





The potential of liquid smoke as an oral ulcer remedies: A proposed mechanism based on systematic review

[El potencial del humo líquido como remedio para las úlceras orales: Un mecanismo propuesto basado en una revisión sistemática]

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Abstract

Context: The standard management of oral ulcer therapy is focused only on symptomatic therapy, such as reducing pain. To date, there is no topical drug that has the pharmacodynamics to intervene in oral ulcer pathogenicity. Liquid smoke is traditionally used as a safe natural preservative. The liquid smoke is highly phenolic and compound rich. It is presumed to have analgesic and anti-inflammatory effects with potentially promising therapeutic effects on oral ulcers.

Aims: To describe the possible pharmacodynamics or action mechanism of liquid smoke as a promising remedy for oral ulcer therapy.

Methods: A comprehensive literature review on PubMed, ScienceDirect, Scopus and Embase was performed using Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA). The keywords used included 'liquid smoke', 'wood vinegar', 'liquid pyrolysis' and 'oral ulcer'. A screening process, including titles, abstracts and full texts, was performed. Eight related articles were selected to describe the possible pharmacodynamics or mechanism action of the liquid smoke originating from coconut shells and rice hulls for oral ulcer remedies.

Results: Liquid smoke from coconut shell and rice husk is highly contained phenol, guaiacol and 2-methoxy-5-methylphenol (2-EMP). These compounds are antioxidants that can bind reactive oxygen species and increase cellular responses, inhibiting nuclear factor-kappa B activation and pro-inflammatory cytokine production while increasing macrophage differentiation to M2. The increase of M2, with help from lymphocytes, can secrete various growth factors, which can accelerate the proliferation of fibroblasts and collagens needed in the healing process of oral ulcers.

Conclusions: Liquid smoke pharmacodynamics inhibit both inflammatory and proliferation pathway stimulation, which promises remedies for oral ulcers.

Keywords: analgesic; anti-inflammatory; coconut shell-liquid smoke; human health; oral ulcer.

Resumen

Contexto: El humo líquido es altamente fenólico y rico en compuestos. Se presume que este tiene efectos analgésicos y antiinflamatorios con prometedores efectos terapéuticos potenciales sobre las úlceras orales.

Objetivos: Describir la posible farmacodinámica o mecanismo de acción del humo líquido como un remedio prometedor para la terapia de úlceras orales.

Métodos: Se realizó una revisión exhaustiva de la literatura sobre PubMed, ScienceDirect, Scopus y Embase utilizando elementos de informes preferidos para revisiones sistemáticas y metaanálisis (PRISMA). Las palabras clave utilizadas incluyeron "humo líquido", "vinagre de madera", "pirólisis líquida" y "úlcera oral". Se realizó un proceso de selección, que incluyó títulos, resúmenes y textos completos. Se seleccionaron ocho artículos relacionados para describir la posible farmacodinámica o mecanismo de acción del humo líquido procedente de las cáscaras de coco y de arroz para remediar las úlceras orales.

Resultados: El humo líquido de la cáscara de coco y la cáscara de arroz tiene un alto contenido de fenol, guayacol y 2-metoxi-5-metilfenol (2-EMP). Estos compuestos son antioxidantes que pueden unirse a especies reactivas de oxígeno y aumentar las respuestas celulares, inhibiendo la activación del factor nuclear kappa B y la producción de citocinas proinflamatorias al tiempo que aumentan la diferenciación de macrófagos a M2. El aumento de M2, con la ayuda de los linfocitos, puede secretar varios factores de crecimiento, que pueden acelerar la proliferación de fibroblastos y colágenos necesarios en el proceso de curación de las úlceras orales.

Conclusiones: La farmacodinamia del humo líquido inhibe la estimulación de las vías de proliferación e inflamación, lo que promete remedios para las úlceras orales.

Palabras Clave: analgésico; antiinflamatorio; humo líquido de cáscara de coco; salud humana; úlcera oral.

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INTRODUCTION

Oral ulcers are a condition in which the epithelial tissue loses its continuity due to molecular necrosis, which manifests in the oral cavity (Lewis and Wilson, 2019). To date, oral ulcer treatment has only focused on symptomatic therapy, such as relieving the pain, which occurs as a symptom of the oral ulcer. In general, the conventional therapy for oral ulcers is divided into topical and systemic therapies (Tarakji et al., 2015).

Topical therapy is the first line of defense in oral ulcer management and is usually administered as a mouth spray or mouthwash with chlorhexidine, benzydamine hydrochloride (BHCl), povidone-iodine and triamcinolone acetonide as the active agents (Fani et al., 2012; Lewis and Wilson, 2019; Teixeira, 2019). However, topical therapy shows less efficiency in the absorption process, which causes inaccuracies of the dosages used (Bhattarai and Gupta, 2016). In addition, the only purpose of topical therapy is to relieve pain symptoms, and in some long-term cases, it can cause side effects, such as brownish staining of the teeth and yellowish tongue (Prasad et al., 2015; Pandya et al., 2017).

