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Modulation of Secondary Metabolites among Mexican Medicinal Plants by Using Elicitors and Biotechnology Techniques

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Abstract

Medicinal plants are being utilized as raw material and the use has increased in recent decades due that these biosynthesize compounds with several pharmacological activities. Some plant species with biological potential are of interest to the industry for preparation of drugs, phytodrugs, or food supplements. This causes overexploitation and deforestation, which endangers plant species-of-interest. In recent years, alternatives have been sought to eradicate this problem. A solution that was give and is maintained is plant biotechnology, which favors the production of active Secondary Metabolites (SMt). Plant biotechnology allows us to increase the yield of a compound-of-interest, reduces its production times and costs, and allows constant and controlled production of the raw material, and while aiding in the protection of medicinal plants that are found in danger of extinction. In the scientific literature, procuring the SMt by means of biotechnological processes is described, highlighting the study of four species from Mexican traditional medicine (*Lopezia racemosa*, *Galphimia glauca*, *Cnidoscolus chayamansa*, *Sphaeralceae angustifolia* and *Buddleja cordata*), and the main biological activities are as follows: anti-inflammatory, hepatoprotector, neuroprotector, anxiolytic, antitumoral, antibacterial, and antioxidant, among others.

Keywords: Elicitors, biotechnology, Mexican medicinal plants, plant tissue culture, secondary metabolites, phenolic compounds

1. Introduction

Ever since our ancestors, humanity has been dependent on the consumption of plants as a source of food, health, and for construction/ornamental. In addition, plants have developed a complex defense system against biotic and antibiotic stress: therefore, they can produce diverse secondary metabolites (SMt). The stress to which plants are submitted under natural conditions is caused by different factors, among

which stand out: attack by diverse insects and/or microorganisms (viruses, bacteria, and fungi) competition for soil, light, and nutrients, and exposure to sunlight [1].

SMt are compounds that do not play a fundamental role in the vital processes of plants, but they are important as mechanisms of defense. They are responsible for organoleptic and protective properties, such as odor, flavor, color, and consistency. These SMt also act as chemoattractants or chemorepellents. In addition, they are of great interest in industry for the preparation of food additives, agrochemicals, essences, biodiesel, narcotics, insecticides, cosmetics, and aromatics, and one of the most important of these is for the production of substances with pharmaceutical interest. Frequently, the production of SMt wild-collected plant is very low (less than 1% of the plant's Dry Weight -DW-), and this depends specifically on the plant's physiological state, the geographic location, the climate, among other factors [2].

Due to the low yield of SMt in wild plants and considering its important biological activity, alternatives or tools are currently being sought to increase its yield. One of these alternatives is the application of several biotechnology processes, a discipline that is oriented toward the development and innovation of technologies that involve the management of biological material for the production of a good or service [3].

One of the advantages in the use of biotechnological processes is to increase the production of bioactive SMt and also reduce the production time, which favors their availability [4]. The purpose of this paper is to summarize all the information that exists on the use of biotechnological processes for the production of bioactive compounds from Mexican medicinal plants.

2. Products with pharmaceutical importance

Plants constitute a huge reservoir of chemical structures, the most economically important are medicinal plants, due to their diverse biological activities; which over the years have favored human survival thanks to their use in Traditional Medicine (TM) [5–7]. TM is widely used in some developing countries, where their health system is still growing and is of great economic importance. In Africa, up to 80% of the population employs TM to help satisfy its health needs. In Asia and in Latin America, the populations continue to use TM because of historical circumstances and cultural beliefs. In China, TM is of great importance due to the large percentage of population that utilizes it, being higher than 60%. In some developed countries, the percentage of the population that uses TM is 48% in Australia, 70% in Canada, 42% in the USA, 38% in Belgium and 75% in France [8].

Currently, Medicinal Plants (MP) are employed by 80% of the world population; therefore, these are overexploited not only because are source of active ingredients, also due to the high nutritional, wood, cosmetic, agricultural, and/or medicinal value that many of these have. For example, it is estimate that China exports 120,000 tons of MP and India, some 32,000 tons while Europe imports 400,000 tons of MP. This leads to overexploitation of the species and many of them are in danger of extinction [9, 10].

The World Conservation Union and the World Nature Fund report that there are between 350,000 and 550,000 species of MP in the world, of which only approximately 20% possess documented investigation of their biological potential, and nearly 15,000 species are in danger of extinction due to the overexploitation and destruction of habitats [10, 11].

Nowadays, scientific interest in MP has increased due to the high costs and adverse effects that allopathic drugs cause, in addition to the increasing appearance of strains of microorganisms that are resistant to current treatments [12–14]. It is noteworthy that almost 25% of the active principles of allopathic drugs currently used were isolated and/or semisynthesized from plants [9]. In modern medicine, digoxin is used as a cardiotonic and was isolated from *Digitalis purpurea* (purpura, its common name); escin is used as an anti-inflammatory and venotonic and was isolated from *Aesculus hippocastanum* (its common name, horse chestnut). Another compound utilized is ajmalicin, employed for circulatory disorders, and was isolated from *Rauwolfia serpentina*; paclitaxel (an anticancer drug) has been semisynthesized by Bristol-Myers Squibb since 2002, and was obtained from the compound 10-deacetylbatatin III, it was obtained from the cell suspension of the *Taxus baccata*. While diosgenin, a steroidal sapogenin, was obtained from the tubercles of several *Dioscorea* species, which was the raw material for the semisynthesis of progesterone [15].

Guanidine is a natural product with good hypoglycemic activity that was isolated from *Galega officinalis* (L); however, this compound has been reported to be toxic for human consumption. Therefore, this compound was semi-synthesized obtaining metformin (dimethylbiguanide), which is less toxic and has a pharmacologic effect similar to the original molecule and is widely used for the treatment of type II diabetes mellitus. It is worth noting, due to the high demand for SMt on the market; several companies have seen the need to discover novel sources of raw material from MP [16].

On the other hand, at present, the use of medicinal plants and/or phytodrugs is very frequent. The phytodrugs are elaborate with plant material and some derivatives of this. The main ingredient is the aerial or subterranean plant's part; as well as extracts, tinctures, juices, resins, fatty acids, and essential oils presented in pharmaceutical form. The therapeutic effectiveness and safety have been confirmed scientifically [17]. Some examples of these include ginseng, it is obtained from Panax genus (*Panax ginseng* and *P. quinquefolium*) native from Asia and America, respectively. The main biological effect of ginseng "tonic" phytodrug is that it possesses the ability to increase the capacity to tolerate tensions, which leads to increased mental and physical yield. Another phytodrug obtained from St. John's wort (*Hypericum perforatum*) is Hiperikan, which is standardized based on its content of hypericin; its principal pharmaceutical use is against depression. *Ginkgo biloba* (Ginkgo) belongs to the Ginkgoaceae family, the active compounds in the leaf's extracts are gingolides (gingolides A-C, J, and M), along with a mix of sesquiterpene lactones and flavonoids which is used against depression. The majority of commercial preparations from Ginkgo are standardized with approximately 5–7% of terpenic lactones and 22–27% of flavonoids and they are employed mainly for the treatment of the cognitive deterioration associated with alterations in blood circulation in the brain, such as dementia. The phytodrug elaborated with *Echinacea purpurea* is commercially known as EchinaCold (Schwabe Pharma) or Immulone (ATOS Pharma). These are standardized on based of the echinacosides (caffeic acid derivative) content, whose main biological effect is as an immunostimulant [18]. In Oceania region, the extract from *Piper methysticum* (from root and rhizome) has the commercial name Kava-kava (with 30% of kava lactones), and is utilized for their neurotransmitter activity [19]. Another phytodrug is Vitango, obtained from *Rhodiola rosea* (with 3.5% rosavins and 1% salidroside), and it is employed for reducing the stress associated with physical and mental tasks [20]. Plantival has extract mixture from *Valeriana officinalis* (160 mg) and *Melissa officinalis* (80 mg) and is used in the treatment of nervousness, restlessness and insomnia as an anxiolytic and antidepressive [21]. Another phytodrug, known as

Prostasan, is the extract of *Serenoa repens*, standardized at 25% of fatty acids; the dose employed is 160 mg, and its principal effect is antiandrogenic and against benign prostatic hyperplasia [22].

Due to the acceptance and growing use of phytodrugs around the world, PM are raw materials of great attention due to high consumption. In addition, MP biosynthesize several bioactive compounds, which are classified as terpenoids, alkaloids, lactones, flavonoids, coumarin, lignans and phenols, among others; many of these have restrictive taxonomical distribution. Although the SMt functions are not directly associated with the plant's basic function, these compounds carry out some interaction roles in the plant and its environments such as: protection against pathogens, protection against abiotic tensions (ultraviolet radiation radiation), they possess the function of attracting pollinating insects, and they are signaling molecules and active ingredients for drugs [23–25].

