



A Systematic Review About the Contribution of The Genus *Tessaria* (Asteraceae) To Cancer Study and Treatment

Revisión sistemática sobre la contribución del género Tessaria (Asteraceae) al estudio y tratamiento del cáncer

Revisão sistemática sobre a contribuição do gênero Vernonia condensada Baker (Asteraceae) para o estudo e tratamento do câncer

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Abstract

Belonging to the Asteraceae family, *Tessaria* (Ruiz & Pavon, 1753) is a genus of shrubs or small trees distributed in various habitats located from Argentina to the southwestern United States of America. The taxa are composed of five confirmed species, which have been previously reported for their ethnopharmacological uses as hypocholesterolemic, hypoglycemic, antitussive, anti-inflammatory, anticancer, and abortifacient agent; it also has been used in hepatic, renal, pulmonary, and rheumatic diseases. Up to the present, *T. absinthioides*, *T. ambigua*, *T. dodoneifolia*, *T. fastigiata* and *T. integrifolia* have been chemically analyzed. While decoctions, infusions, methanolic, and hydromethanolic extracts are the most studied botanical compounds, its most recognized phytochemical constituents are caffeoylquinic acid, eupatorin, naringenin, protocatechuic, and quercetin. Scientifically, several biomedical properties such as virucidal, antibacterial, leishmanicidal, insecticidal, gastroprotective, antiasthmatic, hypoglycemic, and antiatherogenic were attributed to the genus. Concerning oncologic research, the chemicals produced by *Tessaria* have antitumor activities interfering with carcinogenesis, cell proliferation, metastasis, and angiogenesis. In addition, the natural extracts obtained from *Tessaria* species have biological activities closely related to cancer, acting as antioxidants and anti-inflammatories. In particular, it has been reported that *T. absinthioides* resulted cytotoxic against several cancer cell lines and acts as an antitumoral compound in murine models. Due to the aforementioned antecedents, the *Tessaria* species are considered undervalued within the oncological field; the goal of the current systematic review is to summarize the information available on the genus, relevant for cancer research and treatment.

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Keywords: pájaro bobo; native plants; ethnopharmacology; oncology; herbal medicines; complementary medicine

Resumen

Pertenciente a la familia Asteraceae, *Tessaria* (Ruiz & Pavon, 1753) es un género de arbustos o árboles pequeños distribuidos en diversos hábitats ubicados desde Argentina hasta el suroeste de los Estados Unidos de América. El taxón está compuesto por cinco especies confirmadas, las cuales han sido previamente reportadas por sus usos etnofarmacológicos como hipocolesterolemiante, hipoglucemiante, antitussivo, antiinflamatorio, anticanceroso y abortivo; así como en enfermedades hepáticas, renales, pulmonares y reumáticas. Hasta el momento, se han analizado químicamente *T. absinthioides*, *T. ambigua*, *T. dodoneifolia*, *T. fastigiata* y *T. integrifolia*. Mientras que las decocciones, infusiones, extractos metanólicos e hidrometanólicos son los compuestos botánicos más estudiados; sus constituyentes fitoquímicos más reconocidos son el ácido cafeoilquínico, eupatorina, naringenina, protocatchuic y quercetina. Científicamente, se atribuyeron al género varias propiedades biomédicas como virucida, antibacteriano, leishmanicida, insecticida, gastroprotector, antiasmático, hipoglucemiante y antiaterogénico. En relación con la investigación oncológica, las sustancias químicas producidas por *Tessaria*, tienen actividades antitumorales que interfieren con la carcinogénesis, la proliferación celular, la metástasis y la angiogénesis. Además, los extractos naturales obtenidos de estas especies tienen actividades biológicas estrechamente relacionadas con el cáncer y actúan como antioxidantes y antiinflamatorios. En particular, se ha informado que *T. absinthioides* resulta citotóxico contra varias líneas de células cancerosas y actúa como compuesto antitumoral en modelos murinos. Debido a los antecedentes mencionados, se considera la especie *Tessaria* como infravalorada dentro del campo oncológico. El objetivo de la presente revisión sistemática es resumir la información disponible del género, relevante para la investigación y el tratamiento del cáncer.