Systemic therapy may be suggested if topical therapy has not shown significant results in the healing of oral ulcers, especially in moderate to severe oral ulcers. Systemic therapy, such as amlexanox and corticosteroids, are usually administered orally (Lewis and Wilson, 2019). However, the mechanism of action of amlexanox is still unknown, and long-term use of corticosteroids has side effects on other organs, such as the liver and the digestive tract, and may increase the incidence of *Candida* infection in the oral cavity (Abbasi et al., 2016; Hernawati et al., 2019; Lewis and Wilson, 2019).

Therefore, other therapeutic agents, which are able to safely intervene in the pathogenicity of oral ulcers, able to improve the healing process of the ulcers and act as more than just a pain reliever are needed. A natural ingredient that has been proven to have various benefits in the wound healing pro-

cess is liquid smoke. There are several studies that observe liquid smoke and its potential for human health, including liquid smoke from coconut shells (Surboyo et al., 2019a), rice hull (Budhy et al., 2021), cocoa shell (Budaraga et al., 2019), and cocoa pod shell (Desvita et al., 2021). Liquid smoke is a natural substance that has the potential to accelerate wound healing (Tarawan et al., 2017) and heal oral ulcers (Arundina et al., 2021a; Surboyo et al., 2019a); it is antibacterial (Arundina et al., 2020a), analgesic (Surboyo et al., 2012), anti-inflammatory (Kim et al., 2011) and anti-diabetes (Yang et al., 2012a). In some of the research, liquid smoke also possesses mechanisms that increase the migration and response of macrophages, thereby accelerating the inflammatory process in oral ulcers (Surboyo et al., 2019a; 2019b), increasing the number of fibroblasts (Tarawan et al., 2017) and increasing collagen formation (Surboyo et al., 2017). All the promising effects of liquid smoke are expected to stimulate oral ulcer healing. This study aims to describe the possible pharmacodynamics of liquid smoke as a promising remedy for oral ulcer treatment.

MATERIAL AND METHODS

Review methodology

A systematic review of published studies on the topic of liquid smoke for oral ulcer therapy was conducted according to the PRISMA method (Selcuk, 2019). The focus question in this review was 'How do the pharmacodynamics or mechanism action of liquid smoke promise remedies for oral ulcer therapy?'

The Population, Intervention, Comparison, Outcome (PICO) statement used for this study is the population included in all the studies investigating the potential effect of liquid smoke on oral ulcers. Intervention is defined as the topical application of liquid smoke. Any type of comparison (placebo, another topical treatment or no control) was included. All clinical outcomes were considered, including any parameter – cellular, molecular or clinical – related to oral ulcers.

Information sources

A comprehensive literature search was conducted on the following databases: PubMed (US National Library of Medicine, USA), Science Direct (Elsevier, Netherlands), Scopus, and Embase for studies published between 2010 and April 2021.

Search strategy

The terms 'liquid smoke', 'wood vinegar', 'liquid pyrolysis' and 'oral ulcer' were used as keywords in the research. Results were limited to studies published in English. Review articles were not included in this review.

Study design and selection process

All studies found on those databases and fitting the criteria below were grouped together, and any duplicates were removed. The remaining studies were then filtered according to the title, followed by the abstract. Studies that did not match were excluded at this stage. The remaining studies were screened at the final stage according to their full text, and those that did not meet the inclusion criteria were excluded. Microsoft® Excel for Mac Version 16.47.1 was used to organise the study titles and abstracts and identify duplicates. This process was conducted by four independent investigators: KNA, PHC, AABN, and MDCS. In the case of any disagreements, the investigators reached their decision through discussion.

The inclusion criteria for this review included studies about liquid smoke, studies describing the effect on or the potential of liquid smoke for oral ulcers, the dose and duration of the treatment, the marker analyzed, and the analysis methods described. This process, documented by Microsoft® Excel for Mac Version 16.47.1, was performed in the following order: the name of the first author, publication year, country, study design, and results.

The exclusion criteria were *in vitro* studies and review articles. The diagram flow for the study design and selection process is shown in (Fig. 1). The final selected articles were analyzed in detail to assess the most current and relevant infor-

mation about the potential mechanism of action of liquid smoke on oral ulcers.

RESULTS

Study selection

Flow of the studies

After using a combination of keywords, 560 articles were found in the three databases. The titles were screened, and the duplicates were removed, resulting in 175 remaining articles. After reading the abstracts, only 28 articles were included in the next step of assessing the full text for eligibility. After this process, only 16 articles were included in the next step. The full texts of eight articles were included in the systematic review (Fig. 1).

