clonazepam.** Both oral ingestion and topical application of clonazepam have showed a favourable result on BMS pain relief up from 2–6 months (19,20,39). The association of the peripheral or central nervous system in BMS pathogenesis explained the use of anti-epileptic and antidepressant drugs. Continuous nociceptive peripheral neuropathy input will eventually lead to central sensitisation and changes. Pharmacological drugs such as clonazepam demonstrated their analgesic ability by inhibiting neurological transduction and transmitting the pain signal. Clonazepam, a benzodiazepine anticonvulsant drug, acts as an agonist modulator on GABA-A receptors and activates the descending pain inhibitory pathway of the peripheral (PNS) and central nervous system (CNS) by facilitating the opening of the chloride channel. It antagonises the neuron hyperexcitability
transmission by generating a continuous hyperpolarisation, thus preventing depolarisation and post deafferentation neuronal firings (44). GABA-A receptors are found in the oral mucosa, mandible, palate, salivary gland, and taste pathway. GABA agonist could reverse the dysfunction of peripheral chorda tympani nerve and taste loss in BMS patients (45). Clonazepam could provide fast and continuous pain relief due to its rapid absorption and 90% bioavailability of clonazepam within 1–4 h after oral administration and its long half-life of 30–40 h.

Meanwhile, intraoral topical clonazepam has shown to be superior to oral ingestion in providing much rapid pain analgesia but a shorter duration of action. Patients reported rapid positive effects within 10 min upon dissolving the clonazepam tablet intraorally and recurrence of pain in 3–4 hours (19). The topical clonazepam route is simple with a rapid and shorter duration of action, which allows repetitive use and lower risk of common systemic adverse effects such as drowsiness, dizziness, and unsteadiness. It allows patients to have better self-control over pain relief magnitude in their daily activities. Inevitably, some of the topical clonazepam will be absorbed systemically through the oral mucosa and affect the CNS pain modulation. This is reported in a study assessing patient’s post topical clonazepam serum concentration, which was similar between 5 h post sucking a 1 mg clonazepam tablet and after sucking the tablets three times daily for 14 days (46).

The use of amitriptyline, a tricyclic antidepressant, commonly used to treat chronic neuropathic pain, has not been widely mentioned in BMS studies. This may be the result of the frequent xerostomia induced by amitriptyline, which aggravates the pre-existing BMS-related xerostomia. A retrospective study has reported a superior rapid decrease of VAS pain scores outcome for clonazepam drops (n = 23) compared to amitriptyline drops (n = 16) at 6 weeks but no statistical difference between them (47).

Gabapentin and pregabalin. Gabapentin and pregabalin have been the favourable drug choice in treating neuropathic pain conditions such as diabetic neuropathy and postherpetic neuralgia due to its hepatic safety profile (48). Similar advantages in BMS pain were achieved with the use of gabapentin and pregabalin in short- and long-term assessment (25, 39). Gabapentin mediates pain attenuation by binding to the z2δ-1 subunit of the voltage calcium channels and inhibits the release of neurotransmitters such as glutamate, CGRP, and substance P; the development of chronic pain (49, 50) correlates BMS as a neuropathic pain that may involve both central and peripheral mechanisms.

The benefits of gabapentin in BMS with peripheral neuropathy disorders may suggest using adjunct dietary supplements such as ALA to enhance the pain attenuation without increasing the synthetic drug’s needs. However, a more extensive sample size study is recommended to test the efficacy of gabapentin and its adverse effects. Cinar et al. compared the use of systemic pregabalin (150 mg) with clonazepam (2 mg), and both drugs show similar significant efficacy in reducing pain score (39). A third of patients in both study groups had common adverse effects, but no patients withdrew from the study. The absence of a placebo group in the study failed to give a definitive superiority outcome between pregabalin and clonazepam (39).

Antidepressants

BMS has been strongly associated with depression and anxiety, and the lack of clarity between them is unsettling. This neurophysiological mechanism in BMS was shown in functional magnetic resonance imaging (fMRI) (51) and quantitative somatosensory testing (QST) study (52). fMRI study has reported an increase in the region’s functional neural activity regulating depression and anxiety in BMS patients (51). It is known that chronic anxiety and depression may disturb neuroprotective steroid productions (53). As pain could be a somatic trait, the use of an antidepressant has suggested the role of anxiety and depression in BMS pathogenesis.

Trazodone. Trazodone is a second-generation antidepressant that has been considered a multifunctional drug and acts as a serotonin reuptake inhibitor. Trazodone has been used in treating anxiety and pain symptoms, including fibromyalgia (54). However, in this review, trazodone use did not significantly affect pain reduction and had a high placebo effect. The reported high adverse effects on dizziness and drowsiness limit its use (21).

Citalopram. Citalopram has shown to be able to reduce pain intensity (30). A review of SSRIs such as zimelidine, sertraline, citalopram, paroxetine, and fluoxetine has suggested it for the treatment of chronic pain conditions (55). The SSRI citalopram has similar antidepressant and analgesic properties to tricyclic antidepressants but with significantly fewer side effects and better tolerability (56). Serotonin is a neurotransmitter that plays a role in both central and peripheral nociception and mood regulation. SSRIs inhibit serotonin’s reuptake and prolong its availability in the synaptic cleft. There was inconclusive effectiveness in treating chronic pain with SSRIs. Inconclusive results were observed from various studies on its use for chronic somatoform pain and fibromyalgia. As there
is no placebo group in comparing the efficacy of citalopram in reducing burning mouth and less than 50% of patients recovered from depression and anxiety, there is limited evidence to support its use (30). Clinical trials with better methodology and low risk bias are needed to conclude the effect of SSRI as a treatment for chronic pain conditions.

**Phytomedicine**

The perspective of using herbal medicine or phytomedicines has been established and increased in primary health care (57). The efficacy of phytomedicines such as capsaicin, herbal catuama, umPEA and hypericum perforatum have demonstrated their analgesia ability, with capsaicin having a tremendous number of patients in responding to it. Through well-designed randomised control trials and observational studies, phytomedicine has a tremendous future to be used solely or as adjunct therapy in treatment therapeutic strategies and products (58).