Keywords: Pájaro bobo; plantas autóctonas; etnofarmacología; oncología; medicamentos herbarios; medicina complementaria

Resumo

Pertencente à família Asteraceae, a *Vernonia* (Ruiz & Pavon, 1753) é um gênero de arbustos ou pequenas árvores distribuídas em vários habitats localizados da Argentina ao sudoeste dos Estados Unidos da América. O táxon é composto por cinco espécies confirmadas, que foram previamente relatadas para seus usos etnofarmacológicos como hipocolesterolemiante, hipoglicêmico, antitussivo, anti-inflamatório, anticâncer e abortivo; assim como em doenças hepáticas, renais, pulmonares e reumáticas. Até agora, foram analisados quimicamente *T. absinthioides*, *T. ambigua*, *T. dodoneifolia*, *T. fastigiata* e *T. integrifolia*. Por outro lado, as decoções, infusões, extratos metanólicos e hidrometanólicos são os compostos botânicos mais estudados; seus constituintes fitoquímicos mais reconhecidos são: ácido cafeoilquínico, eupatorina, naringenina, protocatchuico e quercetina. Cientificamente, várias propriedades biomédicas foram atribuídas ao gênero como virucida, antibacteriana, leishmanicida, inseticida, gastroprotetor, antiasmática, hipoglicêmica e antiaterogênica. Com relação à pesquisa sobre câncer, os produtos químicos produzidos pela *Vernonia* têm atividades antitumorais que interferem na carcinogênese, proliferação celular, metástase e angiogênese. Além disso, extratos naturais obtidos dessas espécies têm atividades biológicas intimamente relacionadas com o câncer e atuam como antioxidantes e anti-inflamatórios. Em particular, *t. absinthioides* tem sido relatado como citotóxico contra várias linhas de células cancerosas e age como um composto antitumoral



em modelos murinos. Devido ao passado supracitado, a espécie *Vernonia* é considerada desvalorizada dentro do campo oncológico. O objetivo desta revisão sistemática é resumir as informações do gênero disponíveis relevantes para a pesquisa e o tratamento do câncer.

Keywords: Boldo baiano; plantas nativas; etnofarmacologia; oncologia; fitoterápicos; medicina complementar

Introduction

Cancer is a leading cause of death in the world. Despite synthetic drugs used in current therapies have improved patient prognosis, the toxicity and development of secondary resistance remain a serious concern for researchers (Qazi *et al.*, 2018). It is generally accepted that plant-derived products provide health-related benefits, specifically for the prevention and treatment of several diseases. Crude herbal or botanical preparations have shown promising utility for chronic illnesses such as oncologic disorders, diabetes, heart diseases, and neurodegenerative processes, among others. The herbal bioactive compounds can provoke changes in plasma nutrient availability, therefore, in the cell microenvironment. By this mode of action, botanicals became a valuable source of anticancer compounds, which can affect tumor growth by modifications induced in antitumor immune response, cancer cell proliferation, its survival, and spread (Kanarek *et al.*, 2020). The mentioned properties are supported by numerous preclinical studies, which provide ample evidence that botanicals regulate multiple cancer hallmark pathways, including cell cycle, apoptosis, angiogenesis, invasion, and metastasis (Puccinelli & Stan, 2017).

In addition, these natural compounds can also interfere with the early stages of tumor development, acting as chemopreventive agents (Butt *et al.*, 2013). Carcinogenesis is a

biological process hallmarked by its diffuse and multifocal presence, with high statistical chances to progress to malignancy, altered DNA synthesis, and changes in cell protection mechanisms (Ryan & Faupel-Badger, 2016). This process, in which the normal cell is transformed into a cancer cell, is commonly divided into 3 different stages: initiation, promotion, and progression (Weinstein *et al.*, 1984). Among an increased list of chemical and environmental carcinogens, inflammation is a factor widely related to neoplastic transformation. In the interplay between inflammation and cancer, reactive oxygen/nitrogen species, cytokines and prostaglandins act as promoters of carcinogenesis by induction of DNA damage. The interference with the inflammatory process and its intermediaries will reduce the tumor transformation, neoplastic progression, and the development of metastases and recurrences (Piotrowski *et al.*, 2020). For this reason, anti-inflammatory and anti-oxidant properties of botanic extracts could represent a crucial aid for the prevention of carcinogenesis mediated by the inflammatory processes (Serrano *et al.*, 2018).