It is estimated that around 50% of the drugs approved by the Federal Drug Administration are products derived from natural sources or analogs deriving from plants or microorganisms [26]. However, raw material can be limited, and its exploitation is one of the main ecological concerns. One of the key objectives of plant biotechnology is the development of large-scale production methods of pharmacologically active products. Additionally, the massive biosynthetic potential of plants has not been completely exploited yet and biotechnology can be employed to generate new chemical compounds that possess unknown biological activities and/or with a different mechanism of action, or a better one, than those in existence [23].

3. Production of SMt by biotechnology tools

There are distinct strategies to optimize the production and modulation SMt in medicinal plants and food. The main strategies are by uses the elicitors (molecules capable of inducing defense in the plant) [1], which are classified as biotic and abiotic. Biotics are of biological origin, while abiotics can be physical or chemical. Some examples of physical abiotics are the weather, bacteria, and plagues, among others, while chemical abiotics possess an intense variety, with those most utilized being jasmonic acid and salicylic acid [27, 28]. One of the advantages of using elicitors treatment is that they function as signaling compounds for the mechanisms of defense; thus, they increase the production of SMt in an effective and rapid manner [29]. There is great specify in the interaction of plant-elicitor species which implies that the adequate one for each culture, the time of adding it, and the concentration for obtaining best response should be selected [30].

There is other technique very used to obtain SMt *in vitro*, it focuses on obtaining the roots, which is known as “hairy roots” or transformed roots; for this, the bacterium *Agrobacterium rhizogenes* is very used. This microorganism transfers the plasmid of the Transfer-DNA (T-DNA) of the T-DNA to the plant cell, to verify whether a root transformation was obtained, this can be confirmed by Southern hybridation analysis (this technique permits the detection of a specific DNA sequence in a complex mixture). A main advantage of these is that they have the capacity of rapid growth without the external administration of Plant Growth Regulators (PGR); the majority of these do not require a light supply, and their yield of metabolites is constant due to their genetic stability [1]. Another internal factor is the culture medium added with macro- and micronutrients, as well the external factors, such as light intensity, temperature, humidity, and stirring speed [31].

In general, formulation of the culture medium begins with the base medium, being the most utilized Muashige & Skoog (MS), B5 of Gamborg and Linsmaier and Skoog (LS), and Nitsch and Nitsch (NN) [32]. These culture mediums contain minerals, vitamins, and a carbon source, normally sucrose and sometimes fructose is used. Although plant cell cultures typically are initiated in solid medium, they require liquid medium for production on a large scale. The mineral content and/or the carbon source in culture medium have a profound impact on biosynthesis of SMt employed in the manufacturing of phytodrugs and/or compound-of-pharmaceutical-interest [33].

Other tools very used to obtain SMt by biotechnological process is through the use of BioCatalyzers; this method has been used to transform polyphenols compounds; for example, *Bouvardia ternifolia* is utilized for the production of a BioC denominated dehydrodiisoeugenol, which was obtained from the supernatant of cells suspension, demonstrating a yield of around 77%. The dehydrodiisoeugenol obtained from *B. ternifolia* allows the production of isoeugenol by biotransformation; it is known that plant peroxidases transform phenols substituted for by a methyl group ortho position to the corresponding O-radical, which, on establishing itself by resonance, produces a C-radical; the latter is that which leads to dimerization, producing a dimer. This biotransformation represents a clean and green alternative with respect to traditional chemical methods, in which oxidative bonding reactions are affected using catalysts such as FeCl₃, K₃(FeCN)₆, and Cu(OH)Cl [34].

Recently, interest in research and development of *in vitro* plant tissue cultures from MP has grown; however, there are scarce studies, to our knowledge, in which the biological activities of these SMt obtained by this process are described. The majority of works published only mention the conditions of the biotechnological process and the final concentrations of the different metabolites produced, but do not evaluate the pharmacological activity of these SMt, and the authors solely cite that these have been reported in previous works.

In **Table 1** and **Figure 1**, some examples are described. It is important to mention that on some occasions is difficult to establish the biotechnological process conditions to induce the biosynthesis of bioactive SMt from a MP.

Species	Products	Use	References
<i>Alanthus altissima</i>	Alkaloids	Antimicrobial	[35]
<i>Ajuga reptans</i>	Antocyanins	Antioxidant	[36, 37]
<i>Alanthus altissima</i>	Cannabinoids alkaloids	Antimicrobial	[38]
<i>Ammi majus</i>	Coumarins	Anticoagulant	[39]
<i>Anchusa officinalis</i>	Rosmarinic acid	Antioxidant	[40, 41]
<i>Anthoceros agrestis</i>	Rosmarinic acid and Glycosides	Antioxidant	[42]
<i>Arachis hypogea</i>	Piceatannol	Antioxidant	[43]
<i>Artemisia annua</i>	Artemisinin	Antimalaric	[30, 44]
<i>Artemisia judaica</i>	Flavonoids	Antioxidant	[45]
<i>Bouvardia ternifolia</i>	Dehydrodiisoeugenol	Biocatalyst	[34]
<i>Beta vulgaris</i>	Betalains	Antioxidant	[46, 47]
<i>Buddleja cordata</i>	Verbascoside (1), linarin (2)	Anti-inflammatory, antioxidant	[48]

Species	Products	Use	References
<i>Buddleja cordata</i>	Phenylpropanoids	Antioxidant	[49]
<i>Caesalpinia pulcherrima</i>	Homo isoflavones	Antimicrobial, antitumoral	[50]
<i>Calophyllum inophyllum</i>	Dipyranocoumarins	AntiHIV	[51]
<i>Camelia sinensis</i>	Thiamine or theanine	Antihypertensive	[52]
<i>Capsicum frutescens</i>	Capsaicin (3)	Irritant	[53]
<i>Cassia acutifolia</i>	Antraquinones	Antimicrobial	[54]
<i>Castilleja tenuiflora</i>	Verbascoside (1), Isoverbascoside, aucubin	Anti-inflammatory, Antispasmodics	[55]
<i>Catharanthus roseus</i>	Ajmalicin	Antihypertensive	[56]
<i>Catharanthus roseus</i>	Vinblastin	Anticancer	[57]
<i>Cecropia obusifolia</i> <i>Cecropia peltata</i>	Chlorogenic acid, Isoorientin	Hypoglycemic	[58]
<i>Cephaelis ipecacuana</i>	Emetin	Antiparasitic	[59]
<i>Cephalo-taxus fortunei</i>	Abietane diterpenoids	Antitumoral	[60]
<i>Choisya ternata</i>	Furanocoumarins	Antitumoral, Antioxidant	[61]
<i>Choisya ternata</i>	Furoquinolin alkaloids	Antitumoral, Antimicrobial	[62]
<i>Cinchona robusta</i>	Robustaquiones	Antimalarial	[63]
<i>Cistanche deserticola</i>	Glycosides	Antioxidant	[64]
<i>Cistanche salsa</i>	Glucophenyletanoids	Aphrodisiac	[65]
<i>Colchium autumnale</i>	Colchicine (4)	Antitumoral	[66]
<i>Coleus forskolii</i>	Forskolin (5)	Asthma	[67]
<i>Comptotheca acuminata</i>	Camptotecin (6)	Antitumoral	[68]
<i>Coptis japonica</i>	Berberin (7)	Intestinal infection	[69]
<i>Cornus kousa</i>	β -glucogallin, (+)-Catechin, (+)-gallo catechin, procyanidin B-3	Hyperglycemic and antimicrobial	[70]
<i>Coscinium fenestratum</i>	Berberin	Antioxidant, Antidiabetic	[71]
<i>Crocus sativus</i>	Crocin	Anticancer	[72]
<i>Cynara cardunculus</i>	Cinarin, Chlorogenic acid	Antioxidant	[73]
<i>Daucus corata</i>	Antocyanins	Lipoperoxidation	[74]
<i>Digitalis lanata</i>	Digoxin (8)	Cardiostimulant	[75]
<i>Dioscorea deltoide</i>	Diosgenin (9)	Steroidal stimulant	[76]
<i>Drosophyllum lusitanicum</i>	Plumbagin (10)	Anticancer, Antimicrobial	[77]
<i>Eleutherococcus sessiliflorus</i>	Eleuteroside	Anti-inflammatory, diuretic, analgesic, antipyretic	[78]