**Capsaicin.** Capsaicin has shown to be an effective pain desensitiser especially with oral topical application for up to 4 months (33). Transient receptor potential vanilloid-1 receptors (TRPV1) are found in the PNS and CNS (59). The numbers of TRPV1 receptors are significantly increased in the mucosa of BMS patients' tongues (60). Activation of TRPV1 at the peripheral terminal fibre endings leads to the release of neuropeptides such as substance P, neurokinin A (NKA) and calcitonin-gene-related peptide (CGRP), which contributes to the onset of hyperalgesia pain and inflammation. Local capsaicin application activates the TRPV1 and modulates the nociceptive transmission of pain impulses from the peripheral stimulation site to the central nervous system by blocking axonal transportation, depleting neuropeptides, and loss of membrane action potential. Hence, capsaicin-induced analgesic effect by desensitisation of the nociceptive fibre (61–62), which is a reversible process (63). The used of topical capsaicin have been suggested in neuropathic pain, such as postherpetic neuralgia and painful HIV associated polyneuropathy (64–66) but not inflammatory pain such as osteoarthritis (67).

A study showed no difference between systemic and topical capsaicin efficacy in BMS (68). However, gastric pain limits systemic capsaicin use (68). The use of topical capsaicin rinse is recommended in BMS due to its rapid action and there being no reported adverse effects, as seen in other synthetic drugs. However, there are no known risks of long-term repeated rinsing of capsaicin, especially in the oral cavity mucosa innervation. Patients should be warned of the initial increase in burning pain induced by topical capsaicin rinse or application followed by the discharge in the C and Aδ nociceptive fibres, but this effect is limited, of short duration, and followed by pain relief. Cutaneous site pre-treatment with anaesthetic cream has been used clinically to reduce the capsicain patch-induced treatment discomfort in patients with peripheral neuropathic pain (69). Hence, a possible hypothetical proposition of a mouth rinse mixture containing both capsaicin and lidocaine may mask this initial burning pain and enhance pain relief effectiveness.

**Ultramicronised palmitoylethanolamide (umPEA).** There is a small reduction of pain score with umPEA but its effect did not sustain (31). Systemic administration of PEA elicits anti-inflammatory, antinociceptive, and neuroprotective effects, both in vivo and in vitro (70,71), as well as in human subjects (72,73). Neurodegeneration could occur due to inflammatory reactions and activation of immune cells. Microglia facilitates the CNS’s inflammatory response, and white mast cells coordinate PNS inflammation. umPEA is an endogenous fatty acid that suppresses the discharge of proinflammatory mediators from mast cells and microglia during inflammation, thus preventing neuronal injury and chronic pain. A meta-analysis study has reported umPEA as a novel treatment in managing chronic neuropathic pain caused by neuroinflammation (74). A study of 40 days umPEA has reported positive benefit in diabetic or traumatic peripheral neuropathic pain (75). The novelty of umPEA efficacy as a primary or adjunct treatment in BMS should be further studied with a larger cohort and follow-up period for its sustainability.

**Herbal catuama.** Three months used of catuama has shown a significant reduction in BMS pain score (29). Catuama is a herb commonly used for mental and physical exhaustion. It has been shown to have antidepressant, antinociceptive and vasorelaxant actions in animal models by acting on the dopaminergic, serotonergic, and opioid pathways and reducing the inflammatory nociception in animal models (76). It is thought that catuama may alleviate the burning pain based on the possible BMS aetiologies of psychologic and neuropathic disorders. A more extended observation on the use of catuama is suggested to ensure its long-term adverse effects and suitability as a pain relief.

**Hypericum perforatum.** The short-term use of hypericum perforatum in BMS has shown a favourable outcome but not significantly better than placebo (28). Hypericum perforatum (St. John’s Wort extracts) has been used as an antidepressant in mild to moderate depression, anxiety and sleep disorders (77) and may be beneficial to BMS patients as they frequently...
experience emotional and mood distress, in which anxiety and depression could be the primary or secondary event. Several active extracts in hypericum perforatum have a strong affinity for γ-aminobutyric acid (GABA), adenosine, serotonin 5HT1, as well as benzodiazepine receptors, and act as monoamine oxidase inhibitors (MAOIs) (78). Its action as a MAOI prevents the reuptake of norepinephrine, serotonin and dopamine neurotransmitters from the brain, providing beneficial antidepressant effects. As a GABA agonist, it induces a temporary hyperpolarisation of the neuronal membrane and ensuing desensitisation and inhibition of neurotransmission, which provides an anxiolytic and analgesic effect (79).

Hypericum perforatum rarely causes any adverse drug reactions, except for dizziness, and is usually well tolerated by the elderly (80). It has comparable efficacy and safety compared to SSRIs in patients with mild to moderate depression (81). However, there is inadequate evidence on its long-term efficacy and safety, especially in patients with severe depression or suicidal risk.

Although it is relatively safe, clinicians should be wary of prescribing hypericum perforatum with other medications as it may elicit severe clinical adverse drug interaction effects. Hypericum perforatum activates the cytochrome P450 enzymes involved in drug metabolism, and reduces the plasma concentration and potency of a number of drugs such as warfarin (risk of thrombosis), cyclosporin (risk of transplant rejection), oral contraceptives (unintended pregnancy), anticonvulsant (uncontrolled seizures), digoxin (cardiac arrhythmia), theophylline (poor asthmatic control), and HIV protease inhibitors and non-nucleoside reverse transcriptase inhibitors (diminution in HIV suppression) (82). Caution should also be taken in combining hypericum perforatum with medications that have serotoninergic effects as it increases the serotoninergic action of serotonin receptor agonists (triptans) as well as of selective serotonin reuptake inhibitors (SSRIs), selective norepinephrine reuptake inhibitors (SNRIs), tricyclic antidepressants (TCAs) and monoamine oxidase inhibitors (MAOIs) (82,83).

Crocin. Crocin is a carotenoid chemical compound found in the flowers crocus and gardenia and is responsible for the colour of saffron. Crocin prevents neuroinflammation and neurodegeneration by decreasing oxidative stress and cell death (84) by inhibiting microglial activation and suppressing inflammatory cytokine production (85). Microglia dysfunction contributes to the disturbance in their protective regulator function on neuroinflammation stimuli and generates an imbalance of reactive oxygen species (ROS) homeostasis and the antioxidant system, creating oxidative stress (86,87). Oxidative stress is associated with neurodegeneration through several cascades of deleterious events on the cells, causing lipid peroxidation, protein oxidation and mitochondrial DNA damage, and mutations (88). The accumulated increased oxidative stress in the aged brain has been thought to be a possible aetiology of neurodegenerative diseases such as Parkinson’s disease and Alzheimer’s disease. There have been reports on BMS occurrence in a patient with Parkinson’s disease (89,90), but there is no study on dysfunction of microglia and mitochondria and the oxidative stress in BMS patients. The brain is much more vulnerable to this oxidative stress due to its high oxygen demand and lipids’ vital role in maintaining neuronal function (91).