Excluded studies

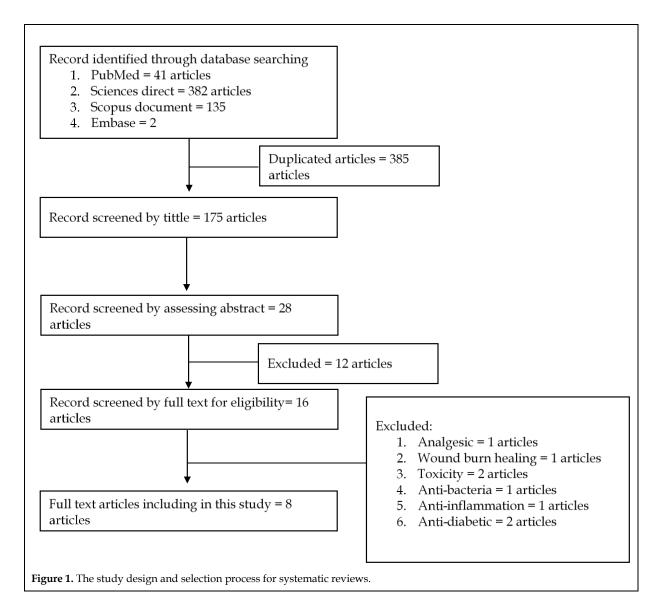
There were eight articles excluded from this review. Two articles assessed *in vitro* applications, three articles assessed *in vivo* applications, and three articles assessed both *in vitro* and *in vivo* applications. The excluded articles assessed the potential of liquid smoke from coconut shells (two articles) and rice hulls (six articles).

The excluded *in vitro* articles contained studies on the potential of liquid smoke from rice hulls and studies assessing its toxicity, antibacterial properties, antioxidant properties, anti-inflammatory properties, and antidiabetic properties (Table 1).

The excluded *in vivo* articles contained studies on the potential of coconut shell-liquid smoke (CS-LS), assessing its analgesic and wound healing properties. These articles also assessed the toxicity, anti-inflammatory properties, and antidiabetic properties of liquid smoke from rice hulls (Table 2).

Study characteristics

Six studies revealed the potential of CS-LS for healing oral ulcers (Table 3), and two studies examined liquid smoke from rice hulls (Table 4). All studies were *in vivo* experiments on the healing of oral ulcers conducted on animal models.



Results of individual studies

Potential of CS-LS

<u>Increased the neutrophils</u>

The topical administration of liquid smoke increased the number of neutrophils compared to a seven-day administration of BHCI (Ernawati et al., 2020).

<u>Increased the macrophages</u>

The topical administration of liquid smoke increased the number of macrophages compared to a five-day administration of BHCI (Surboyo et al., 2020). A seven-day duration increased the number

of macrophages better than five or three days (Ernawati et al., 2020).

Increased the fibroblasts

The topical administration of liquid smoke increased the number of fibroblasts compared to the five-day administration of BHCI (Ayuningtyas et al., 2020). A seven-day duration increased the number of fibroblasts compared to five days (Ernawati et al., 2020).

Increased oral ulcer healing

Topical administration of CS-LS decreased oral ulcer size by more than five- or seven-day administration of BHCI (Surboyo et al., 2019b).

Table 1. The *in vitro* experiment for assessing the potential of liquid smoke.

| 1 | | 0 1 1 | | |
|------------------------|---------------------------|--|---|--|
| Authors | Liquid smoke origin | Marker | Analysis methods | Result |
| Arundina et al., 2020a | Rice hull | Inhibition area of <i>Porphyromonas</i> gingivalis | Standard diffusion and dilution methods | The minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC) on was found in liquid smoke concentration 10% and 12.5% |
| | | Cell viability on osteoblast | MTT assay | The highest viability of osteoblast was analysed liquid smoke concentration 1% |
| Arundina et al., 2021c | Rice hull | Cell viability on baby hamster kidney | MTT assay | The highest viability of osteoblast was analysed liquid smoke concentration 10% |
| Kim et al., 2011 | Rice hull | Cell viability on RBL-2H3 and RAW264.7 | MTT assay | The highest viability of osteoblast was analysed liquid smoke concentration 1% |
| | | Antioxidant | DPPH assay | Liquid smoke exhibited strong anti-oxidative properties |
| | | Ionophore A23187-stimulated β -hexosaminidase release from RBL-2H3 | β-Hexosaminidase Secretion Assay | Liquid smoke inhibited the release of $\beta\mbox{-hexosaminidase}$ |
| | | NO production in RAW264.7 mouse | NO generation assay | Liquid smoke inhibited the NO production |
| | | TNF- α , IL-1 β , and IL-6 level | ELISA | Liquid smoke reduced TNF- α , IL-1 β , and IL-6 level |
| | | PGE ₂ and LTB ₄ level | ELISA | Liquid smoke reduced PGE2 and LTB4 level |
| | | Myeloperoxidase assay | ELISA | Liquid smoke reduced myeloperoxidase activity |
| | | ICAM-1, iNOS, 5-LOX, COX-2, TNF- α , IL-1 β , IL-6 and β -actin | RT-PCR | Liquid smoke modulated this gene expression |
| | | iNOS, 5-LOX, COX-2 and β -actin | Western blot | Liquid smoke modulated this protein expression |
| | | | | |