Species	Products	Use	References
<i>Eriobotrya japonica</i>	Triterpenes	Anti-inflammatory, antidiabetic, antitumoral	[79]
<i>Eucommia ulmoides</i>	Chlorogenic acid (11)	Antimicrobial, Antioxidant	[80]
<i>Fagopyrum esculentum</i>	Rutin (12)	Antioxidant	[81]
<i>Fragaria ananassa</i>	Antocyanins	Antioxidant	[82]
<i>Galphimia glauca</i>	Galphimine B (27)	Central nervous system disorders	[83]
<i>Glehnia littoralis</i>	Antocyanins	Antioxidant	[84, 85]
<i>Gymnema sylvestre</i>	Gymnemicanor	Antidiabetic	[86]
<i>Helianthus tuberosus</i>	Inulin	Antidiabetic	[87]
<i>Hemidesmus indicus</i>	Rutin (12)	Antioxidant	[88]
<i>Hypericum perforatum</i>	Hypericin	Antidepressive	[89]
<i>Hyssopus officinalis</i>	Rosmarinic acid (13)	Antioxidant	[90]
<i>Hyssopus officinalis</i>	Lithospermic acid	Antioxidant	[91]
<i>Ipomoea batatas</i>	Antocyanins	Antioxidant	[92–94]
<i>Larrea divaricata</i>	Nordihydroguayaretic acid, Quercetin	Antiarthritic, digestive, against venereal diseases	[95]
<i>Lavandula vera</i>	Rosmarinic acid (13)	Hepatoprotective	[96, 97]
<i>Leucophyllun frutescens</i>	Coumarins, lactones, flavonids	Antioxidants	[98]
<i>Lithospermum erythrorhizon</i>	Shikonin	Antibacterial	[99]
<i>Lopezia racemosa</i>	6-O-palmitoyl-3-O-β-D-glucopyranosylcampesterol, 6-O-palmi-toyl-3-O-β-D-glucopyranosyl-β-sitosterol	Anti-inflammatory	[100]
<i>Lycium chinense</i>	Cerebroside	Cellular Growth Regulator	[101]
<i>Morinda elliptica</i>	Antraquinones	Antimicrobial	[102]
<i>Mucuna pruriens</i>	L-dihydroxyphenylalanine	AntiParkinson	[103]
<i>Ochrosia elliptica</i>	Elipticin	Antitumoral	[104]
<i>Ocimum basilium</i>	Rosmarinic acid (13)	Antioxidant	[105]
<i>Panax ginseng</i>	Ginkgolides (14)	Cognitive deterioration	[106]
<i>Panax ginseng</i>	Ginsenosides (15)	Immunomodulator	[107]
<i>Papaver somniferum</i>	Codeine (16)	Sedative	[108]
<i>Papaver somniferum</i>	Morphine (17)	Sedative	[109]
<i>Papaver somniferum</i>	Sanguinarin (18)	Platelet stimulator	[110]
<i>Passiflora quadrangularis</i>	Orientin, Isoorientin, Vitexin, Isovitexin	Antioxidant	[111]
<i>Petroselinum sativum</i>	Flavonolides	Antioxidant	[112]

Species	Products	Use	References
<i>Picrasma quassioides</i>	Cuasin	Antiphylogistic	[113]
<i>Piqueria trinerva</i>	Monoterpene	Antifungal	[114]
<i>Podophyllum hexandrum</i>	Podophylotoxin (19)	Antitumoral	[115]
<i>Psoralea corylifolia</i>	Genistein and Daidzein	Tonic	[116]
<i>Rauwolfia serpentina</i>	Reserpin (20)	Antihypertensive	[117]
<i>Rubia tinctorum</i>	Antraquin-ones	Antimicrobial	[118]
<i>Salvia miltiorrhiza</i>	Cryptotanshinone	Antioxidant, antimicrobial	[119]
<i>Salvia miltiorrhiza</i>	Tanshinone	Cardiac problems	[120]
<i>Saussurea medusa</i>	Hispiduline (21), Jaceosidine	Antitumoral	[121]
<i>Silybum marianum</i>	Silymarin	Hepatoprotective	[122]
<i>Solanum malacoxylon</i>	Cholecalcipherol	Aids calcium absorption	[123]
<i>Sphaeralcea angustifolia</i>	Sphaeralcic acid (22), Tomentin (23)	Antiinflammatory	[124]
<i>Swietenia humilis</i> Zucc.	Alkaloids	Cytotoxic	[125]
<i>Tanacetum parthenium</i>	Parthenolide (24)	Anticancer	[126]
<i>Taxus brevifolia</i>	Taxol (25)	Anticancer	[127]
<i>Vitis vinifera</i>	Antocyanins	Antioxidant	[128]
<i>Vitis vinifera</i>	Resveratrol (26)	Antioxidant, Hepatoprotective	[129]
<i>Withania somnifera</i>	Withanolide A	Antioxidant, Antistress	[130]

Table 1.
Secondary metabolites obtained for cellular cultures from medicinal plant tissues in vitro and their biological activity.

4. Anti-inflammatory activity of SMt isolated from vegetal material obtained by biotechnological processes

Some SMt with significant anti-inflammatory activity have been obtain from MP through employment some biotechnological processes. From cell suspension cultures *Sphaeralcea angustifolia*, two compounds with important anti-inflammatory activity (evaluated in murine models) were isolated. The cell suspension was developed in MS medium with total nitrate 2.74 mM, under this condition was obtained scopoletin, sphaeralcic acid (22) and tomentin (23). From the CH₂Cl₂:CH₃OH extract sphaeralcic acid (22) and tomentin (23) were isolated; these compounds showed 58 and 66% anti-inflammatory activity, in the carrageenin model at 45 mg/kg administered by intraperitoneal (i.p.) route. On the other hand, in the topical anti-inflammatory model (TPA, 12-O-TetradecanoylPhorbol-13-Acetate), tomentin (225 mM/ear) exhibited 57% inhibition in the formation of auricular edema, while sphaeralcic acid (174 mM/ear) revealed 86% inhibition with a dose-dependent effect and one half of the Effective Dose (ED₅₀) = 93 mM. Sphaeralcic acid is the most active compound in both models (topical as well as systemic) [124, 131].

Figure 1.
Chemical structure of some polyphenols and other SMT with biological activity.

In another study, the anti-inflammatory activity of the cell suspension culture from *S. angustifolia* is described. In this case, aseptic-leaf explants and Naphthalene Acetic Acid (NAA, such as auxin) in several concentrations (0, 0.5, 1.0, and 2.0 mg/L) in combination with a constant concentration of Kinetin (KIN) were used. For the cell suspension culture, they utilized 4% initial inoculum in MS medium with 2.74 mM of the total nitrates, 1 mg/L of NAA and 0.1 mg/L of KIN and supplemented with 30 g/L of sucrose. The main SMt identified in this suspension cultures were the same compounds (scopoletin, tomentin, and sphaeralcic acid). Scopoletin was excrete in the culture medium, although it also accumulated in the biomass. For evaluation of the anti-inflammatory activity, the authors prepared the CH₂Cl₂:MeOH extract of the cell's suspension from *S. angustifolia* and this extract was administered i.p. in male ICR mice (35 g) employing the carrageenin model. This extract showed ED₅₀ = 137.63 mg/kg; sphaeralcic aid and tomentin at 45 mg/kg inhibited 67 and 62%, respectively on carrageenan assay and sphaeralcic acid at 1 mg/ear was more active in TPA assay, showed ED₅₀ = 93 mM and tomentin showed 48% of inhibition at 1 mg/ear [132, 133]. In addition, the same extract from biomass of cells in suspension of *S. angustifolia* at 100 mg/kg (with 0.10 mg of scopoletin, 0.10 mg of tomentin and 0.19 mg of sphaerelcic acid), as well as tomentin (20 mg/kg) were active as anti-inflammatory agent and reduced the mean body weight lost in Freund adjuvant- and kaolin/carrageenan-induced arthritis, respectively. In this assay, the organic extract and tomentin reduced the levels of pro-inflammatory interleukins such as IL-1 β , IL-6 and TNF- α and increased levels of IL-4 and IL-10 (anti-inflammatory cytokines) [133].

In parallel with obtaining cells *in vitro* of *S. angustifolia*, the authors performed a preclinical phase study (in rats). The CH₂Cl₂ extract of the aerial parts of *S. angustifolia* (wild material) was tested in chronic inflammation model induced with complete Freund's adjuvant (polyarthritis) The administration of the extract at 100 mg/kg/day during 8 days showed sustained and significant inhibition of edema, being of 62.6% [134]. A double-blind clinical phase study with the extract of *S. angustifolia* (wild material) standardized at 1% hydroxycoumarin content was conducted; the experiment was performed on 130 patients diagnosed with osteoarthritis. 55 of them were treated with standardized extract of *S. angustifolia* (gel) and 75 patients were treated with Diclofenac (2%). The therapeutic effectiveness of the gel administered topically for 4 weeks was 89%, while that of the control group (Diclofenac) was 91.3%; it was highlighted in the study that patients who received the treatment (gel of the standardized extract) did not exhibit adverse effects and did show an improvement in their disorder [135].