Neuroprotective effects of crocin have been shown in an experimental animal model (84), but not in more extensive human clinical trials on its long-term safety and benefits. This review shows a significant improvement in crocin pain score but no significant superiority over citalopram (30). A three-arm study design with placebo control group comparison is advised to compare crocin and citalopram’s superiority.

**Lycopene and virgin olive oil (VOO).** Lycopene is naturally found in red carotenoid pigmented food, such as in tomatoes. It has antioxidant, anti-inflammatory, and anti-apoptotic properties. These benefits have been seen in reducing cancer and cardiovascular risk with the consumption of lycopene and VOO (92,93). Combination of lycopene and VOO is thought to provide a synergistic effect of antioxidative and anti-inflammatory mechanisms. The ingestion of lycopene with olive oil will increase bioavailability (94). The application of topical lycopene and VOO may protect the oral mucosa’s peripheral neurons from oxidative stress, while VOO provides a lubricant effect. However, lycopene and VOO are not superior to placebo in improving pain score and health quality (27).

**Alpha lipoic acid (ALA)**

ALA is the most studied treatment in BMS. Although the VAS findings from the pooled ALA analysis suggested there was no significant reduction in pain intensity relative to placebo treatment, a significantly higher proportion of patients reported pain reduction with ALA. As such, it suggests ALA as a treatment for BMS, but the evidence is not conclusive due to the variability of the studies treatment regimens and short- and long-term study results (9–11).

ALA is a naturally occurring compound found in the body and vegetables such as tomatoes, potatoes, broccoli, and brussels sprouts. It acts as an enzymatic cofactor for pyruvate dehydrogenase and α-ketoglutarate dehydrogenase complexes in glucose and lipid
metabolism. ALA is a robust universal antioxidant and can chelate and remove heavy metals from the body. Thus, it reduces oxidative stress-induced inflammation and damage to the nerve. ALA’s advantages and safety were demonstrated in the treatment of diabetic polyneuropathy pain and paraesthesia by preventing nerve fibre degeneration (95,96). Hence, the possible goal of administering ALA in BMS patients is to treat patients with peripheral neuropathy as the pathogenesis. The bioavailability of oral ALA is strongly affected by its formulation and its regime due to its reduced solubility and stomach instability. ALA in liquid form is preferred over solid for better absorption and should be taken pre-meal. Age influences the bioavailability of ALA. Patients aged above 75 years have better absorption rates than 18 and 45 years, but there was no difference in gender (97). As BMS commonly occurs in the fifth to seventh decade of age, ALA may be a beneficial adjunct supplement to ease the pain. In this review, the mean age reported ranged between 45 to 67 years.

ALA and gabapentin have shown a superior result, with mild adverse effects reported (25). Combined ALA use as an adjunct supplement to pharmacotrophic drugs may benefit the patients in minimising the drug’s adverse effects by reducing the prescribed frequency and dosage. However, studies with larger sample sizes and longer follow-ups of a minimum of 6 months with better methodology design should be conducted to validate the use of ALA.

**Melatonin**

There was insufficient evidence of the benefit of melatonin in BMS. The relationship between pain and sleep is inextricable, in which poor sleep quality is a risk factor for chronic pain development, and pain disrupts the sleep pattern (98). Melatonin is a neurohormone that regulates the circadian biological rhythms. Melatonin has antioxidant, anti-inflammatory, anticancer, anxiolytic and antinociceptive activities (99). It has been shown to reduce chronic pain in fibromyalgia (100) and temporomandibular joint disorders (101). The analgesic effect of melatonin in neuropathic pain has been demonstrated in animal models (102,103). The use of exogenous melatonin in neuropathic pain is controversial due to multiple complex analgesic mechanistic pathways (104). A notable 40% drop-out rate was seen using melatonin due to heavy tremor, sexual disturbances, blurred vision, and headache (18), despite the claim that melatonin is well tolerated and safe at high doses (105). As sleep disturbances are uncommon in BMS patients, this may in part explain the poor treatment response of BMS-related pain to melatonin.

**Low-level laser therapy**

Photobiomodulation with low-level laser therapy (LLLT) effectively reduces chronic pain such as low back pain, temporomandibular joint disorder, and osteoarthritis (106). LLLT facilitate analgesia via its anti-inflammatory effects by increasing the secretion of serotonin, endorphins and adenosine triphosphate, augmentation of the cell membrane potential and suppressing impulse conduction velocity (107). The infrared laser has a longer wavelength compared to the red laser. It will penetrate tissue deeper, reaching the nerve fibres (108). This is observed in Sapanberg et al., where the infrared laser has a higher, significant difference in the reduction of pain score compared to placebo, but the red laser showed no difference to the control group (34). Increasing the intensity of the laser therapy application has remarkably augmented the significance of pain score improvement compared to placebo as seen in IRW3 with three sessions per week compared to IRW1 with one session a week. In summary, LLLT seems to be able to contribute to BMS patients’ pain relief and the possibility of being used along with pharmacological and psychological treatments for a better outcome. The beneficial effect of LLLT is sustained from 1–4 months after 10 sessions of LLLT (36). It is suitable for use in medically compromised or patients on polymedication for pain as it is a non-invasive technique with no known reported adverse effects.

**Saliva substitute – Biotene and urea**

BMS patients often complain of dry mouth discomfort (109). The lower salivary flow rate and thicker saliva froth may disturb the taste function (110). Urea and lysozyme lactoperoxidase (Biotene) are topical anti-xerostomic medication (saliva replacement). De Silva et al. studied urea as an adjunct therapy in BMS patients who were concurrently treated with amitriptyline (32). Amitriptyline is the first line of drug used in treating chronic neuropathic pain (111) and is known to cause dry mouth. There was no beneficial improvement seen in burning pain, taste and somatosensory despite increased oral cavity moisture and lubrication with urea or Biotene. BMS patients have decreased unstimulated salivary flow rate but not stimulated saliva. There was no objective hyposalivation observed, which explain the lack of oral cavity lubricants efficacy in reducing the pain intensity (110,112) and the possibility of central neuropathy as the pathogenesis. Caution should be taken due to the small participant size of less than 20 in both studies (32,33).

Anecdotal patient claims suggest regular sips of ice water help elevate the pain, which may be due to
stimulation of transient receptor potential melastatin 8 (TRPM8) cold receptors or antagonist effect on TRPV1 found in the oral mucosa. The role of TRPM8 in pain analgesia has been widely contradictory debated, which may depend on its anatomical site and degree of activation (113).