The plant kingdom represents an endless supply of bioactive compounds with potential activity to control diseases. In particular, the Asteraceae family includes more than 20 thousand species around the world and is considered the most evolved botanical family due to the floral structure and its chemical composition. In America, it is within the most important families of



plants in relation to the number of species reported with medicinal properties (Thomas *et al.*, 2009). *Tessaria* (Ruiz & Pavon, 1753) is a genus of the Asteraceae family, sometimes considered as *Pluchea*, composed of up to 17 proposed species. It is distributed from the southwest of the United States to Argentina; including Brazil, Bolivia, Chile, Colombia, Costa Rica, Ecuador, Panamá, Paraguay, Perú, Uruguay, and Venezuela (Tropicos.org, 1982). Currently, there are only 5 accepted and confirmed species: *T. absinthioides* (Hook. & Arn.) DC.; *T. ambigua* DC.; *T. dodoneifolia* (Hook. & Arn.) Cabrera; *T. fastigiata* (Griseb.) Cabrera and *T. integrifolia* Ruiz & Pav (The Plant List, 2013). All these species are reported in the bibliography by their health care implications and, often, scientific studies demonstrate novel valuable biological properties. Because of the mentioned antecedents, considering the *Tessaria* species as undervalued within the oncologic field, the goal of the current systematic review is to summarize the available genus information relevant to cancer research and treatment.

Material and Methods

A literature search on the *Tessaria* genus was performed to identify texts describing species with ethnopharmacological reports of use and other papers related to its scientific studies. The current review includes 77 references selected from websites (The plant list and Tropicos.org) and scientific databases as PubMed, Science Direct, SciELO, Google Scholar, LILACS, and Library Genesis. The main descriptors used were *Tessaria* and the constitutive genus species (especially, *T. absinthioides*, *T. ambigua*, *T. dodoneifolia*, *T. fastigiata*, and *T. integrifolia*), cancer, cytotoxicity, antitumoral,

carcinogenesis, tumor growth, metastasis, angiogenesis, and, finally, other words related to the chemical characterization as phytochemicals, phenolic compounds, and all the specific compounds mentioned in Table 3. Altogether, more than 4500 articles and documents were reviewed.

Distribution and Ethnopharmacological Uses of *Tessaria spp.*

Cancer is a group of diseases that represent a worldwide problem and, often, conventional therapy is limited by the cost and side effects of used drugs. With few exceptions, folk medicine has not reported information about cancer diagnosis and treatment. Because of this, exploring plants with registered ethnobotanical properties resulted an important strategy to find effective natural products for oncologic purposes. For this reason, during the last decades, many plants with reports of ethnomedicinal use were studied to develop anticancer plant-based drugs with improved potency and better tolerance by the patients (Tariq *et al.*, 2017).

Because the natural distribution of *Tessaria spp.* includes regions and cultures of Mesoamerica and South America, the ethnopharmacological properties of the species are widely registered (Table 1). Among others, there are reports of biological actions related to inflammation, cell proliferation, immune system response, and liver and kidney protection. Because of the relation between these properties and the tumor growth or treatment toxicity, the mentioned attributes make the genus a valuable candidate for plant-based cancer research.

Table 1
Ethnopharmacological Information of Tessaria Tessaria spp.

<i>Species</i>	<i>Popular names</i>	<i>Ethnopharmacological uses</i>	<i>Product</i>	<i>Geographical distribution</i>	<i>Ref.</i>
T. absinthioides	<i>Pájaro bobo, chilca, suncho rosado, suncho negro, brea Sorona, brea</i>	<i>Hypocholesterolemiant, balsamic, expectorant Diabetes</i>	<i>Dried leaves infusion</i> <i>Leaves infusion</i>	<i>Argentina</i> <i>Chile</i>	<i>Barboza et al., 2009</i> <i>Madaleno & Delatorre-Herrera, 2013</i>
	<i>Pájaro bobo, suncho negro</i>	<i>Empacho (digestive disorder)</i>	<i>Leaves infusion</i>	<i>Argentina</i>	<i>Campos-Navarro & Scarpa, 2013</i>
	<i>Pájaro bobo, Sorona, hierba de zorra</i>	<i>Rheumatism, prostate illness, cancer,</i>	<i>Leaves infusion</i>	<i>Perú, Bolivia, Chile, Argentina</i>	<i>Torres-Carro et al., 2017</i>
T. ambigua	<i>Pájaro bovo</i>	<i>Antitussive, hepatic, tonic, depurative, laxative</i>	<i>Part not specified</i>	<i>Argentina</i>	<i>Barboza et al., 2009</i>
T. dodonefolia	<i>hierba dulce, ka'a he'é</i>	<i>sweetening</i>	<i>Young shoots</i>	<i>Paraguay</i>	<i>Nanayakkara et al., 1988</i>
	<i>Chilca dulce, suncho, chilca, chilca negra, suncho negro</i>	<i>abortifacient, vaginal mycosis, anuria, urin with blood, emmenagogue</i>	<i>Leaf</i>	<i>Argentina</i>	<i>Barboza et al., 2009</i>
T. fastigiata	<i>Uri uri</i>	<i>antiinflammatory</i>	<i>Part not specified</i>	<i>Bolivia</i>	<i>Parejo et al., 2003</i>
T. integrifolia	<i>Pájaro bobo</i>	<i>hepatic and renal insufficiency, hepatitis.</i>	<i>Leaves</i>	<i>Perú</i>	<i>Feo et al., 1990</i>
	<i>Aliso del río, aliso, aliso bobo, bobo, buibé, pájaro bobo, palo bobo</i>	<i>Diuretic, asthma, febrifuge, astringent, cicatrizant</i>	<i>Aereal parts infusion</i>	<i>Argentina</i>	<i>Barboza et al., 2009</i>
		<i>Antigonorrheal, anti allergic, antiasthmatic, antiinflammatory, diuretic</i>	<i>Leaf and flowers</i>	<i>Argentina, Peru</i>	<i>Barboza et al., 2009, Peluso, et al., 1995</i>
	<i>---</i> <i>pájaro bobo, huapariu, tseco</i>	<i>Malnutrición</i> <i>asthma, antipiretic, antiinflammatory, diuretic</i>	<i>All plant</i> <i>Part not specified</i>	<i>Bolivia</i> <i>Perú</i>	<i>Feo et al., 1990</i> <i>Silva-Correa et al., 2018</i>