Another plant utilized in Mexican ethnomedicine is *Lopezia racemosa* Cav. Callus cultures in MS medium were obtain with variable amounts of NAA, 2,4-Dichlorophenoxyacetic acid (2,4-D) and 6-BenzylAminoPurine (BAP). The authors carried out 10 treatments with the previously mentioned PGR. In this case, they employed three types of explants (hypocotyl, stem nodule, and leaf) and several treatments. The combination of 1.0 mg/L of 2,4-D plus 0.5 mg/L of BAP was the best. From these callus material two novel compounds: 6-O-palmitoyl-3-O- β -D-glucopyranosylcampesterol (174.0 μ g/g of biomass) and 6-O-palmitoyl-3-O- β -D-glucopyranosyl- β -sitosterol were isolated. When quantifying these compounds, the authors observed that the wild plant contains less quantity than the callus. The topical anti-inflammatory activity of the biomass obtained from the callus was evaluate in the TPA model on CD-1 male mice at 1 mg/ear. Three extracts (hexanic, CH₂Cl₂, and methanol), was tested and showed 48.74, 57.14, and 16. 81% of inhibition, respectively. The CH₂Cl₂ extract was the most active, with a half-maximal Inhibitory

Concentration (IC_{50}) = 0.93 mg/ear. On the other hand, the pure compound (6-O-palmitoyl-3-O- β -D-glucopyranosyl-campesterol) was tested in the same model at 1 mg/ear showing a 57.14% inhibition, with IC_{50} = 0.45 mg/ear [100].

The lipophilic extract containing beta-carotene (LMBC) from plant cell cultures of *Cleome spinosa* was evaluated in two *in vivo* models to determine the anti-inflammatory and antinociceptive activities in Swiss Webster (SW) mice of both sexes. The callus culture was obtained of the MS medium supplemented with 1 mg/L of 4- amino-3,5,6-trichloropicolinic acid (picloram) and sub-cultured to culture medium with the same composition at 4-week intervals. The anti-inflammatory activity in carrageenan model at 10 mg/kg by i.p. via was evaluated. LMBC was inactive with respect to extract from whole plant, which showed more than 50% inhibition of edema at the same dose. On the other hand, the LMBC (at 50 mg/kg) showed around 68% decrease in writhes, these data were very similar to that shown in wild plant, and the effect was better than dipyrone (at 100 mg/kg) used as positive control. The authors concluded that the results of LMBC are particularly important; since this active SMt of medicinal interest can be continuously obtain from callus cultures [136].

Buddleja cordata is other medicinal species utilized to treat diseases related with inflammation. This cell suspension was obtained in MS medium supplemented with NAA (9.05 μ M) and Kin (2.32 μ M). The anti-inflammatory activity of the extracts from wild plant and of the cell suspension cultures were describe. In both extracts, the verbascoside content was quantified by HPLC methods. The extract of the cells suspension has 87.48 mg verbascoside/g Dry Matter (DM), while that the same extract from wild plant only contained 47.34 mg of verbascoside/g DM. In addition, acute toxicity in Balb/C mice of the both extracts were also determined, with half of a Lethal Dose (LD_{50}) of >2 g/kg. On the other hand, the topical anti-inflammatory effect of the wild plant extract and of the cell suspension was assay. The ED_{50} values was 3.93 and 1.26 mg/ear, respectively, cell suspension extract was the most active due to its greater content of verbascoside. Evaluation of both extracts in the carrageenan model (systemic inflammation), showed ED_{50} = 251.26 and 204.62 mg/kg for wild plant and cells suspension extracts, respectively; in this case, the latter extract was more active. In the chronic inflammation model (the arthritis model induced with complete Freud's adjuvant), both extracts showed moderate anti-inflammatory activity (<35%) and favored weight increase in animals with arthritis. The authors concluded that the cell suspension culture of *Buddleja cordata* obtained through the biotechnological process contained a better anti-inflammatory activity; therefore, it represents a source for obtaining this type of secondary metabolite-of-pharmacological-interest [48].

Cnidocolus chayamansa is medicinal plant whit anti-inflammatory, antiprotozoal, hepatoprotective, hypoglycemic and antimycobacterial activities [137–139]. Recently, a biotechnology processes was described to obtain callus using BAP (5 mg/L) and 2,4-D (2.5 or 5 mg/L), this callus was used as a biotechnological alternative for *in vitro* propagation of this plant [140]. After that, this callus was use to establish a cell suspension culture. From the cell suspension, organic extract was prepared and its antioxidant, antibacterial and anti-inflammatory activities were determined, as well as the main SMt was quantified by HPLC analysis. In cell suspension, lupeol acetate (38.1 mg/g DW) was obtained as a main constituent and scopoletin (3.6 mg/g DW) was also quantified; in wild material, both compounds were isolated in low quantity. The organic extract was active against *Staphylococcus aureus*, *S. coagulase* and *Listeria monocytogenes*, and a moderate antioxidant and anti-inflammatory activities (in TPA and carrageenan models) showed [28, 141].

5. Antineoplastic activity of the plant material obtained by biotechnological cultures

From the callus culture of *Eriobotrya japonica*, nine triterpenes (ursolic acid; oleanolic acid; maslinic acid; tormentic acid; 2 α ,19 α -dihydroxy-3-oxo-urs-12-en-28-oic acid; 2 α -hydroxyursolic acid; hyptadienic acid, and the mixture of 3-O-*cis*-p-coumaroyl tormentic acid and 3-O-*trans*-p-coumaroyl tormentic acid) were isolated. The main triterpenes of the callus tissues were tormentic acid (50 mg/g DW) and 2 α ,19 α -dihydroxy-3-oxo-urs-12-en-28-oic acid (11.8 mg/g DW), the latter compound (2 α ,19 α -dihydroxy-3-oxo-urs-12-en-28-oic acid) is known as a potent protease inhibitor of the human immunodeficiency virus. All these triterpenes were tested in two cell lines (HSC-2 and HSC); seven of the nine triterpenes were active. Showing mean cytotoxic concentration (CC₅₀) between 10 and 48 μ g/ml, while the oleanolic acid and 2 α ,19 α -dihydroxy-3-oxo-urs-12-en-28-oic acid exhibited weak cytotoxic activity. Additionally, the authors evaluated the *in vivo* antitumor activity of the 2 α ,19 α -dihydroxy-3-oxo-urs-12-en-28-oic acid in female ICR mouse skin ($n = 15$) during two stages of carcinogenesis; in this assay, carcinogenesis was induced topically with (+)-(E)-4-methyl 2[(E)-hydroxyimino]-5-Nitro-6-methoxy-3-hexenamide (NOR1) at a dose of 90 μ g/0.1 mL of acetone. One week after NOR1 administration started, TPA (1 μ g/0.1 mL of acetone) was administered twice weekly, yielding as a result a weak inhibition of the carcinogenesis. On the other hand, the authors mention that 2 α ,19 α -dihydroxy-3-oxo-urs-12-en-28-oic acid is an antiproliferative agent and that the number of papillomas diminished by 40% in 20 weeks, indicating that this compound possesses potential for the delay of carcinogen in mouse skin [142].

6. Biological effect of SMt isolated from cell cultures of *Galphimia glauca*

Galphimia glauca is widely used in Mexican traditional medicine. From this species, some triterpenes such as Galphimine-A, B and E (27) have been isolated. These compounds showed a neuroprotective effect, when these were evaluated in mice convulsions model. To induce the seizure, the authors used strychnine or pentylene-tratrazole administered by i.p. or subcutaneous route. In the study's results, the depressor effects observed on motor activity directed toward an objective or an aim [143]. The pharmacological effect of galphimine B (G-B) was due to selectively inhibiting the discharge of dopaminergic neurons in the central area in *in-vivo* models [144]. Due to its therapeutic importance of G-B, the authors proceeded to induce the production of this homogeneous raw material through a biotechnological process.