**Transcranial magnetic stimulation (rTMS)**

Neuroimaging studies have demonstrated BMS patients to have similar brain pain matrix changes with increased functional connectivity and reduced grey matter volume as seen in other chronic pain imaging studies, indicating dysfunction of pain regulation at the CNS level (51,114). It has been established that unilateral stimulation of primary motor cortex (M1) and dorsal lateral prefrontal cortex (DLPFC) with rTMS generates a diffuse analgesic effect in both experimental and clinical pain studies (115,116). The extent rTMS induced analgesic effects depend on the stimulation patterns such as the frequency and magnitudes and coil position. A single stimulation session could provide several days of analgesia, and this effect is reinforced with echoing rTMS sessions (116). This was demonstrated in Umezaki et al. with a rapid decrease in VAS scores at day 8 and 15 of rTMS treatment and a stable pain reduction score for 2 months (35). However, a peculiar finding of a temporary increase of pain score on day 30 followed by a reduction in pain score on day 60 was explained by the author as possible psycho-pathophysiological disease differences (perception of pain and duration of diseases) of each patient. Further statistical analysis shows a lack of significant improvement in the mean pain score difference for short-term rTMS used (35). rTMS is a non-invasive neuromodulation technique that could be a novel treatment in chronic pain either solely or as a complement to medication and could be useful in refractory cases. However, standardisation of therapy protocol should be established in experimental animal models before its clinical implication.

**Oral appliance (tongue protector)**

A tongue protector has been shown to reduce discomfort and improve oral health and quality of life (38). BMS often presents in the anterior two-thirds of the tongue, dorsal and lateral surfaces of the tongue, anterior hard palate, lip mucosa and gingiva (4). It was thought that parafunctional habits such as tongue thrust or continuous habitual rubbing over the teeth or denture and lip, cheek, or tongue biting contribute to BMS pain (117), but this contradicts the definition of BMS (1). It is hypothesised that chronic hyperactivity of trigeminal nociceptive pathways will produce intense pain response and occurrence, or a burning mouth feeling. The use of a tongue protector may avoid other triggering factors such as dietary stimulants (hot and spicy food, citrus food) or accidental tongue irritation on the pain site. It may create a self-false psychological security belief that the appliance protects the tongue.

**Cognitive therapy**

Bergdahl et al. reported an impressive reduction of three units of pain scores for both short and long-term assessment (37). The study has clearly defined its BMS patients as similar to the current ICOP recommendation (1), despite being an early years study and proven CBT benefits (37). BMS has frequently been associated with psychological disorders such as depression, anxiety, hypochondriasis and cancerphobia (4). It remains unclear whether anxiety and depression precede BMS or if they are a consequence of chronic pain. Treatment-resistant patients may have a contributing psychological factor. Cognitive behavioural therapy (CBT) is a common psychotherapeutic intervention for patients with chronic pain, and its effectiveness is influenced by the level of empathy received by the patient. Interestingly, females have commonly better outcomes than males. CBT improves the patient’s quality of life by allowing them to perform their daily activities without limitation and diverts their concentration on the pain, changing the thought and coping adaptive behaviours (118,119). A combination of psychopharmacological treatment may help the patient avoid the possibility of drug abuse and adverse effects. However, a larger sample size should be obtained to establish the benefit of CBT and to rule out the attention placebo effect, as the patient was reviewed more frequently.

In summary, the statistical analysis on the RCTs comparing intervention with placebo suggests a strong favourable outcome (SMD > 1.000) for cognitive behavioural therapy, capsaicin, topical clonazepam, and laser therapy (highest to lowest) in both short- and long-term assessment. There was some evidence on the use of phytomedicines such as umPEA, herbal catuama and hypericum perforatum in short-term pain score reduction. There were negligible changes in short term pain improvement in both trazodone and ALA (pooled effects) studies. However, the positive effects of ALA increase in long-term assessment. Although the pooled effect of ALA pain score improvement is low, the number of patients responding to ALA and its combination with gabapentin or a vitamin were high in both short- and long-term assessments. Capsaicin, topical clonazepam and saliva substitute lysozyme lactoperoxidase showed consistent treatment effectiveness or improvement in pain comparing with placebo in both short- and long-term analysis.
Acupuncture

There is emerging interest in acupuncture as an adjunct therapy to pharmacological treatment for BMS patients due to its encouraging analgesic results on significant VAS score reduction within the first 2 months of therapy (120–125). Long-term follow-up, between 18 and 24 months after the initial acupuncture treatment, suggests a decreased level of burning sensation and improved quality of life are maintained (122,125). Scardina et al. proposed that acupuncture increases BMS patients’ lip microcirculation, which in turn reduces the localised collection of inflammatory mediators, hence providing respite from the burning pain (125). Acupuncture was not included in this review as, disappointingly, studies of this treatment to date have either been non-randomised clinical trials recruiting cohorts of consecutive BMS patients, lacked a control group, and/or administered follow up less than 2 months post-treatment. A further detailed study on the potential of acupuncture as a complementary therapy to reduce medication loading and increase patient compliance with medications is warranted.

Limitations

There was a substantial amount of heterogeneity in the therapeutic intervention types and method of delivery. None of the included studies has a high-grade quality of evidence in both short- and long-term outcome assessment. Short-term changes in pain score, quality of life, and adverse therapy effects may not reflect the clinical practice’s real implication. Long-term outcomes data availability was minimal, with reports only on cognitive therapy, ALA, capsaicin, umPEA, topical clonazepam, and low-level laser therapy. There were other trials with similar or other treatments reported in this review but these were not included mainly due to short-term assessment of as little as 2 weeks (46,68). Publication limitation and error in the statistical study data led to limited statistical analysis comparing treatment and placebo groups. The significant efficacy of psychology and LLLT studies should be interpreted with caution due to unreported adverse effects (34,36,37). Varoni et al. is a cross-over trial assuming a sufficient wash-over period of melatonin 4 weeks before the next intervention (18). The small study samples for each group (range 10 to 33) do not provide a robust statistical power in their results. The definition of improvement or reduction in pain for categorical data analyses (RRs) were varied across the studies as some studies may have meant almost or complete recovery while other may have meant a range of numerical decrease in VAS scores.

Conclusion

In perspective, it is suggested that multicentre trials investigate various therapeutic techniques in regulating BMS pain and increase participant numbers to conclude the treatment guidelines for BMS. The sustainability of pain reduction or remission is not adequately studied due to a short assessment period of less than a year. No treatment achieves a 50% pain remission in BMS. Investigating the influence of BMS’s biopsychosocial and neurophysiological mechanisms will provide a robust framework for integrating its various con founding aetiology factors. Studies should be ideally designed with multi-arm comparison for various pharmacological and non-pharmacological treatments to grade the treatment efficacy based on the universally accepted BMS disease’s diagnosis criteria. Likewise, a greater volume for sample size, multicentre studies, and longitudinal follow-up studies will enhance BMS treatment strategies’ value. The beneficial effects exhibited for neuroprotective and analgesic auxiliary therapies such as phytomedicine and rTMS, and the behavioural therapy CBT, could be valuable alternatives or applied in conjunction with synthetic systemic drugs, with a lesser risk of adverse drug effects, and tailored, holistic individual patient treatment, rather than the disease itself.