Note: derived from research.





Reported Biomedical Properties Of *Tessaria* Species Different From Cancer

Considering that cancer is a very heterogeneous group of diseases, it is not possible to predict when future discoveries will report other activities of *Tessaria spp.* relevant for oncology. For this reason, Table 2 summarizes the available information about other biomedical, scientifically tested properties of the genus.

Antioncologic Effects Of *Tessaria spp.*

Cancer is caused by a multistep process that results in uncontrolled cell proliferation. The available information about *Tessaria* species effects concerning cancer establishment and progression is schematized in Figure 1. To the present, only 4 species have been scientifically studied for biological activities with oncological relevance; in these

cases, the natural compounds demonstrated anticarcinogenic, cytotoxic, and antitumoral properties. According to our knowledge, *T. ambigua* has no previous reports of biological properties related to cancer.

The Role of the *Tessaria* Genus in Carcinogenesis

In some opportunities, the carcinogenesis process is driven by oxidative stress and inflammation, which determines the aberrant gene expression of tumor cells and those sites surrounding the lesion (Lechner & Stoner, 2019). As with other plants, *Tessaria* species are a source of phenolic compounds reported as natural antioxidants with clear inhibitory effects of carcinogenesis.

T. absinthioides is the most studied specimen because of its anti-inflammatory and antioxidant properties. Torres Carro *et al.* (2015; 2017) evidenced the capability of the hydromethanolic extracts obtained from the plant aerial parts to interfere with the inflammation process. Extracts act by a reduction in

Table 2
Biomedical properties of Tessaria spp. not cancer related.

Specie	Activity	Plant source	Cite
<i>T.abs.</i>	Insecticidal and repelent	Sesquiterpenes from aerial parts.	García <i>et al.</i> , 2003; García <i>et al.</i> , 2017
	Virucidal	Essential oils from leaves.	García <i>et al.</i> , 2003
		Organic extract (dichloromethane: methanol)	Visintini Jaime <i>et al.</i> , 2013
<i>T.amb.</i>	Gastric cytoprotection	Sesquiterpenes from aerial parts.	Donadel <i>et al.</i> , 2005
	Antibacterial	Methanolic extract from leaves	Romero <i>et al.</i> , 2016
	Hypoglycemic and antiatherogenic	Aqueous extract from leaves	Quesada <i>et al.</i> , 2021
<i>T.dod.</i>	Insecticidal	Penduletin from aerial parts	Sosa <i>et al.</i> , 2000
<i>T.fas.</i>	Antifungal	Flavonones from aerial parts	Soberón <i>et al.</i> , 2020
<i>T.int.</i>	Antiasthmatic, analgesic, immunomodulatory	Casticin from leaves	Chan <i>et al.</i> , 2018
	Antispasmodic	Aqueous extract from aerial parts	Silva-Correa, 2011
	Gastric cytoprotection	Ethanollic extract from leaves	Correa <i>et al.</i> , 2014
	Leishmanicidal	Sesquiterpenes from leaves	Silva-Correa <i>et al.</i> , 2018

T. abs.: *T. absinthioides*; *T. amb.*: *T. ambigua*; *T. dod.*: *T. dodoneifolia*; *T. fas.*: *T. fastigiata*; *T. int.*: *T. integrifolia*.