A first step was to obtain callus from hypocotyl explants in MS medium for 30 days with a combination of NAA and KIN; under these conditions, only great cell growth was obtained, and with 2,4-D at 4 mg/L the G-B production was stimulated with a yield of 0.154 mg/g DW. In addition, under this condition, G-E was also obtained but at less concentration (0.057 mg/g DW). Also, friable callus from suspension culture in MS medium with NAA and KIN (2:2 mg/L) was obtain, denominating this line as ggxl. By means of a growth kinetic, galphimines were shown to be produced in the culture's stationary stage [83, 145]. The next step was to carry out the scaling of galphimine production in the 5-liter *airlift* bioreactor and in one with mechanical stirring; the growth indices were 11.66 and 1.7, respectively. However, the authors observed that neither the biomass production, nor the time exerted an influence on the yield of G-B. Because the *airlift* produced a greater biomass but with lower yield of G-B (255 mg/L), while the stirring bioreactor at day 10 shown an intracellular as well as an extracellular

content of 1381 mg of G-B/L, 5.4-times higher than the *airlift* at day 25 [146]. Once the biotechnological conditions for the production of G-B were established, this allowed having raw material to carry out the pharmacological evaluation in different models.

7. Toxicologic effect of Galphimines

Aqueous extract from material obtained by bioreactor was prepared, whose galphamine content was G-A, G-B and G-E = 0.6, 1.034 mg/g, and 1.12 mg/g, respectively. Meanwhile, the content of these galphamine in the ethanolic extract was G-A = 5.35 mg/g, G-B = 18.8 mg/g, and G-E = 17.49 mg/g and the MeOH extract content G-A = 7.29 mg/g, G-B = 17.47 mg/g, and G-E = 11.6 mg/g. Afterward, each extract was administered to Balb/C male and female mice for 28 days (2.5 g/kg). During the study period, there were no deaths, and in the histopathological analysis of the different organs; the latter did not present alterations. Also, analyzed the behavioral parameters, demonstrating a reduction in spontaneous activity. Administration of these extracts for 56 days (2.5 g/kg) in mice did not cause any change in liver-function biochemical parameters. With regard to the cytotoxic evaluation in KB, UISO, and OVCAR-5 cell lines, no cytotoxic effects were found, but all of these extracts specifically inhibited growth of the colon-cancer cell line with ED₅₀ of <2 µg/mL. On the genotoxicity test *in vitro*, the extracts were evaluated at three concentrations (250, 100, and 50 mg/mL) and none of the three *G. glauca* extracts showed a genotoxic effect [147].

8. Evaluation of the MeOH extract of *Galphimia glauca* in Behavioral models of anxiety

The anxiolytic and anti-depressive effects were evaluated for the *G. glauca* MeOH extract (wild material) standardized with content of G-B (8.3 mg/g), using the elevated light–dark labyrinth and forced swimming in albino (ICR) mice. The extract, administered orally, three times (24, 18, and 1 h prior to the test) at doses of 125, 250, 500, 1,000, and 2,000 mg/kg was capable of significantly increasing ($p < 0.05$) the number of entries, as well as time spent on the elevated labyrinth's open arms, which indicates an anti-anxiolytic effect. A similar effect was observed in the light–dark paradigm test: time spent in the light box increased in treated mice. However, this treatment was not able to change any parameter in the forced swimming test [148].

9. Conclusions

The MP form part of the daily life of the worldwide population. It is currently of scientific interest due to its high consumption, as an alternative treatment and/or co-administered with allopathic treatments for the improvement of chronic-degenerative diseases. On the other hand, the population has been responsible for affording a great boost to the use of MP; therefore, its consumption generates a great demand and consequently overexploitation. This overexploitation is a danger in the extinction of species of pharmaceutical interest. Another problem regarding the consumption of MP is that not all the population has access to species that are endemic and that have

great biological potential. All the above led to the search for methods to achieve the production and induction of SMt biosynthesis with important biological activity in less time, with constant, controlled and standardized production. Besides helping to preserve plant species without altering the ecosystem.

In some cases, has been reported that cell suspension cultures increase by up to 300% the production of SMt with biological interest respect to wild plant material. In addition, to the increase in SMt production, these are obtained in less complex mixtures, which facilitates the purification process. In the present work, we describe several SMt obtained for biotechnological processing; however, many of these SMt have not been submitted to *in vivo* studies that prove their potential biological activity. Therefore, it is necessary to develop projects aimed at obtaining metabolites by biotechnological processes and demonstrate their biological activity in *in vivo* models.

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Ethical approval

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Nomenclature

SMt	Secondary Metabolites
SMBB	Sociedad Mexicana de Biotecnología y Bioingeniería
TM	Tradicional Medicine
MP	Medicinal Plants
DNA	Deoxyribonucleic acid
PGR	Plany Growth Regulators
BioC	BioCatalyzers
TPA	12-O-Tetradecanoyl Phorbol 13-Acetate
ED ₅₀	Half of the Effective Dose
NAA	Naphthalene Acetic Acid
KIN	Kinetina
2,4-D	Dichlophenoxiacetic acid
BAP	6-BenzylAminoPurine

IC ₅₀	Hal-maximal Inhibitory concentration
DM	Dry Metter
LD ₅₀	Half of a Lethal Dose
DW	Dry Metter
CC ₅₀	Mean Cytotoxic Concentration
i.p.	Intraperitoneal
S.C.	Subcutaneous
G-A	Galphimine A
G-B	Galphimine B
G-E	Galphimine E
C.N	Kinetine

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References

- [1] Rao SR, Ravishankar GA (2002) Plant cell cultures: chemical factories of secondary metabolites. *Biotechnol Adv* 20:101-153. doi: 10.1016/s0734-9750(02)00007-1
- [2] Akula R, Ravishankar GA (2011) Influence of abiotic stress signals on secondary metabolites in plants. *Plant Signal Behav* 6:1720-1731. doi: 10.4161/psb.6.11.17613
- [3] SMBB, Sociedad Mexicana de Biotecnología y Bioingeniería. 2016. <https://smbb.mx/> Access: january 2021.
- [4] Yue W, Ming QL, Lin B, Rahman K, Zheng CJ, Han T, Qin LP (2016) Medicinal plant cell suspension cultures: pharmaceutical applications and high-yielding strategies for the desired secondary metabolites. *Crit Rev Biotechnol* 36:215-232. doi: 10.3109/07388551.2014.923986
- [5] Vanisree M, Tsay HS (2007) Chapter 12: Plant cell cultures: production of biologically important secondary metabolites from medicinal plants of Taiwan. In: Kayser O, Quax WJ (Eds) *Medical Plant Biotechnology: from Basic Research to Industrial Applications*. Willey Verlag Gmb H & Co. K GaA, pp 267-285. doi: 10.1002/9783527619771.ch12
- [6] Vanisree M, Lee CY, Lo SF, Nalawade SM, Lin CY, Tsay HS (2004) Studies on the production of some important secondary metabolites from medicinal plants by plant tissue cultures. *Bot Bull Acad Sin* 45:1-22. <https://www.researchgate.net/publication/228860673>
- [7] Newman DJ, Cragg GM (2020) Natural Products as Sources of New Drugs over the Nearly Four Decades from 01/1981 to 09/2019. *J Nat Prod* 83: 770-803. <https://doi.org/10.1021/acs.jnatprod.9b01285>
- [8] WHO. Traditional Medicine Strategy 2002-2005. World Health Organization; Geneva, Switzerland: 2002. <https://apps.who.int/iris/handle/10665/67163>
- [9] Beyene B, Beyene B, Deribe H (2016) Review on application and management of medicinal plants for the livelihood of the local community. *J Resour Develop Managem* 22:33-39. <https://iiste.org/Journals/index.php/JRDM/article/view/31747/32619>
- [10] Chen SL, Yu H, Luo HM, Wu Q, Li CF, Steinmetz A (2016) Conservation and sustainable use of medicinal plants: problems, progress, and prospects. *Chin Med* 11:1-37. doi:10.1186/s13020-016-0108-7
- [11] Sen T, Samanta SK (2015) Medicinal plants, human health and biodiversity: a broad review. *Adv Biochem Eng Biothechnol* 147:59-110. doi: 10.1007/10_2014_273
- [12] Bahare S, Konovalov DA, Fru P, Kapewangolo P, Peron G, Ksenija M, Pereira OR, Nigam M, Nicola S, Pignata G, Rapposelli S, Sestito S, Anil Kumar NV, Cadiz_Gurrea ML, Segura-Carretero A, Mishra AP, Sharifi-Rad M, Cho WC, Taheri Y, Setzer WN, Sharifi-Rad J (2020) Areca catechu- From farm to food and biomedical applications. *Phytother Res* 34:2140-2158
- [13] Bahare S, Selamoglu Z, Sener B, Kilic M, Kumar Jugran A, de Tommasi N, Sinisgalli C, Milella L, Rajkovic J, Morais-Braga MFB, Bezerra CF, Rocha JE, Coutinho HDM, Oluwaseun AA, Khan SZ, Ahnmad SJ, Erol E, Ali Z, Ostrander EA, Sharifi-Rad