Article highlights

- This paper systematically reviews the evidence base for medicines in treating BMS based on the recent ICOP definition.
- This reviews RCTs with a minimum follow-up of 2 months, which had not been conducted by any previous systematic review.
- There is evidence on the benefit of topical oral clonazepam and capsaicin and alternative medicines such as neuroprotective agents and cognitive behavioural therapy.
- There is still insufficient long term follow up on the sustainable benefits of each treatment and its side effects.
Declaration of conflicting interests

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References

1. International Classiﬁcation of Orofacial Pain, 1st edition (ICOP). Cephalalgia 2020; 40(2): 129–221.
2. Kohorst JJ, Bruce AJ, Torgerson RR, et al. The prevalence of burning mouth syndrome: A population-based study. Br J Dermatol 2015; 172: 1654–1656.
3. Bergdahl M and Bergdahl J. Burning mouth syndrome: Prevalence and associated factors. J Oral Pathol Med 1999; 28: 350–354.
4. Scala A, Checchi L, Montevacchi M, et al. Update on burning mouth syndrome: Overview and patient management. Crit Rev Oral Biol Med 2003; 14: 275–291.
5. Forssell H, Teerijoki-Oksa T, Kotiranta U, et al. Pain and pain behavior in burning mouth syndrome: A pain diary study. J Orofac Pain 2012; 26: 117–125.
6. Schiavone V, Adamo D, Ventrella G, et al. Anxiety, depression, and pain in burning mouth syndrome: First chicken or egg? Headache 2012; 52: 1019–1025.
7. Jääskeläinen SK and Woda A. Burning mouth syndrome. Cephalalgia 2017; 37: 627–647.
8. Sardella A, Lodi G, Demarosi F, et al. Burning mouth syndrome: Retrospective study investigating spontaneous remission and response to treatments. Oral Dis 2006; 12: 152–155.
9. Liu YF, Kim Y, Yoo T, et al. Burning mouth syndrome: A systematic review of treatments. Oral Dis 2018; 24: 325–334.
10. De Souza IF, Mármore BC, Rados PV, et al. Treatment modalities for burning mouth syndrome: A systematic review. Clin Oral Investig 2018; 22: 1893–1905.
11. McMillan R, Forssell H, Buchanan JA, et al. Interventions for treating burning mouth syndrome. Cochrane Database Syst Rev 2016; 11: CD002779.
12. Moher D, Liberati A, Tetzlaff J, et al. Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. PLoS Med 2009; 6: e1000097.
13. Higgins JP, Altman DG, Gøtzsche PC, et al; Cochrane Bias Methods Group; Cochrane Statistical Methods Group. The Cochrane Collaboration’s tool for assessing risk of bias in randomised trials. BMJ 2011; 343: d5928.
14. Brozek JL, Akl EA, Alonso-Coello P, et al. Grading quality of evidence and strength of recommendations in clinical practice guidelines. Part 1 of 3. An overview of the GRADE approach and grading quality of evidence about interventions. Allergy 2009; 64: 669–677.
15. Altman DG and Dore CJ. Randomisation and baseline comparisons in clinical trials. Lancet 1990; 335: 149–153.
16. Pagano M and Gauvreau K. Principles of biostatistics, 2nd edn. Belmont, CA: Brooks/Cole, 2000, p.592.
17. Deeks JJ and Higgins JPT. Statistical algorithms in Review Manager 5 2010, https://training.cochrane.org/handbook/statistical-methods-revman5 (2010, accessed 3 January 2021).
18. Varoni EM, Lo Faro AF, Lodi G, et al. Melatonin treatment in patients with burning mouth syndrome: A triple-blind, placebo-controlled, crossover randomised clinical trial. J Oral Facial Pain Headache 2018; 32: 178–188.
19. De Rivera Campillo ER, López-López J and Chimenos-Küstner E. Response to topical clonazepam in patients with burning mouth syndrome: A clinical study. Bull Group Int Rech Sci Stomatol Odontol 2010; 49: 19–29.
20. Heckmann SM, Kirchner E, Grushka M, et al. A double-blind study on clonazepam in patients with burning mouth syndrome. Laryngoscope 2012; 122: 813–816.
21. Tammla-Salanen T and Forssell H. Trazodone in burning mouth pain: A placebo-controlled, double-blind study. J Orofac Pain 1999; 13: 83–88.
22. Femiao F and Scully C. Burning mouth syndrome (BMS): Double blind controlled study of alpha-lipoic acid (thioctic acid) therapy. J Oral Pathol Med 2002; 31: 267–269.
23. Carbone M, Pentenero M, Carrozzo M, et al. Lack of efficacy of alpha-lipoic acid in burning mouth syndrome: A double-blind, randomized, placebo-controlled study. Eur J Pain 2009; 13: 492–496.
24. López-Jornet P, Camacho-Alonso F and Leon-Espinosa S. Efficacy of alpha lipoic acid in burning mouth syndrome: A randomized, placebo-treatment study. J Oral Rehabil 2009; 36: 52–57.
25. López-D’ Alessandro E and Escovich L. Combination of alpha lipoic acid and gabapentin, its efficacy in the treatment of Burning Mouth Syndrome: A randomized, double-blind, placebo controlled trial. Med Oral Patol Oral Cir Bucal 2011; 16: e635–e640.
26. Palacios-Sánchez B, Moreno-López LA, Cerero-Lapi edra R, et al. Alpha lipoic acid efficacy in burning
mouth syndrome. A controlled clinical trial. Med Oral Patol Oral Cir Bucal 2015; 20: e435–e440.
27. Cano-Carrillo P, Pons-Fuster A and López-Jornet P. Efficacy of lycopene-enriched virgin olive oil for treating burning mouth syndrome: A double-blind randomised. J Oral Rehabil 2014; 41: 296–305.
28. Sardella A, Lodi G, Demarosi F, et al. Hypericum perforatum extract in burning mouth syndrome: A randomised placebo-controlled study. J Oral Pathol Med 2008; 37: 395–401.
29. Spanemberg JC, Cherubini K, De Figueiredo MA, et al. Effect of an herbal compound for treatment of burning mouth syndrome: Randomized, controlled, double-blind clinical trial. Oral Surg Oral Med Oral Pathol Oral Radiol 2012; 113: 373–377.