Note: derived from research.

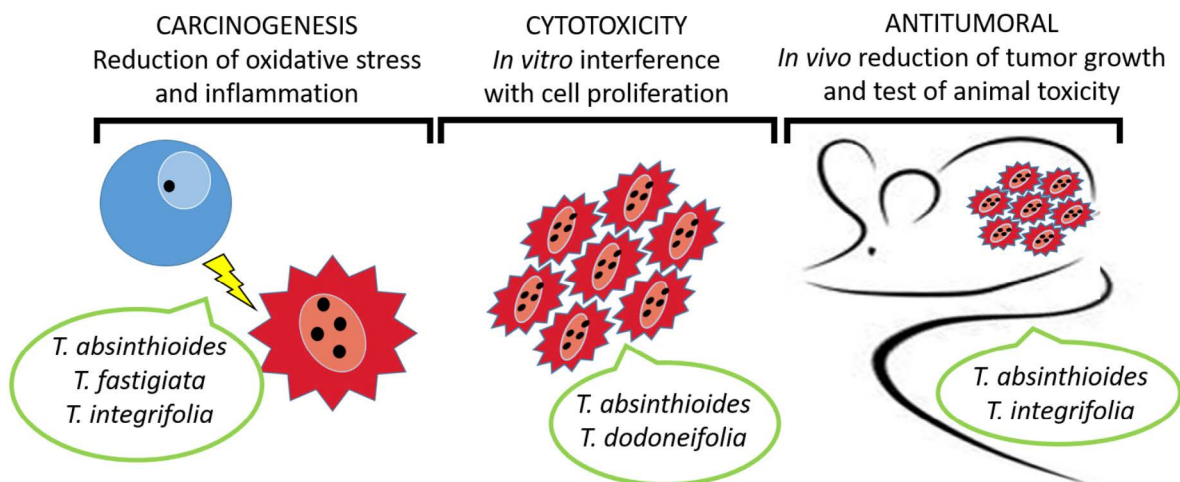


Figure 1. Antioncologic Effects of *Tessaria* spp.
Note: derived from research.

the activity of the pro-inflammatory enzymes lipoxygenase (LOX), cyclooxygenase (COX-2), secretory phospholipase A2 (PLA2), and hyaluronidase. Moreover, diminished production of nitric oxide (NO) by a reduction in the activity of nitric oxide synthase enzyme (iNOS) and the stabilization of human red blood cells membrane was also demonstrated. In the same studies, the antioxidant capability of hydromethanolic extract was established by determining the iron-chelating capacity and the 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid) (ABTS) radical scavenging effect. Later, Gómez *et al.* (2019) confirmed the antioxidant power of Argentinian and Chilean *T. absinthioides* decoctions by 2,2-diphenyl picrylhydrazyl (DPPH) scavenging activity, ferric reducing antioxidant power assay (FRAP), Trolox Equivalent Antioxidant Activity (TEAC), and reduction of lipid peroxidation in human erythrocytes. The above-mentioned biological properties were attributed to the presence of caffeoylquinic acid derivatives, as well as vanillic acid, protocatechuic, taxifolin, chlorogenic acid, quercetin, and rutin, among other phenolic compounds evidenced in the extracts.

In *T. integrifolia*, the antioxidant and anti-inflammatory properties were demonstrated by studying changes in the migration and the superoxide anion secretion of activated human macrophages. In this work, the biological properties were also related to the caffeoylquinic acid present in the aerial parts of the plant (Peluso *et al.*, 1995). Later, Ono *et al.* (2000) described the antioxidants effects of eudesmane derivatives present in the methanolic extract obtained from leaves, evidenced by the ferric thiocyanate method and DPPH scavenging assay.

T. fastigiata is the other species in the genus reported as an antioxidant. The unique available study used DPPH scavenging activity, nitroblue tetrazolium (NBT)/hypoxanthine superoxide assay, and the hydroxyl radical scavenging activity to evidence the biological activity attributable to the phenolic compounds present in the leaves hydromethanolic extract.

Altogether, the presented evidence makes *Tessaria* plants a very important source of natural compounds to prevent biological oxidations and inflammation, reducing the impact of both key factors of carcinogenesis.