Table 1. The *in vitro* experiment for assessing the potential of liquid smoke (continued...)

| Authors | Liquid smoke origin | Marker | Analysis methods | Result |
|--------------------|---------------------------|--|--|--|
| Yang et al., 2012a | Rice hull | Oxidative stress | Non fluorescent probe, DCF-DA | Liquid smoke suppressed the intracellular peroxide level |
| | | Cell viability on rat insulinoma β-cell line INS-1 | MTT assay | Liquid smoke showed the highest cell viability |
| | | NO production on INS-1 cell | Membrane-permeable fluorescent indicator DAF- 2/DA | Liquid smoke decreased the NO production |
| | | Nitric oxide (NO) generation | Pro-inflammatory cytokine genes | Liquid smoke suppressed IL-1 β , IL-6, and TNF- α genes |
| | | iNOS gene | RT-PCR | Liquid smoke modulated the iNOS gene expression |
| | | iNOS protein | Western blot | Liquid smoke modulated the iNOS expression |
| | | TNF- α , IL-1 β , IL-6 and β -actin gene expression | RT-PCR | Liquid smoke modulated this gene expression |
| | | Insulin release on INS-1 cell | ELISA | Liquid smoke increased insulin level |
| | | Blood glucose | Rapid test | Liquid smoke lowered blood glucose level |
| | | Serum insulin | ELISA kit | Liquid smoke increased serum insulin level |
| | | Glycogen | weight | Liquid smoke restored glycogen content |
| | | GOT and GTP | Colorimetric kit | Liquid smoke lowered the level of GOT and GTP |
| | | C6, PEPCK and GCK gene expression | RT-PCR | Liquid smoke suppressed this gene expression |
| | | C6, PEPCK and GCK level | ELISA | Liquid smoke suppressed these protein level |
| Yang et al., 2012b | Rice hull | GOT and GTP | Colorimetric assay | Liquid smoke reduced serum levels of GOT and GTP |
| | | C6 Pase, GCK, PEPCK | Spectrophotometric | Liquid smoke lowered this enzyme |
| | | GLUT2 and PPAR-γ | RT-PCR | Liquid smoke increased the mRNA expression in hepatic and adipose |
| | | TNF- α , IL-1 β and IL-6 level | ELISA | Liquid smoke decreased these cytokines serum and adipose tissue |

Table 2. *In vivo* experiments (animal models) for assessing the potential of liquid smoke.

| Authors | Liquid smoke origin | Marker | Analysis methods | Result |
|------------------------|---------------------|------------------------------------|-------------------------------------|--|
| Tarawan et al., 2017 | Coconut shell | Wound contraction Fibroblast | Histopathology | Liquid smoke promotes burn wound healing by increasing the number of fibroblasts and wound contraction |
| Surboyo et al., 2012 | Coconut shell | Analgesic properties | Writhing reflex induced acetic acid | Liquid smoke concentration 100% has showed analgesic properties |
| Arundina et al., 2020b | Rice hull | Toxicity | Acute toxicity test | The toxicity dose of liquid smoke >15000mg/kg body weight and no toxicity symptoms in animals |
| Kim et al., 2011 | Rice hull | Ear thickness | Changes in ear thickness | Liquid smoke reduced ear thickness |
| Yang et al., 2012a | Rice hull | Liver | Histology using HE staining | Liquid smoke ameliorated liver damaged |
| | | Langerhans islet of the pancreas | Histology using HE staining | Liquid smoke restored the size and damage Langerhans islet |
| Yang et al., 2012b | Rice hull | Bodyweight | | Liquid smoke suppressed body weight gain induced by a high-fat diet |
| | | Liver weight | | Liquid smoke reduced the liver weight |
| | | White adipose tissue weight | | Liquid smoke reduced the white adipose tissue |
| | | Blood glucose | Rapid test | Liquid smoke suppressed blood glucose level |
| | | Serum insulin | ELISA | Liquid smoke increased serum insulin level |
| | | Oral glucose tolerance | Oral glucose tolerance test | Liquid smoke decreased oral glucose loading |
| | | Serum lipid | commercial kit | Liquid smoke lowered serum triglycerides |
| | | Liver tissue | Histopathology using HE staining | Liquid smoke ameliorated liver damage |
| | | Pancreatic islets of Langerhans | Histopathology using HE staining | Liquid smoke decreased islets of Langerhans |
| | | Pancreatic islets of Langerhans | IHC | Liquid smoke lowered insulin production |

Table 3. The properties of liquid smoke from coconut shell (CS-LS) for promotes oral ulcer healing.