30. Pakfetrat A, Talebi M, Dalirsani Z, et al. Evaluation of the effectiveness of crocin isolated from saffron in treatment of burning mouth syndrome: A randomized controlled trial. Avicenna J Phytomed 2019; 9: 505–516.
31. Ottaviani G, Rupel K, Gobbo M, et al. Efficacy of ultramicronized palmitoylethanolamide in burning mouth syndrome-affected patients: A preliminary randomized double-blind controlled trial. Clin Oral Investig 2019; 23: 2743–2750.
32. De Silva LA, Siqueira JT, Teixeira MJ, et al. The role of xerostomia in burning mouth syndrome: A case-control study. Arg Neuropsiquiatr 2014; 72: 91–98.
33. Marino R, Torretta S, Capaccio P, et al. Different therapeutic strategies for burning mouth syndrome: Preliminary data. J Oral Pathol Med 2010; 39: 611–616.
34. Spanemberg JC, López López J, De Figueiredo MA, et al. Efficacy of low-level laser therapy for the treatment of burning mouth syndrome: A randomized, controlled trial. J Biomed Opt 2015; 20: 098001.
35. Umezaki Y, Badran BW, DeVries WH, et al. The efficacy of daily prefrontal repetitive transcranial magnetic stimulation (rTMS) for burning mouth syndrome (BMS): A randomized controlled single-blind study. Brain Stimul 2016; 9: 234–242.
36. De Pedro M, López-Pintor RM, Casonas E, et al. Effects of photobiomodulation with low-level laser therapy in burning mouth syndrome: A randomized clinical trial. Oral Dis 2020; 26: 1764–1776.
37. Bergdahl J, Anneroth G and Perris H. Cognitive therapy in the treatment of patients with resistant burning mouth syndrome: A controlled study. J Oral Pathol Med 1995; 24: 213–215.
38. López-Jornet P, Camacho-Alonso F and Andujar-Mateos P. A prospective, randomized study on the efficacy of tongue protector in patients with burning mouth syndrome. Oral Dis 2011; 17: 277–282.
39. Çınar SL, Kartal D, Pergel T, et al. Effectiveness and safety of clonazepam, pregabalin, and alpha lipoic acid for the treatment of burning mouth syndrome. Erciyes Med J 2018; 40: 35–38.
40. Imamura Y, Shinozaki T, Okada-Ogawa A, et al. An updated review on pathophysiology and management of burning mouth syndrome with endocrinological, psychological and neuropathic perspectives. J Oral Rehabil 2019; 46: 574–587.
41. Galli F, Lodi G, Sardella A, et al. Role of psychological factors in burning mouth syndrome: A systematic review and meta-analysis. Cephalalgia 2017; 37: 265–277.
42. Jääskéliäinen SK. Pathophysiology of primary burning mouth syndrome. Clin Neurophysiol 2012; 123: 71–77.
43. Kim JY, Kim YS, Ko I, et al. Association between burning mouth syndrome and the development of depression, anxiety, dementia, and Parkinson disease. JAMA Otolaryngol Head Neck Surg 2020; 146: 561–569.
44. Griffin CE 3rd, Kaye AM, Bueno FR, et al. Benzodiazepine pharmacology and central nervous system-mediated effects. Ochsner J 2013; 13: 214–223.
45. Bartoshuk LM, Snyder DJ, Grushka M, et al. Taste damage: Previously unsuspected consequences. Chem Senses 2005; 30: i218–i219.
46. Gremu-Richard C, Woda A, Navez ML, et al. Topical clonazepam in stomatodynia: A randomised placebo-controlled study. Pain 2004; 108: 51–57.
47. Fenelon M, Quinque E, Arrive E, et al. Pain-relieving effects of clonazepam and amitriptyline in burning mouth syndrome: A retrospective study. Int J Oral Maxillofac Surg 2017; 46: 1505–1511.
48. Wiffen PJ, Derry S, Bell RF, et al. Gabapentin for chronic neuropathic pain in adults. Cochrane Database Syst Rev 2017; 6: CD007938.
49. Takasuishi T and Yaksh TL. The effects of intrathecal and systemic gabapentin on spinal substance P release. Anesth Analg 2011; 112: 971–976.
50. Quintero JE, Dooley DJ, Pomerleau F, et al. Ampereometric measurement of glutamate release modulation by gabapentin and pregabalin in rat neocortical slices: Role of voltage-sensitive Ca2+ x2-1 subunit. J Pharmacol Exp Ther 2011; 338: 240–245.
51. Khan SA, Keaser ML, Meiller TF, et al. Altered structure and function in the hippocampus and medial prefrontal cortex in patients with burning mouth syndrome. Pain 2014; 155: 1472–1480.
52. Yang G, Su S, Jie H, et al. Somatosensory profiling of patients with burning mouth syndrome and correlations with psychologic factors. J Oral Facial Pain Headache 2019; 33: 278–286.
53. Zorunski CF, Paul SM, Covey DF, et al. Neurosteroids as novel antidepressants and anxiolytics: GABA-A receptors and beyond. Neurobiol Stress 2019; 11: 100196.
54. Morillas-Arques P, Rodriguez-Lopez CM, Molina-Barea R, et al. Trazodone for the treatment of fibromyalgia: An open-label, 12-week study. BMC Musculoskelet Disord 2010; 11: 204.
55. Patetsos E and Horjales-Araujo E. Treating chronic pain with SSRIs: What do we know? Pain Res Manag 2016; 2016: 2020915.
56. Dharmshaktu P, Tayal V and Kalra BS. Efficacy of antidepressants as analgesics: A review. J Clin Pharmacol 2012; 52: 6–17.
57. World Health Organization. *WHO global report on traditional and complementary medicine*. Geneva: WHO, 2019, p.226.

58. Khan H. Brilliant future of phytomedicines in the light of latest technological developments. *J Phytomedic* 2015; 4: 58–60.

59. Pan YZ and Pan HL. Primary afferent stimulation differentially potentiates excitatory and inhibitory inputs to spinal lamina II outer and inner neurons. *J Neurophysiol* 2004; 91: 2413–2421.

60. Yilmaz Z, Renton T, Yiango Y, et al. Burning mouth syndrome as a trigeminal small fibre neuropathy: Increased heat and capsaicin receptor TRPV1 in nerve fibres correlates with pain score. *J Clin Neurosci* 2007; 14: 864–871.