Cytotoxicity of *Tessaria* Genus Against *in Vitro* Cells

In recent times, an important scientific effort was focused on discovering novel, effective, and affordable anticancer agents from natural sources. Because of this, a large number of botanicals have been explored for their cytotoxic potential against *in vitro* models of cancer cells (Dehelean *et al.*, 2021). In this section, the available evidence of cytotoxicity is analyzed in relation to the capability of make interference in cancer cell metabolism, the inhibition of cell proliferation, and the cell death induction.

Between the reviewed *Tessaria* species, the evidence clearly indicates that botanicals exert a selective cytotoxic effect against cancer cells. By these results it was demonstrated that the treatment with natural compounds of normal, non-tumoral cells induces lower or null toxicity; in contrast, when cancer cell lines were assayed, the cytotoxic effects were potent, similar to those evidenced by the conventional chemotherapeutic drugs used as a positive control.

Persia *et al.* (2017) demonstrated the selective cytotoxic effects of *T. absinthioides* leaves aqueous extract. The study reported a dose-response effect on HeLa (cervix cancer), Gli-36 (glioblastoma), HCT-116 (colorectal cancer) and MCF-7 (breast cancer) human cancer cell lines. The extract potency, measured as CV50 (50% of cell viability) in all cases, was similar to 5-fluoracile, a chemotherapeutic agent used as a positive control. Interestingly, in the same study and conditions, the cytotoxicity determined by the extract on non-tumoral HBL-100 cell line was significantly lower in relation to the other cancer cell lines tested; also, in these cell lines, the measured toxicity was notably diminished in relation to the effect induced by 5-fluoracile.

In another study, eudesmane semi-synthetic derivatives from *T. absinthioides* affected the proliferation of A2780 (ovarian), HeLa (cervix), SW1573 (non-small cell lung), T47D (breast) and WiDr (colorectal) human solid tumor cell lines. In a dose-response experimental design, the treatment determined metabolic cytotoxicity (measured by SRB assay) and cell cycle arrest in G2/M phase. In the study, 5-fluoracil was used as positive control, and its potency, measured as GI50 (50% of growth inhibition), was always lower than the plant-derived compound (León *et al.*, 2009).

There are, at least, other 3 works of *in vitro* studies related to the cytotoxicity of *Tessaria*'s compounds. In the course of these studies, not directly related to cancer research, were tested caffeoylquinic derivatives and flavonoids from *T. absinthioides* (Torres Carro *et al.*, 2015), caffeoylquinic derivatives from *T. integrifolia* (Peluso *et al.*, 1995) and the ethanolic extract of *T. dodoneifolia* with content of naringenin and pinocembrin (Soberón *et al.*, 2020). In all of these studies was reported none or slight toxicity induced by treatment on cultured non-tumoral murine macrophages and human peripheral blood lymphocytes (PBL).

The presented evidence in this section for the *Tessaria* derived compounds coincides with the observation of the selective cytotoxicity against cancer cells, originally proposed by Persia *et al.* (2017).

Preclinical Evidence About *Tessaria* sp. Antitumoral Effects

Often, the therapeutic potential of some natural products is limited by the presence of xenobiotics' effects. In other cases, the *in vitro* evidenced cytotoxic properties cannot be reproduced *in vivo* because of bioavailability limitations of the bioactive phytoconstituents



(Piroozmand *et al.*, 2020). In the case of *T. absinthioides*, both limitations were analyzed on the aqueous extract by the study of oral toxicity and by the determination of its antitumoral effects against colorectal-induced cancer. The *T. absinthioides* aqueous extract oral toxicity was tested and discarded in males and females of Sprague Dawley rats. At doses up to 2000 mg kg⁻¹, a single administration of *T. absinthioides* did not determine acute toxic effects. No animals died immediately or within 14 days after administration and were not evidenced changes of body weight nor other clinical signs of toxicity. After euthanasia, necropsy did not evidence changes on tissues or organs. On the other hand, the study of 28 oral repeated doses up to 1000 mg kg⁻¹ d⁻¹ did not show toxic evidence either. After administration, no significant changes were registered in body weight, organs weight or organs histological appearance. Neither changes were present in blood cell counting nor blood serum biochemistry (Persia *et al.*, 2017).

T. integrifolia inflorescences infusion was also tested by its oral toxicity in the 28 days repeated doses experimental design. By the use of *Rattus norvegicus* var. albina, the study concluded that no significant toxic effects were observed in the males or females analyzed. After administration of 500 mg kg⁻¹ day⁻¹ doses, the histopathologic analysis showed neither cell damage nor necrosis in the liver, lungs, stomach, brain, ovary or testis. Only a mild to moderate glomerular congestion was evidenced and was attributable to the sesquiterpene lactones present in the plant sample. In conclusion, the study demonstrated that there was not significant toxicity determined by oral administration of *T. integrifolia* during 28 days and for the long term administration of the infusion, specific studies need to be performed to discard

potential kidney damages (Julián Dávalos & Vásquez Muñoz, 2016).