J, Cadiz.Gurrea ML, Taheri Y, Martorell M, Segura-Carretero A, Cho WC (2019a) Berberis Plants-Drifting from Farm to Food Applications, Phytotherapy and Phytopharmacology. Foods 8:522

[14] Bahare S, Sener B, Kilic M, Sharifi-Rad J, Naz R, Yousaf Z, Nixwell MF, Tosouh KPV, Ezzat SM, Bishbishy MH, Taheri Y, Lucariello G, Durazzo A, Lucarini M, Rasul Suleira HA, Santini A (2019) Dioscorea Plants: A genus rich in vital nutraceuticals-A review. Iran J Pharm Res 18:68-69

[15] Farag M, Mohammed MS, Foud I, Ahmed WJ, Malik S, Mohamed MS (2015) The role of natural products in drug discovery and development. World J Pharm Res 4:22-33. <https://www.researchgate.net/publication/291830543>

[16] Krentz AJ, Bailey CJ (2005) Oral antidiabetic agents: current role in type 2 diabetes mellitus. Drugs 65:385-411. doi: 10.2165/00003495-200565030-00005

[17] Romero-Cerecero O, Reyes-Morales H, Torija-Barrio IT, Herrera-Arellano A, Tortoriello J (2005) Conocimiento sobre fitofármacos en médicos de atención primaria del estado de Morelos. Rev Med Inst Mex Seguro Soc 43:281-286. <https://www.medigraphic.com/cgi-bin/new/resumen.cgi?IDARTICULO=4919>

[18] Briskin PD (2000) Medicinal plants and Phytomedicines. Linking plant biochemistry and physiology to human health. Plant Physiol 124:507-514. doi: 10.1104/pp.124.2.507

[19] Ulbricht C, Basch E, Boon H, Ernst E, Hammerness P, Sollars D, Tsourounis C, Woods J, Bent S (2005) Safety review of kava (*Piper methysticum*) by the Natural Standard Research Collaboration. Expert Opin

Drug Saf 4:779-794. doi: 10.1517/14740338.4.4.779

[20] Iheozor-Ejiofor P, Dey ES (2009) Extraction of rosavin from *Rhodiola rosea* root using supercritical carbon dioxide with water. J Supercrit Fluids 50:29-32. <https://doi.org/10.1016/j.supflu.2009.04.011>

[21] Müller S F, Klement S (2006) A combination of valerian and lemon balm is effective in the treatment of restlessness and dyssomnia in children. Phytomedicine 13:383-387. doi: 10.1016/j.phymed.2006.01.013

[22] Vela-Navarrete R, Escribano-Burgos M, Farré AL, García-Cardoso J, Manzarbeitia F, Carrasco C (2005) *Serenoa repens* treatment modifies bax/bcl-2 index expression and caspase-3 activity in prostatic tissue from patients with benign prostatic hyperplasia. J Urol 173:507-510. doi: 10.1097/01.ju.0000150533.94952.25

[23] Gandhi SG, Mahajan V, Bedi YS (2015) Changing trends in biotechnology of secondary metabolism in medicinal and aromatic plants. Planta 241:303-317. doi: 10.1007/s00425-014-2232-x

[24] Sharifi-Rad J, Taheri Y, Abdulmajid AS, Naderi N, Anil Kumar NK, Koirala N, Khadka S, Karazhan N, Shahinozzaman Md, Sen S, Acharya K, Dey A, Martorell M, Martins N (2020) Biological activities and health-promoting effects of *Pyracantha* genus: A key approach to the phytochemical's potential. Cell Mol Biol 66:20-27

[25] Sharifi-Rad J, Rayess YE, Abi Rizk A, Sadaka C, Zgheib R, Zam W, Sestito S, Rapposelli S, Neffe-Skocińska K, Zielińska D, Salehi B, Setzer WN, Dosoky NS, Taheri Y, Beyrouthy ME, Martorell M, Ostrander EA, Rasul Suleira HA, Cho WC, Mroyi A, Martins

N (2020). Turmeric and Its Major Compound Curcumin on Health: Bioactive Effects and Safety Profiles for Food, Pharmaceutical, Biotechnological and Medicinal Applications. *Front Pharmacol* 11:01021

[26] Vuorela P, Leinonen M, Saikku P, Tammela P, Rauha JP, Wennberg T, Vuorela H (2004) Natural products in the process of finding new drug candidates. *Curr Med Chem* 11: 1375-1389. doi: 10.2174/0929867043365116

[27] Patel H, Krishnamurthy R (2013) Elicitors in plant tissue culture. *J Pharmacog Phytochem* 2:60-65. https://www.phytojournal.com/vol2Issue2/Issue_july_2013/9.1.pdf

[28] Pérez-González MZ, Juárez-Vázquez MdelC Jiménez-Arellanes MA (2020) Establishment of a cell suspension culture with anti-inflammatory activity and the effect of salicylic acid on the production of callus from *Cnidioscolus chayamansa*. *Int J Appl Microb Biotechnol Res* 8:40-50. doi.org/10.33500/ijambr.2020.08.005

[29] DiCosmo F, Misawa M (1995) Plant cell and tissue culture: alternatives for metabolite production. *Biotechnol Adv* 13:425-453. doi: 10.1016/07349750(95)02-005-n

[30] Paniego NB, Giuletta AM (1995) *Artemisia annua* L.: dedifferentiated and differentiated cultures. *Plant Cell Tissue Organ Cul* 36 (2), 163-168 doi: 10.1007/BF00037715

[31] Schmitz C, Fritsch L, Fischer R, Schillberg S, Rasche S (2016) Statistical experimental designs for the production of secondary metabolites in plant cell suspension cultures. *Biotechnol Lett* 38: 2007-2014. doi: 10.1007/s10529-016-2206-0

[32] Gaurav N, Singh AP, Kumar A, Som D, Kumar D, Komal HSG (2016) *In vitro* tissue culture studies from nodal and petiole explants of wild and cultivated traits of *Withania somnifera* in MS and B5 medium. *J Med Plant Stud* 4: 92-96. <https://www.researchgate.net/publication/328699276>

[33] McCoy E, O'Connor SE (2008) Natural products from plant cell cultures. *Prog Drug Res* 65:329-370. <https://pubmed.ncbi.nlm.nih.gov/18084920/>

[34] Hernández-Vázquez L, Olivera-Flores M, de Jesús T, Luna H, Navarro-Ocaña A (2017) Obtención de dehidrodiisoeugenol por dimerización de isoeugenol con cultivos celulares de *Bouvardia ternifolia* (trompetilla) TIP. *Rev Esp Cienc Quim-Biol* 20:15-22. <http://dx.doi.org/10.1016/j.recqb.2017.04.002>

[35] Crespi-Perellino N, Guicciardi A, Malyszko G, Arlandi E, Ballobio E, Ballobio, M, Minghetti A (1986) Occurrence of indole alkaloids in *Ailanthus altissima* cell cultures. *J Nat Prod* 49:1010-1014. <https://doi.org/10.1021/np50048a007>

[36] Callebaut A, Hendrickx G, Voets AM, Motte JC (1990) Anthocyanins in cell cultures of *Ajuga reptans*. *Phytochemistry* 29:2153-2158. doi: 10.1016/0031-9422(90)83027

[37] Terahara N, Callebaut A, Ohba R, Nagata T, Ohnishi-Kameyama M, Suzuki M (2001) Acylated anthocyanidin 3-sophoroside-5-glycosides from *Ajuga reptans* flowers and the corresponding cell cultures. *Phytochemistry* 58: 493-500. doi: 10.1016/s0031-9422(01)00172-8

[38] Anderson LA, Harris A, Phillipson JD (1983) Production of cytotoxic

canthin-6-one alkaloids by *Ailanthus altissima* plant cell cultures. J Nat Prod 46:374-378. doi: 10.1021/np50027a014

[39] Staniszevska I, Królicka A, Maliński E, Łojkowska E, Szafranek J (2003) Elicitation of secondary metabolites in *in vitro* cultures of *Ammi majus* L. Enzyme Microb Technol 33: 565-568. [https://doi.org/10.1016/S0141-0229\(03\)00180-7](https://doi.org/10.1016/S0141-0229(03)00180-7)

[40] De-Eknamkul W, Ellis BE (1984) Rosmarinic acid production and growth characteristics of *Anchusa officinalis* cell suspension cultures. Planta Med 50: 346-350. doi: 10.1055/s-2007-969728

[41] Su WW, Lei F, Su LY (1993) Perfusion strategy for rosmarinic acid production by *Anchusa officinalis*. Biotechnol Bioeng 42:884-890. doi: 10.1002/bit.260420713

[42] Vogelsang K, Schneide B, Petersen M (2006) Production of rosmarinic acid and a new rosmarinic acid 3'-O-β-D-glycoside in suspension cultures of the hornwort *Anthoceros agrestis* Paton. Planta 223:369-373. doi: 10.1007/s00425-005-0089-8

[43] Ku KL, Chang PS, Cheng Y C, Lien CY (2005) Production of stilbenoids from the callus of *Arachis hypogaea*: a novel source of the anticancer compound piceatannol. J Agric Food Chem 53:3877-3881. doi: 10.1021/jf050242o

[44] Baldi A, Dixit VK (2008) Yield enhancement strategies for artemisinin production by suspension cultures of *Artemisia annua*. Bioresour Technol 99: 4609-4614. doi: 10.1016/j.biortech.2007.06.061

[45] Liu CZ, Murch SJ, El-Demerdash M, Saxena PK. (2004) *Artemisia judaica* L.: micropropagation and antioxidant

activity. J Biotechnol 110:63-71. doi: 10.1016/j.jbiotec.2004.01.011.