61. Game S, Petsche U, Lembeck F, et al. Capsaicin applied to peripheral nerve inhibits axoplasmic transport of substance P and somatostatin. *Brain Res* 1982; 239: 447–462.

62. Anand P and Bley K. Topical capsaicin for pain management: Therapeutic potential and mechanisms of action of the new high-concentration capsaicin 8% patch. *Br J Anaesth* 2011; 107: 490–502.

63. Nolano M, Simone DA, Wendelschafer-Crabb G, et al. Topical capsaicin in humans: Parallel loss of epidermal nerve fibers and pain sensation. *Pain* 1999; 81: 135–145.

64. Derry S, Rice AS, Cole P, et al. Topical capsaicin (high concentration) for chronic neuropathic pain in adults. *Cochrane Database Syst Rev* 2017; 1: CD007393.

65. Maihofner C and Heskamp ML. Prospective, non-interventional study on the tolerability and analgesic effectiveness over 12 weeks after a single application of capsaicin 8% cutaneous patch in 1044 patients with peripheral neuropathic pain: First results of the QUEPP study. *Curr Med Res Opin* 2013; 29: 673–683.

66. Mou J, Paillard F, Turnbull B, et al. Efficacy of Qutenza® (capsaicin) 8% patch for neuropathic pain: A meta-analysis of the Qutenza Clinical Trials Database. *Pain* 2013; 154: 1632–1639.

67. Altman RD and Barthel HR. Topical therapies for osteoarthritis. *Drugs* 2011; 71: 1259–1279.

68. Petruzzi M, Lauritano D, De Beneditis M, et al. Systemic capsaicin for burning mouth syndrome: Short-term results of a pilot study. *J Oral Pathol Med* 2004; 33: 111–114.

69. Jensen TS, Hoye K, Fricová J, et al. Tolerability of the capsaicin 8% patch following pretreatment with lidocaine or tramadol in patients with peripheral neuropathic pain: A multicentre, randomized, assessor-blinded study. *Eur J Pain* 2014; 18: 1240–1247.

70. Skaper SD. Impact of inflammation on the blood-neural barrier and blood-nervous: From review to therapeutic preview. *Int Rev Neurobiol* 2017; 137: 29–45.

71. Scuderi C, Bronzuoli MR, Facchinetti R, et al. Ultramicronized palmitoylethanolamide rescues learning and memory impairments in a triple transgenic mouse model of Alzheimer’s disease by exerting anti-inflammatory and neuroprotective effects. *Transl Psychiatry* 2018; 8: 32.

72. Passavanti MB, Fiore M, Sansone P, et al. The beneficial use of ultramicronized palmitoylethanolamide as add-on therapy to Tapentadol in the treatment of low back pain: A pilot study comparing prospective and retrospective observational arms. *BMC Anesthesiol* 2017; 17: 171.

73. Artukoglu BB, Beyer C, Zuloff-Shani A, et al. Efficacy of Palmitoylethanolamide for pain: A meta-analysis. *Pain Physician* 2017; 20: 353–362.

74. Paladini A, Fusco M, Cenacchi T, et al. Palmitoylethanolamide, a special food for medical purposes, in the treatment of chronic pain: A pooled data meta-analysis. *Pain Physician* 2016; 19: 11–24.

75. Cocito D, Peci E, Ciaramitaro P, et al. Short-term efficacy of ultramicronized palmitoylethanolamide in peripheral neuropathic pain. *Pain Res Treat* 2014; 2014: 854560.

76. Quintão NL, Ferreira J, Beirith A, et al. Evaluation of the effects of the herbal product Catuama in inflammatory and neuropathic models of nociception in rats. *Phytomedicine* 2008; 15: 245–252.

77. Coppock RW and Dziwenka M. Chapter 45 – St. John’s Wort. In: Gupta RC (ed) *Nutraceuticals*. Amsterdam: Academic Press/Elsevier, 2016, pp.619–631.

78. Müller WE, Rolli M, Schäfer C, et al. Effects of hypericum extract (LI 160) in biochemical models of antidepressant activity. *Pharmacopsychiatry* 1997; 30: 102–107.

79. McCarson KE and Enna SJ. GABA pharmacology: The search for analogics. *Neurochem Res* 2014; 39: 1948–1963.

80. Hippius H. St John’s Wort (Hypericum perforatum): A herbal antidepressant. *Curr Med Res Opin* 1998; 14: 171–184.

81. Ng QX, Venkatanarayanan N and Ho CY. Clinical use of Hypericum perforatum (St John’s wort) in depression: A meta-analysis. *J Affect Disord* 2013; 29: 171–184.

82. Henderson L, Yue QY, Bergquist C, et al. St John’s Wort and depression: In depth, outcomes. *Br J Clin Pharmacol* 2017; 137: 29–45.

83. National Center for Complementary and Integrative Health. St. John’s Wort and depression: In depth, outcomes. *Br J Clin Pharmacol* 2002; 54: 349–356.

84. The protective effects of crocin in the management of neurodegenerative diseases: A review. *Am J Neurodegener Dis* 2018; 7: 1–10.

85. Yorgun MA, Rashid K, Aslanidis A, et al. Crocin, a plant-derived carotenoid, modulates microglial reactivity. *Biochem Biophys Rep* 2017; 12: 245–250.

86. Li Q and Barres BA. Microglia and macrophages in brain homeostasis and disease. *Nat Rev Immunol* 2018; 18: 225–242.

87. Simpson DSA and Oliver PL. ROS generation in microglia: Understanding oxidative stress and inflammation in neurodegenerative disease. *Antioxidants (Basel)* 2020; 9: 743.
88. Singh A, Kukreti R, Saso L, et al. Oxidative stress: A key modulator in neurodegenerative diseases. *Molecules* 2019; 24: 1583.

89. Clifford TJ, Warsi MJ, Burnett CA, et al. Burning mouth in Parkinson’s disease sufferers. *Gerodontology* 1998; 15: 73–78.

90. Kim JY, Kim YS, Ko I, et al. Association between burning mouth syndrome and the development of depression, anxiety, dementia, and Parkinson disease. *JAMA Otolaryngol Head Neck Surg* 2020; 146: 561–569.

91. Cobley JN, Fiorello ML and Bailey DM. 13 reasons why the brain is susceptible to oxidative stress. *Redox Biol* 2018; 15: 490–503.

92. Gaforio JJ, Visioli F, Alarcón-de-la-Lastra C, et al. Virgin olive oil and health: Summary of the III international conference on virgin olive oil and health consensus report. *JAEN (Spain)* 2018. *Nutrients* 2019; 11: 2039.