About the *in vivo* antitumoral effects, *T. absinthioides* aqueous extract was tested in a colorectal cancer model induced by dimethylhydrazine (DMH). In BALC/c mice, the oral administration of 300 mg animal⁻¹ day⁻¹ significantly increased the median survival of animals. While the median survival in *Tessaria* treated animals was 30 weeks, in the untreated group survival was significantly lower (24 weeks). It is important to note that 5-fluoracil was used as a positive control drug; in this group, the median survival was 27 weeks. In spite of the fact that no statistical differences were observed between the survival of *Tessaria* and 5-fluoracil groups, the *in vivo* results confirm the similar potency evidenced by both compounds *in vitro*. To finish, animals treated with *Tessaria* did not evidence toxic symptoms related to the oral administration of the extract during the entire assay (up to 38 weeks) (Persia *et al.*, 2017).

Only preliminary evidence exists about *T. absinthioides* efficiency against *in situ* and metastatic murine syngeneic melanomas (personal observations); until now, no other *Tessaria* species were reported in the bibliography by its antitumoral properties.

***Tessaria* Genus as the Origin of Anticancer Phytochemicals**

The phytochemicals are bioactive non-nutrient vegetal compounds that have health-related effects. More than 5,000 phytochemicals have been identified; if well, their health benefits are still to be fully understood. Several studies have strongly demonstrated that phytochemicals have many different mechanisms of action related to cancer (Liu, 2004).

Table 3
Tessaria spp. Phytochemicals and Its Anticancer Properties.

Phytochemical	Source	Anticancer effect by target...				Ref
		carcinogenesis	tumor growth	metastasis	angiogenesis	
Amyrin	<i>T. amb.</i>		yes			Wen et al., 2018
Artemisinin	<i>T. abs.</i>	yes	yes		yes	Slezakova & Ruda-Kucerova, 2017
Caffeoylquinic acid and derivatives	<i>T. abs.</i>		yes	yes		In et al., 2016
Caryophyllene oxide	<i>T. abs.</i>	yes	yes			Fidy et al., 2016
Casticin	<i>T. abs.</i>	yes	yes	yes	yes	Ramchandani et al., 2020
Chrysosphanol	<i>T. int.</i>	yes	yes	yes		Hsu et al., 2020
Citric acid	<i>T. abs.</i>		yes			Ying et al., 2013
Enoic acid derivatives	<i>T. abs.</i>	yes	yes			Oliveira et al., 2018
Eriodictyol	<i>T. dod.</i>	yes	yes	yes		Li et al., 2020
Eudesmane derivatives	<i>T. abs.</i>	yes	yes	yes	yes	Liang et al., 2017
Eudesmol (gamma)	<i>T. abs.</i>		yes			Furtado et al., 2018
Eugenol	<i>T. int.</i>		yes			Fathy et al., 2019
Eupatorin	<i>T. abs.</i>		yes	yes	yes	Razak et al., 2019
Gallic acid	<i>T. int.</i>		yes			Rezaei-Seresht et al., 2019
Ginnalin A	<i>T. abs.</i>	yes	yes			Bi et al., 2018
Gurjunene (alfa)	<i>T. abs.</i>	yes	yes			Yongram et al., 2019
Hyperyn	<i>T. int.</i>		yes			Li et al., 2012
Illicic acid	<i>T. abs.</i>		yes			León et al., 2009
Linalool	<i>T. int.</i>		yes			Pan & Zhang, 2019
Mannoheptulose	<i>T. abs.</i>		yes			Board, et al., 1995
Naringenin	<i>T. dod.</i>	yes	yes	yes	yes	Joshi, et al., 2018
Pinoresinol	<i>T. int.</i>		yes	yes		Ning et al., 2019
Protocatechuic	<i>T. abs.</i>		yes	yes		Deng et al., 2020
Quercetin	<i>T. dod.</i>	yes	yes	yes	yes	Tang et al., 2020
Rhamnetin	<i>T. abs.</i>	yes	yes			Lan et al., 2019
Sakuranetin	<i>T. dod.</i>	yes	yes			Stompor, 2020
Taxifolin	<i>T. abs.</i>	yes	yes		yes	Wang et al., 2020
Terpinen-4-ol	<i>T. abs.</i>		yes			Shapira et al., 2016
Tessaric acid and derivatives	<i>T. abs.</i>		yes			León et al., 2009
Trifolin	<i>T. int.</i>		yes			Kim et al., 2016
Vanillic acid	<i>T. abs.</i>	yes	yes		yes	Gong et al., 2019

T. abs.: *T. absinthioides*; *T. amb.*: *T. ambigua*; *T. dod.*: *T. dodoneifolia*; *T. fas.*: *T. fastigiata*; *T. int.*: *T. integrifolia*

Note: derived from research.