| Authors | Marker | Analysis methods | Duration of administration | Result | P-value |
|--------------------------|---------------------|---|----------------------------|---|----------------|
| Surboyo et al., 2019a | NF-κB expression | Immunohistochemistry | 3 days 5 days | Topical administration of CS-LS lowered the NF-κB expression | <0.05 |
| | TNF-a expression | Immunohistochemistry | 3 days | Topical administration of CS-LS lowered the TNF- α expression | < 0.05 |
| Surboyo et al., 2019b | Oral ulcer diameter | Ulcer diameter using digital calipers | 5 days 7 days | Topical administration of CS-LS decreased the oral ulcer size | 0.005 0.004 |
| Surboyo et al., 2017 | Collagen | Histology by Masson trichome staining | 5 days | Topical administration of CS-LS increased the collagen numbers | 0.006 |
| Ernawati et al., 2020 | Fibroblast | Histology by hematoxylin-eosin staining | 7 days | Topical administration of CS-LS increased the fibroblast numbers | <0.01 |
| | Macrophages | Histology by hematoxylin-eosin staining | 7 days | Topical administration of CS-LS increased the macrophage numbers | <0.01 |
| | Neutrophils | Histology by hematoxylin-eosin staining | 7 days | Topical administration of CS-LS decreased the neutrophil numbers | <0.01 |
| | Lymphocytes | Histology by hematoxylin-eosin staining | 7 days | Topical administration of CS-LS increased the neutrophil numbers | <0.01 |
| Surboyo et al., 2020 | Macrophages | Histology by hematoxylin-eosin staining | 5 days | Topical administration of CS-LS decreased the macrophage numbers | 0.000 |
| | NF-κB expression | Immunohistochemistry | 5 days | Topical administration of CS-LS decreased the NF- κB expression | 0.000 |
| Ayuningtyas et al., 2020 | Fibroblast | Histology by hematoxylin-eosin staining | 5 days | Topical administration of CS-LS increased the fibroblast numbers | 0.028 |
| | FGF-2 expression | Immunohistochemistry | 7 days | Topical administration of CS-LS increased the FGF-2 expression | 0.000 |
| | VEGF expression | Immunohistochemistry | 5 days 7 days | Topical administration of CS-LS increased the VEGF expression | 0.019 |

Table 4. The properties of liquid smoke (LS) from rice hull that promote oral ulcer healing.

| Authors | Marker | Analysis methods | Duration of administration | Result | P-value |
|------------------------|------------------|---|----------------------------|--|---------|
| Arundina et al., 2021a | Macrophages | Histology by hematoxylin-eosin staining | 3 days | Topical administration of LS increased the | 0.005 |
| | | | 5 days | macrophage numbers | 0.022 |
| | | | 7 days | | 0.022 |
| | Lymphocytes | Histology by hematoxylin-eosin staining | 3 days | Topical administration of LS increased the | 0.001 |
| | | | 5 days | lymphocyte numbers | 0.000 |
| | Fibroblast | Histology by hematoxylin-eosin staining | 3 days | Topical administration of LS increased the | 0.032 |
| | | | 5 days | fibroblast numbers | 0.001 |
| | | | 7 days | | 0.025 |
| | IL-6 expression | Immunohistochemistry | 5 days | Topical administration of LS increased the IL-6 expression | 0.001 |
| | TGF-β expression | Immunohistochemistry | 3 days | Topical administration of LS increased the | 0.034 |
| | | | 7 days | TGF-b expression | 0.019 |
| Arundina et al., 2021b | FGF-2 expression | Immunohistochemistry | 5 days | Topical administration of LS increased the | 0.000 |
| | | | 7 days | FGF-2 expression | 0.009 |
| | VEGF expression | Immunohistochemistry | 7 days | Topical administration of LS increased the VEGF expression | 0.002 |
| | COL-1 expression | Immunohistochemistry | 5 days | Topical administration of LS increased the | 0.001 |
| | | | 7 days | COL-1 expression than control | 0.000 |
| | PDGF expression | Immunohistochemistry | 3 days | Topical administration of LS increased the | 0.002 |
| | | | 5 days | PDGF expression | 0.003 |
| | | | 7 days | | 0.003 |

<u>Increased the density of collagens</u>

The topical administration of liquid smoke increased the density of collagen compared to the five-day administration of BHCI (Surboyo et al., 2012).

Inhibited NF-κB expression

Topical administration of CS-LS showed lower NF-κB expression than three- or five-day administration of BHCI (Surboyo et al., 2019a). Five-day administration showed increased NF-κB expression (Surboyo et al., 2020).

<u>Inhibited TNF-α expression</u>

The topical administration of CS-LS showed lower TNF- α expression than three-day topical administration of BHCI (Surboyo et al., 2019a).

Increased FGF-2 expression

The topical administration of CS-LS increased the FGF-2 expression by more than seven-day administration of BHCI (Ayuningtyas et al., 2020).

Increased VEGF expression

The topical administration of CS-LS showed a higher expression of VEGF than five- or seven-day administration of BHCI (Ayuningtyas et al., 2020).

Potential of liquid smoke from rice hull

Increased the macrophages

The topical administration of liquid smoke increased the number of macrophages compared to three-, five- or seven-day administration of a control (Arundina et al., 2021a).