[46] Hunter CS, Kilby NJ (1999) Betalains. Their accumulation and release in vitro. In: Dall RD (Eds) Plant Cell Culture Protocols. Methods in Molecular Biology, vol 11. Humana Press, pp 403-410. doi: 10.1385/1-59259-583-9:403

[47] Pavlov A, Kovatcheva P, Tuneva D, Ilieva M, Bley T (2005a) Radical scavenging activity and stability of betalains from *Beta vulgaris* hairy root culture in simulated conditions of human gastrointestinal tract. Plant Foods Hum Nutr 60:43-47. <https://doi.org/10.1007/s11130-005-5098-z>

[48] Gutiérrez-Rebolledo GA, Estrada-Zúñiga ME, Nieto-Trujillo A, Cruz-Sosa F, Jiménez-Arellanes MA (2018) *In vivo* anti-inflammatory activity and acute toxicity of methanolic extracts from wild plant leaves and cell suspension cultures of *Buddleja cordata* Kunth (Buddlejaceae). Rev Mex Ing Quim 17: 317-330. <http://rmiq.org/ojs311/index.php/rmiq/article/view/42/26>

[49] Estrada-Zúñiga ME, Aarland RC, Rivera-Cabrera F, Bernabé-Antonio A, Buendía-González L, Cruz-Sosa F (2016) Micropropagation of *Buddleja cordata* and the content of verbascoside and total phenols with antioxidant activity of the regenerated plantlets. Rev Mex Ing Quim 15:333-346. <http://rmiq.org/ojs311/index.php/rmiq/article/view/1003>

[50] Zhao P, Iwamoto Y, Kouno I, Egami Y, Yamamoto H (2004) Stimulating the production of homoisoflavonoids in cell suspension cultures of *Caesalpinia pulcherrima* using cork tissue. Phytochemistry 65:2455-2461. <https://doi.org/10.1016/j.phytochem.2004.08.004>

[51] Pawar KD, Thengane SR (2009) Influence of hormones and medium

components on expression of dipyrano-coumarins in cell suspension cultures of *Calophyllum inophyllum* L. Process Biochem 44: 916-922. <https://doi.org/10.1016/j.procbio.2009.03.005>

[52] Orihara Y, Furuya T (1990) Production of theanine and other γ -glutamyl derivatives by *Camellia sinensis* cultured cells. Plant Cell Rep 9:65-68. doi: 10.1007/BF00231550

[53] Sudha G, Ravishankar GA (2003) Influence of methyl jasmonate and salicylic acid in the enhancement of capsaicin production in cell suspension cultures of *Capsicum frutescens* Mill. Curr Sci 85:1212-1217. <https://www.jstor.org/stable/24108623>

[54] Nazif NM, Rady MR, El-Nasr M S (2000) Stimulation of anthraquinone production in suspension cultures of *Cassia acutifolia* by salt stress. Fitoterapia 71:34-40. [https://doi.org/10.1016/S0367-326X\(99\)00101-X](https://doi.org/10.1016/S0367-326X(99)00101-X)

[55] Cortes-Morales JA, Lopez-Laredo AR, Zamilpa A, Bermudez-Torres K, Trejo-Espino JL, Trejo-Tapia G (2018) Morphogenesis and secondary metabolites production in the medicinal plant *Castilleja tenuiflora* benth. Under nitrogen deficiency and starvation stress in a temporary immersion system. Rev Mex Ing Quim 17:229-242. <https://doi.org/10.24275/uam/izt/dcbi/revmexingquim/2018v17n1/Cortes>

[56] Ten Hoopen HJG, Vinke JL, Moreno PRH, Verpoorte R, Heijnen JJ (2002) Influence of temperature on growth and ajmalicine production by *Catharantus roseus* suspension cultures. Enzyme Microb Technol 30:56-65. doi: 10.1016/S0141-0229(01)00456-2

[57] Endo T, Goodbody A, Misawa M (1987) Alkaloid production in root and

shoot cultures of *Catharantus roseus*. Planta Med 53:479-482. doi:10.1055/s-2006-962777

[58] Nicasio-Torres M, Erazo-Gómez J C, Cruz-Sosa F (2009) *In vitro* propagation of two antidiabetic species known as guarumbo: *Cecropia obtusifolia* and *Cecropia peltata*. Acta Physiol Plant 31: 905-914. <https://doi.org/10.1007/s11738-009-0304-5>

[59] Jha S, Sahu NP, Sen J, Jha TB, Mahato SB (1991) Production of emetine and cephaeline from cell suspension and excised root cultures of *Cephaelis ipecacuanha*. Phytochemistry 30: 3999-4003. [https://doi.org/10.1016/0031-9422\(91\)83452-Q](https://doi.org/10.1016/0031-9422(91)83452-Q)

[60] Xu XH, Zhang W, Cao XP, Xue S (2011) Abietane diterpenoids synthesized by suspension-cultured cells of *Cephalotaxus fortunei*. Phytochem Lett 4:52-55. <https://doi.org/10.1016/j.phytol.2010.12.003>

[61] Kitamura Y, Ikenaga T, Ooe Y, Hiraoka N, Mizukami H (1998) Induction of furanocoumarin biosynthesis in *Glehnia littoralis* cell suspension cultures by elicitor treatment. Phytochemistry 48: 113-117. [https://doi.org/10.1016/S0031-9422\(97\)00849-2](https://doi.org/10.1016/S0031-9422(97)00849-2)

[62] Creche J, Trémouillaux-Guiller J, Gröger D, Chénieux J.C, Rideau M (1993) *Choisya ternate* (Mexican Orange): In Vitro Culture and the Production of Alkaloids. In: Bajaj Y.P.S. (Eds) Medicinal and Aromatic Plants V. Biotechnology in Agriculture and Forestry, vol 24. Springer, Berlin, pp 107-121. https://doi.org/10.1007/978-3-642-58062-8_8

[63] Schripsema J, Ramos-Valdivia A, Verpoorte R (1999) Robustaquinones, novel anthraquinones from an elicited *Cinchona robusta* suspension culture.

Phytochemistry 51:55-60. [https://doi.org/10.1016/S0031-9422\(98\)00470-1](https://doi.org/10.1016/S0031-9422(98)00470-1)

[64] Cheng XY, Wei T, Guo B, Ni W, Liu CZ (2005) *Cistanche deserticola* cell suspension cultures: phenylethanoid glycosides biosynthesis and antioxidant activity. *Process Biochem* 40:3119-3124. doi: 10.1016/j.procbio.2005.03.012

[65] Liu JY, Guo ZG, Zeng ZL (2007) Improved accumulation of phenylethanoid glycosides by precursor feeding to suspension culture of *Cistanche salsa*. *Biochem Eng J* 33:88-93. <https://doi.org/10.1016/j.bej.2006.09.002>

[66] Yoshida K, Hayashi T, Sano K (1988) Colchicoside in *Colchicum autumnale* bulbs. *Agr Biol Chem* 52:593-594. <https://doi.org/10.1080/00021369.1988.10868674>

[67] Mukherjee S, Ghosh B, Jha S (2000) Establishment of forskolin yielding transformed cell suspension cultures of *Coleus forskohlii* as controlled by different factors. *J Biotechnol* 76:73-81. doi: 10.1016/S0168-1656(99)00181-9

[68] Pasqua G, Monacelli B, Mulinacci N, Rinaldi S, Giaccherini C, Innocenti M, Vinceri FF (2005) The effect of growth regulators and sucrose on anthocyanin production in *Camptotheca acuminata* cell cultures. *Plant Physiol Biochem* 43:293-298. <https://doi.org/10.1016/j.plaphy.2005.02.009>