Table 3 presents the phytochemicals derived from *Tessaria spp.* with reported anticancer actions. Until now, the 5 confirmed species of the genus (*T. absinthioides*, *T. ambigua*, *T. dodonaeifolia*, *T. fastigiata*, and *T. integrifolia*), were chemically analyzed and their phytochemical constituents described (Torres-Carro *et al.*, 2017; García *et al.*, 2003a; Gómez *et al.*, 2019; Ono *et al.*, 2007; Guerreiro *et al.*, 1990; Bailac *et al.*, 1998; Caballero Palacios, 2014). By these studies, more than 30 phenolic compounds were identified in *Tessaria* with reported anticancer efficacy, including sesquiterpenes, flavonoids, phenolic acids, and lignans. All of these chemicals affect tumor growth, modifying proliferation or viability; while 15 compounds interfere with carcinogenesis, 10 reduce the metastasis process, and 8 decrease angiogenesis.

The above-mentioned phytochemicals and their scientifically proven effects make the genus *Tessaria* a valuable source of natural compounds for future cancer research and treatment.

Conclusions

Botanicals, nutraceuticals, and herbals are plant-derived materials with medical benefits that aim for disease prevention or treatment. They represent a particular promise for cancer prevention due to their efficacy and safety profile. The wide chemical diversity features together with available epidemiological, preclinical, and clinical studies suggest an undeniable role of natural products in various approaches related to cancer prevention and treatment. Many of these natural compounds are responsible for antioxidant, anti-inflammatory, chemopreventive, and anticancer activities.

Some botanical constituents as polyphenols, phytoalexins, carotenoids, and flavonoids are specifically related to the expression and activity of multiple proteins such as epidermal growth factor receptor (EGFR), nuclear factor-kappa B (NF- κ B), tumor necrosis factor-alpha (TNF- α), protein kinase B (PKB/AKT), mitogen-activated protein kinase (MAPK) and p53, between others targets. By the modification of these molecular targets, botanicals play a critical role in signal transduction pathways related to carcinogenesis, cell proliferation inhibition, invasion, and angiogenesis (Rahman *et al.*, 2010).

Due to these important properties, it is urgent to perform a scientific validation of regional medicinal plants to precise their toxicological and pharmacological profiles with the goal of ensuring both the effectiveness and the safety in the use of ethnobotanicals. This information is highly valuable and necessary to improve the therapeutic approach of pathologies with unsatisfactory or toxic treatments, mainly cancer. In this field, the botanical complementary treatments rise as a promissory area to improve the potency of available therapies and/or reduce their toxic collateral effects.

The present review systematically summarizes the information available for the *Tessaria* genus related to cancer research and treatment. Then, it is imperative to move forward to complete the preclinical evidence related to its molecular mechanistic mode of action, pharmaceutical presentation and standardization, and the study of pharmacological interactions with current chemotherapeutics. In accordance with the presented evidence, based on its ethnopharmacological reports, biomedical explored properties, and phytochemical composition, it is possible to affirm that *Tessaria*



spp represent a promissory source of botanicals for oncologic purposes, especially in the complementary treatment approach. To conclude, based on the folkloric reports of uses and the recently validated scientific information, the present revision intends to encourage new and deep research destined to promote the *Tessaria* derived botanicals as anticancer compounds.

Conflict of Interest

The authors declare no competing interests.

Author Contribution Statement

All the authors declare that the final version of this paper was read and approved.

The total contribution percentage for the conceptualization, preparation, and correction of this paper was as follows: A.S.L. 25 %, M.B.H 25 % and C.G.L. 50 %.

Data Availability Statement

Data sharing is not applicable since no new data was created or analyzed in this study.

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A Systematic Review About the Contribution of The Genus *Tessaria* (Asteraceae) To Cancer Study and Treatment (Arianna Sosa-Lochedino • María Belén Hapon • Carlos Gamarra-Luques)

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