Increased the lymphocytes

The topical administration of liquid smoke increased the number of lymphocytes compared to the three- and five-day administration of a control (Arundina et al., 2021a).

Increased the fibroblasts

The topical administration of liquid smoke increased the number of fibroblasts compared to three-, five- or seven-day administration of a control (Arundina et al., 2021a).

<u>Increased FGF-2 expression</u>

Topical administration of liquid smoke increased the expression of FGF-2 compared to fiveand seven-day administration of a control (Arundina et al., 2021b).

Increased VEGF expression

The topical administration of liquid smoke increased VEGF expression compared to the sevenday administration of a control (Arundina et al., 2021b).

Increased IL-6 expression

Topical administration of liquid smoke increased the expression of IL-6 compared to the five-day administration of a control (Arundina et al., 2021a).

Increased TGF-β expression

Topical administration of liquid smoke increased TGF- β expression compared to three- and seven-day administration of a control (Arundina et al., 2021a).

<u>Increased PDGF expression</u>

Topical administration of liquid smoke increased the expression of PDGF compared to three-, five- and seven-day administration of a control (Arundina et al., 2021b).

Increased COL-1 expression

The topical administration of liquid smoke increased COL-1 expression compared to the fiveand seven-day administration of a control (Arundina et al., 2021b).

DISCUSSION

Oral ulcers have a complex healing process that includes the phases of inflammation, granulation, and tissue cell regeneration (Sunarjo et al., 2016). To promote the healing process of oral ulcers and act as more than just a pain reliever, a new therapy that is able to intervene in the pathogenicity of oral ulcers is needed. However, to date, there are no topical drugs that have the pharmacodynamics to intervene in the pathogenicity of oral ulcers.

Both CS-LS and liquid smoke from rice hulls are rich in phenolic components. The physical characteristics are similar: CS-LS has an acidity of 2.39 (Surboyo et al., 2019b), and liquid smoke from rice hull has an acidity of 2.296 (Arundina et al., 2021c). Liquid smoke has a pH range between 2.5 and 3.1 and is presumed to have carcinogenic properties; however, it has been shown to have non-toxic and therapeutic effects (Maryam, 2015). CS-LS is 36.6% phenolic components, 25.2% 2methoxyphenols (guaiacol), 5.2% EMP, 3.5% 4ethyl-2-methoxyphenols and 28 other minor components (Surboyo et al., 2019a). Liquid smoke from rice hull contains 13.45% guaiacol, 13.45% mequinol, and 10.52% phenol (Arundina et al., 2021c). In the analyzed studies, there were eight that explored the potential of liquid smoke for the topical therapy of oral ulcers, especially traumatic ulcers.

Liquid smoke has the potential to accelerate the healing of oral ulcers. The healing process requires a faster inflammatory response and a shorter inflammatory phase to provide faster tissue regeneration. The application of CS-LS is able to increase the number of fibroblasts, macrophages, and lymphocytes continuously after three days, up to seven days, except for the number of neutrophils that increase after three days, up to five days, but then decrease by the seventh day (Ernawati et al., 2020). In addition, the topical administration of liquid smoke from rice husk for three, five, and seven days showed a higher number of macrophages, lymphocytes, and fibroblasts (Arundina et al., 2021a). During the inflammatory phase, neutrophils, macrophages, fibroblasts, and endothelial cells produce ROS to compensate for the inflammatory response and protect from microorganism invasion. The overproduction of ROS can slow the healing process. Phenolic compounds contained in liquid smoke, such as EMP and guaiacol, have antioxidant abilities that can increase the cellular response, such as the recruitment of lymphocytes and macrophages. Increased ROS causes oxidative stress on the lymphocytes, which causes apoptosis; hence, with ROS inhibition, the number of lymphocytes will increase. When the inflammatory responses decrease, neutrophils cleanse bacteria and foreign materials from the ulcer area to provide a good environment for the oral ulcer healing process.

Liquid smoke has been shown to increase the recruitment of macrophages in the tissue. In oral ulcer healing, the topical administration of liquid smoke is able to increase fibroblast formation increased macrophage through recruitment (Surboyo et al., 2020). An in vivo study by Ernawati et al. (2020) confirms that topical administration of liquid smoke on oral ulcers for seven days increases the recruitment of macrophages. Macrophages play an important role in the wound healing process through M1 polarization in the early stages, followed by M2 polarization at the final stage (Okizaki et al., 2015). M2 plays a role in the formation of new vessels, and an increase in the number of macrophages correlates with a high microvessel density. A higher number of macrophages was also proven by Arundina et al. (2020) in vivo study, which shows that topical administration of liquid smoke from rice husk increases macrophages and produces growth factors.