93. Gupta S, Jawanda MK, Arora V, et al. Role of lycopene in preventing oral diseases as a nonsurgical aid of treatment. *Int J Prev Med* 2015; 6: 70.

94. Garrido M, González-Flores D, Marchena AM, et al. A lycopene-enriched virgin olive oil enhances antioxidant status in humans. *J Sci Food Agric* 2013; 93: 1820–1826.

95. Ametov AS, Barinov A, Dyck PJ, et al. SYDNEY trial study group. The sensory symptoms of diabetic polyneuropathy are improved with alpha-lipoic acid: The SYDNEY trial. *Diabetes Care* 2003; 26: 770–776.

96. Ziegler D, Low PA, Litchy WJ, et al. Efficacy and safety of antioxidant treatment with z-lipoic acid over 4 years in diabetic polyneuropathy: The NATHAN 1 trial. *Diabetes Care* 2011; 34: 2054–2060.

97. Keith DJ, Butler JA, Bemer B, et al. Age and gender dependent bioavailability of R- and R,S-z-lipoic acid: A pilot study. *Pharmacol Res* 2012; 66: 199–206.

98. Vaughan R, Galley H and Kanakarajan S. Assessment of sleep in patients with chronic pain. *BJA* 2018; 120: e10.

99. Wilhelmsen M, Amirian I, Reiter RJ, et al. Analgesic effects of melatonin: A review of current evidence from experimental and clinical studies. *J Pineal Res* 2011; 51: 270–277.

100. Hussain SA, Al-Khalifa IF, Jasim NA, et al. Adjuvant use of melatonin for treatment of fibromyalgia. *J Pineal Res* 2011; 50: 267–271.

101. Vidor LP, Torres IL, De Souza ICC, et al. Analgesic and sedative effects of melatonin in temporomandibular disorders: A double-blind, randomized, parallel-group, placebo-controlled study. *J Pain Symptom Manage* 2013; 46: 422–432.

102. Ulugol A, Dokmeci D, Guray G, et al. Antihyperalgesic, but not antiallodynic, effect of melatonin in nerve-injured neuropathic mice: Possible involvements of the L-arginine-NO pathway and opioid system. *Life Sci* 2006; 78: 1592–1597.

103. Ambriz-Tututi M and Granados-Soto V. Oral and spinal melatonin reduces tactile allodynia in rats via activation of MT2 and opioid receptors. *Pain* 2007; 132: 273–280.

104. Zhu C, Xu Y, Duan Y, et al. Exogenous melatonin in the treatment of pain: A systematic review and meta-analysis. *Oncotarget* 2017; 8: 100582–100592.

105. Danilov A and Kurganova J. Melatonin in chronic pain syndromes. *Pain Ther* 2016; 5: 1–17.

106. Bjordal JM, Couppé C, Chow RT, et al. A systematic review of low level laser therapy with location-specific doses for pain from chronic joint disorders. *Aust J Physiother* 2003; 49: 107–116.

107. Pandeshwar P, Roa MD, Das R, et al. Photobiomodulation in oral medicine: A review. *J Investig Clin Dent* 2016; 7: 114–126.

108. Prockt AP, Takahashi A and Pagnoncelli RM. Use of low intensity laser therapy in maxillofacial surgery. *Rev Port Estomatol Cir Maxilofac* 2008; 49: 247–255.

109. Nasri-Heir C, Zagury JG, Thomas D, et al. Burning mouth syndrome: Current concepts. *J Indian Prosthodont Soc* 2015; 15: 300–307.

110. Imura H, Shimada M, Yamazaki Y, et al. Characteristic changes of saliva and taste in burning mouth syndrome patients. *J Oral Pathol Med* 2016; 45: 231–236.

111. Moore RA, Derry S, Aldington D, et al. Amitriptyline for neuropathic pain in adults. *Cochrane Database Syst Rev* 2015; 7: CD008242.

112. Lee YC, Hong IK, Na SY, et al. Evaluation of salivary function in patients with burning mouth syndrome. *Oral Dis* 2015; 21: 308–313.

113. Fernandez-Pena C and Viana F. Targeting TRPM8 for pain relief. *Open Pain J* 2013; 6: 154–164.

114. Tan Y, Wu X, Chen J et al. Structural and functional connectivity between the amygdala and orbital frontal cortex in burning mouth syndrome: An fMRI study. *Front Psychol* 2019; 10: 1700.

115. Passard A, Attal N, Benadhira R, et al. Effects of unilateral repetitive transcranial magnetic stimulation of the motor cortex on chronic widespread pain in fibromyalgia. *Brain* 2007; 130: 2661–2670.

116. Moisset X, De Andrade DC and Bouhassira D. From pulses to pain relief: An update on the mechanisms of rTMS-induced analgesic effects. *Eur J Pain* 2016; 20: 689–700.

117. López-Jornet P, Camacho-Alonso F and Leon-Espinosa S. Burning mouth syndrome, oral parafuncions, and psychological profile in a longitudinal case study. *J Eur Acad Dermatol Venereol* 2009; 23: 363–365.

118. Lim JA, Choi SH, Lee WJ, et al. Cognitive-behavioral therapy for patients with chronic pain: Implications of gender differences in empathy. *Medicine (Baltimore)* 2018; 97: e10867.

119. Hofmann SG, Asnaani A, Vonk IJ, et al. The efficacy of cognitive behavioral therapy: A review of meta-analyses. *Cognit Ther Res* 2012; 36: 427–440.

120. Sardella A, Lodi G, Tarozzi M, et al. Acupuncture and burning mouth syndrome: A pilot study. *Pain Pract* 2013; 13: 627–632.

121. Jurisic Kveisic A, Zavoreo I, Basic Kes V, et al. The effectiveness of acupuncture versus clonazepam in patients with burning mouth syndrome. *Acupunct Med* 2015; 33: 289–292.
122. Franco FR, Castro LA, Borsatto MC, et al. Combined acupuncture and auriculotherapy in burning mouth syndrome treatment: A preliminary single-arm clinical trial. *J Altern Complement Med* 2017; 23: 126–134.

123. Iris Z, Miroslav S, Vanja VB, et al. Efficacy of acupuncture and vitamin C in burning mouth syndrome: A pilot study. *Res J Pharm Biol Chem Sci* 2017; 8: 26–30.

124. Diep CP, Goddard G and Mauro G. Acupuncture improves pain and rest in burning mouth syndrome patients: A clinical cases. *Dental Cadmos* 2019; 87: 548–554.

125. Scardina GA, Ruggieri A, Provenzana F, et al. Burning mouth syndrome: Is acupuncture a therapeutic possibility? *Br Dent J* 2010; 209; E2.