After providing faster cellular responses, liquid smoke also plays a crucial role in the control of the inflammatory phase. This mechanism begins by inhibiting the formation of ROS by phenolic compounds contained in the liquid smoke. Part of the hydroxyl group (-OH) makes up the active site of the phenolic compounds contained in an aromatic hydrocarbon ring. This free-radical resistance mechanism consists of the binding of superoxide radicals (O_2 -) to the phenol hydroxyl group (-OH) so that NF- κ B activation decreases, leading to inhibition of TNF- α production and other pro-

inflammatory cytokines, such as IL-6 (Shahidi and Yeo, 2018). This condition has been confirmed by Surboyo et al. (2019a; 2020), who show that topical administration of liquid smoke for three, five, and seven days can reduce the expression of NF-κB. This mechanism may occur because the phenol content, such as guaiacol and EMP, in liquid smoke, affects the expression of NF-κB in macrophages through a hydroxyl group (-OH), which is able to bind superoxide (O2-) during signaling, resulting in the production of pro-inflammatory cytokines, such as TNF-α and IL-6. Superoxide binding (O₂-) inhibits phosphorylation IκB kinase (IKK), inhibitory degradation of the IκB factor, translocation of NF-κB and phosphorylation of p65 in the cell nucleus. Therefore, the activation and expression of NF-κB are decreased, leading to a reduction in the expression of pro-inflammatory cytokines (Lopes et al., 2019).

The proliferation stage of oral ulcer healing through topical administration of liquid smoke has been shown to stimulate fibroblast formation, increase collagen density, FGF-2 expression and VEGF expression, and accelerate oral ulcer contraction. The topical application of liquid smoke for seven days increases the number of fibroblasts and the expressions of FGF-2 and VEGF in oral ulcers (Ayuningtyas et al., 2020). In addition, growth factor expression and collagen formation are also positive in the administration of liquid smoke from rice husk. The topical administration of liquid smoke from rice husk shows higher TGFβ, FGF-2, VEGF, PDGF and COL-1 expression (Arundina et al., 2021b). The mechanism that might be involved uses anti-inflammatory effects that Inhibit NF-κB and TNF-α, resulting in more dominant M2 polarisation through a switch from M1 to M2. M2 secretes growth factors, including TGF-β, FGF-2, VEGF, PDGF, and COL-1. The result of this secretion is increased fibroblast proliferation, collagen production and new vascular formation, leading to the complete healing process (Arundina et al., 2021a; Ayuningtyas et al., 2020). The complete healing process is also supported by the liquid smoke, significantly increasing collagen synthesis (Surboyo et al., 2017) and being clinically

able to accelerate the contraction of oral ulcers (Surboyo et al., 2019b).

Finally, based on the literature, the role of liquid smoke in the healing of oral ulcers interferes with the cellular responses, inhibits the inflammatory phase and increases the proliferation phases. The purpose of this mechanism is shown in Fig. 2.

There is limited data on the possible mechanism or pharmacodynamics of liquid smoke in oral ulcer healing because most of the research focuses on the potential of liquid smoke as a natural preservative. The research data, excluding articles from this study, which are presented in Tables 1 and 2, also support the potential of liquid smoke in oral ulcer healing.

Liquid smoke has been confirmed as a safe material. It was confirmed that a 10% concentration of liquid smoke derived from rice hull showed the highest living cells *in vitro* (Arundina et al., 2021c). *In vivo*, it was also confirmed that liquid smoke from rice hull had a lethal dose value >15,000 mg/kg body weight, categorizing it as a relatively harmless substance (Arundina et al., 2020b). Despite its toxicity, studies have proven several potentials of liquid smoke derived from rice hulls, such as antibacterial properties and an increase in the proliferation effect (Arundina et al., 2020a).

Moreover, liquid smoke derived from rice hull also has the ability to suppress inducible nitric oxide synthase (iNOS) gene expression and proinflammatory cytokines, such as TNF- α , IL-1 β and IL-6 (Kim et al., 2011; Yang et al., 2012a). It also has an antidiabetic potential due to decreasing blood glucose and cholesterol concentrations, increasing serum insulin and hepar glycogen, and improving glucose tolerance, which may be beneficial for the management of oral ulcers in diabetic patients (Yang et al., 2012a; 2012b).

The other benefit of liquid smoke for oral ulcer healing is its analgesic properties. *In vitro*, liquid smoke from rice hull reduces prostaglandin E₂ (PGE₂) and leukotriene B₄ (LTB₄) levels (Kim et al., 2011), and *in vivo*, CS-LS showed analgesic properties by reducing pain-induced reflexes in experimental animals (Surboyo et al., 2012). The analge-

sic properties in liquid smoke are produced by its phenolic components, which inhibit the cyclooxygenase enzyme in the tissue by binding to the prostaglandin G₂ (PGG₂) and the prostaglandin H₂ (PGH₂) components, causing a decrease in the synthesis of PGE₂. Decreasing PGE₂ levels can interfere with the transduction mechanism of afferent nociceptors so that the pain caused by oral ulcers is reduced (Surboyo et al., 2012). This analgesic property is needed in topical oral ulcer therapy, considering the loss of epithelial tissue is the main complaint of oral ulcer pain.

An *in vivo* study by Tarawan et al. (2017) shows that CS-LS accelerates wound healing by increasing wound contraction and the number of fibroblasts. The components of CS-LS that play a role in the healing process are flavonoids, phenols and tannins. The flavonoids and phenols have antioxidant and anti-microbial effects that mediate wound healing by increasing collagen and protein and decreasing lipid peroxidation in granulation tissue. Tannins play a role in fibroblast proliferation; by secreting fibroblast growth factors from neutrophils, fibroblasts proliferate and synthesize collagen. Fibroblasts migrate and proliferate at the wound site during wound healing. The resulting extracellular matrix strengthens the edges of the wound and causes wound contraction.

The study by Kim et al. (2011) shows that topical application of liquid smoke from rice husk has the ability to change ear thickness or swelling, i.e., the inflammatory response characterized by delayed onset and long-term inflammation associated with leukocytes. The rate of leukocyte infiltration changes in ear thickening or swelling. The liquid smoke from rice husk suppresses leukocyte infiltration in the dermis through inhibition of intracellular adhesion molecule-1 (ICAM-1) expression.

Future research needs to further elaborate on the potential of the pharmacodynamics of liquid smoke as a remedy for oral ulcers. Based on the available data, liquid smoke is a promising development in oral ulcer medicine.

Interpretation

Finally, based on the literature, the role of liquid smoke in oral ulcer healing interferes with cellular responses, inhibiting the inflammatory phase and increasing the proliferation phases. The purpose of this mechanism is provided in Fig. 2.

Limitations of evidence

The available articles are limited to *in vivo* and *in vitro* research models, so the level of evidence provided is limited. Further research on liquid smoke needs to be done to add to the data supporting the mechanisms outlined in this review.

Limitations of review processes

The available articles are limited to *in vivo* and *in vitro* research models, so the level of evidence provided is limited.

Implications

Further research on liquid smoke needs to be done to add to the data supporting the mechanisms outlined in this review.

CONCLUSIONS

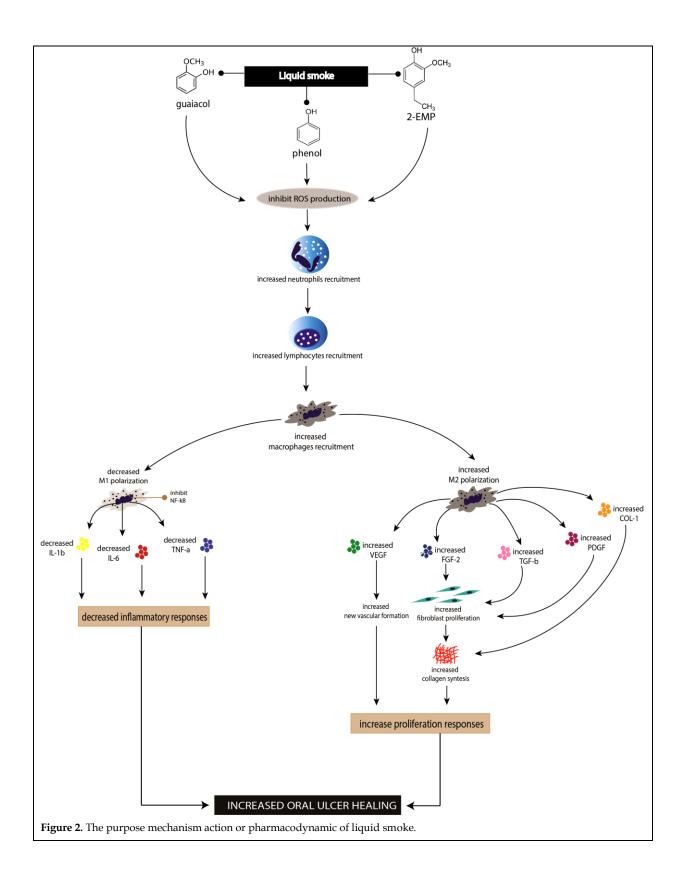
Based on the mechanisms described, it can be concluded that liquid smoke has a mechanism of action or pharmacodynamics that inhibits the inflammatory pathway and stimulates the proliferation pathway so that it is expected to become the topical drug of choice in oral ulcer therapy. However, further research is needed regarding its application and safe dosage in humans so that liquid smoke can be used clinically for oral ulcer therapy.

CONFLICT OF INTEREST

The authors declare no conflicts of interests.

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|------------------------|-------------|-------------|------------|----------|------------|-------------|-----------------|------------|------------|
| Concepts or ideas | x | | | | | | | | |
| Design | x | | | | | | | | |
| Literature search | x | | x | | | | x | x | x |
| Data acquisition | x | | | | | | | | |
| Data analysis | x | | x | x | | | x | x | x |
| Statistical analysis | x | | | | | | | | |
| Manuscript preparation | x | | | | | | x | x | x |
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