[69] Hara Y, Yamagata H, Morimoto T, Hiratsuka J, Yoshioka T, Fujita Y, Yamada Y (1989) Flow cytometric analysis of cellular berberine contents in high-and low-producing cell lines of *Coptis japonica* obtained by repeated selection. *Planta Med* 55:151-154. doi: 10.1055/s-2006-961910

[70] Ishimaru K, Arakawa H, Neera S (1993) Polyphenol production in cell

cultures of *Cornus kousa*. *Phytochemistry* 32:1193-1197. [https://doi.org/10.1016/S0031-9422\(00\)95090-8](https://doi.org/10.1016/S0031-9422(00)95090-8)

[71] Kashyap S, Kapoor N, Kale RD. (2016) *Coscinium fenestratum*: callus and suspension cell culture of the endangered medicinal plant using vermicompost extract and coelomic fluid as plant tissue culture media. *Am J Plant Sci* 7:899-906. doi: 10.4236/ajps.2016.76085

[72] Chen SA, Wang X, Zhao B, Yuan X, Wang Y (2003) Production of crocin using *Crocus sativus* callus by two-stage culture system. *Biotechnol Lett* 25:1235-1238. <https://link.springer.com/article/10.1023/A:1025036729160>

[73] Trajtemberg SP, Apóstolo NM, Fernández G (2006) Calluses of *Cynara cardunculus* var. *cardunculus cardoon* (Asteraceae): determination of cynarine and chlorogenic acid by automated high-performance capillary electrophoresis. *In Vitro Cell Dev Biol-Plant* 42:534-537. <https://doi.org/10.1079/IVP2006803>

[74] Ravindra PV, Narayan MS (2003) Antioxidant activity of the anthocyanin from carrot (*Daucus carota*) callus culture. *Int J Food Sci Nut* 54:349-355. doi: 10.1080/09637480120092134

[75] Kreis W, Reinhard E (1992) 12 β -Hydroxylation of digitoxin by suspension-cultured *Digitalis lanata* cells: production of digoxin in 20-litre and 300-litre air-lift bioreactors. *J Biotechnol* 26:257-273. [https://doi.org/10.1016/0168-1656\(92\)90011-W](https://doi.org/10.1016/0168-1656(92)90011-W)

[76] Tal B, Gressel J, Goldberg I (1982) The effect of medium constituents on growth and diosgenin production by *Dioscorea deltoidea* cells grown in batch cultures. *Planta Med* 44:111-115. doi: 10.1055/s-2007-971414

- [77] Komaraiah P, Ramakrishna SV, Reddanna P, Kishor PK (2003) Enhanced production of plumbagin in immobilized cells of *Plumbago rosea* by elicitation and *in situ* adsorption. J Biotechnol 101:181-187. doi: 10.1016/s0168-1656(02)00338-3.
- [78] Shohael AM, Ali MB, Yu KW, Hahn EJ, Islam R, Paek KY (2006) Effect of light on oxidative stress, secondary metabolites and induction of antioxidant enzymes in *Eleutherococcus senticosus* somatic embryos in bioreactor. Process Biochem 41:1179-1185. <https://doi.org/10.1016/j.procbio.2005.12.015>
- [79] Ho HY, Liang KY, Lin WC, Kitanaka S, Wu JB (2010) Regulation and improvement of Triterpene formation in plant cultured cells of *Eriobotrya japonica* Lindl. J Biosci Bioeng 110:588-592. doi: 10.1016/j.jbiosc.2010.06.009
- [80] Wang J, Liao X, Zhang H, Du J, Chen P (2003a) Accumulation of chlorogenic acid in cell suspension cultures of *Eucommia ulmoides*. Plant Cell Tiss Organ Cult 74:193-195. <https://doi.org/10.1023/A:1023957129569>
- [81] Lee SY, Cho SI, Park MH, Kim YK, Choi JE, Park SU (2007) Growth and rutin production in hairy root cultures of buckwheat (*Fagopyrum esculentum* M.) Prep Biochem Biotechnol 37:239-246. doi: 10.1080/10826060701386729
- [82] Zhang W, Seki M, Furusaki S (1997) Effect of temperature and its shift on growth and anthocyanin production in suspension cultures of strawberry cells. Plant Sci 127:207-214. [https://doi.org/10.1016/S0168-9452\(97\)00124-6](https://doi.org/10.1016/S0168-9452(97)00124-6)
- [83] Osuna L, Pereda-Miranda R, Villarreal ML (2002) *In vitro* production of sedative galphimine B by cell suspension cultures of *Galphimia glauca*. Biotechnol Lett 24:257-261. <https://doi.org/10.1023/A:1014028510775>
- [84] Miura H, Kitamura Y, Ikenaga T, Mizobe K, Shimizu T, Nakamura M, Kato Y, Yamada T, Maitani T, Goda Y (1998) Anthocyanin production of *Glehnia littoralis* callus cultures. Phytochemistry 48:279-283. [https://doi.org/10.1016/S0031-9422\(97\)01115-1](https://doi.org/10.1016/S0031-9422(97)01115-1)
- [85] Kitamura Y, Ohta M, Ikenaga T, Watanabe M (2002) Responses of anthocyanin-producing and non-producing cells of *Glehnia littoralis* to radical generators. Phytochemistry 59:63-68. doi: 10.1016/s0031-9422(01)00428-9
- [86] Praveen N, Murthy HN, Chung IM (2011) Improvement of growth and gymnemic acid production by altering the macro elements concentration and nitrogen source supply in cell suspension cultures of *Gymnema sylvestre* R. Br. Ind Crops Prod 33:282-286. <https://doi.org/10.1016/j.indcrop.2010.12.015>
- [87] Taha HS, El-Kawy AA, Fathalla MA (2012) A new approach for achievement of inulin accumulation in suspension cultures of Jerusalem artichoke (*Helianthus tuberosus*) using biotic elicitors. J Genet Eng Biotechnol 10:33-38. <https://doi.org/10.1016/j.jgeb.2012.02.002>
- [88] Misra N, Misra P, Datta SK, Mehrotra S (2005) *In vitro* biosynthesis of antioxidants from *Hemidesmus indicus* R. Br. Cultures. In Vitro Cell Dev Biol-Plant 41:285-290. <https://doi.org/10.1079/IVP2004627>
- [89] Walker TS, Bais HP, Vivanco JM (2002) Jasmonic acid-induced hypericin production in cell suspension cultures of *Hypericum perforatum* L. (St. John's wort). Phytochemistry 60:289-293. doi: 10.1016/s0031-9422(02)00074-2

- [90] Murakami Y, Omoto T, Asai I, Shimomura K, Yoshihira K, Ishimaru K (1998) Rosmarinic acid and related phenolics in transformed root cultures of *Hyssopus officinalis*. *Plant Cell Tiss Org* 53:75-78. <https://doi.org/10.1023/A:1006007707722>
- [91] Kochan E, Wysokinska H, Chmiel A, Grabias B (1999) Rosmarinic acid and other phenolic acids in hairy roots of *Hyssopus officinalis*. *Z Naturforsch C* 54c: 11-16. <https://doi.org/10.1515/znc-1999-1-204>
- [92] Konczak-Islam I, Yoshimoto M, Hou D. X, Terahara N, Yamakawa O (2003) Potential chemopreventive properties of anthocyanin-rich aqueous extracts from *in vitro* produced tissue of sweet potato (*Ipomoea batatas* L.). *J Agric Food Chem* 51:5916-5922. <https://doi.org/10.1021/jf030066o>
- [93] Terahara N, Konczak I, Ono H, Yoshimoto M, Yamakawa O (2004) Characterization of acylated anthocyanins in callus induced from storage root of purple-fleshed sweet potato, *Ipomoea batatas* L. *J Biomed Biotechnol* 2004:279-286. doi: 10.1155/S1110724304406056
- [94] Terahara N, Konczak-Islam I, Nakatani M, Yamakawa O, Goda Y, Honda T (2000) Anthocyanins in callus induced from purple storage root of *Ipomoea batatas* L. *Phytochemistry* 54: 919-922. [https://doi.org/10.1016/S0031-9422\(00\)00099-6](https://doi.org/10.1016/S0031-9422(00)00099-6)
- [95] Palacio L, Cantero JJ, Cusidó RM, Goleniowski ME (2012) Phenolic compound production in relation to differentiation in cell and tissue cultures of *Larrea divaricata* (Cav.). *Plant Sci* 193-194: 1-7. doi: 10.1016/j.plantsci.2012.05.007
- [96] Georgiev MI, Kuzeva SL, Pavlov AI, Kovacheva EG, Ilieva MP (2007) Elicitation of rosmarinic acid by *Lavandula vera* MM cell suspension culture with abiotic elicitors. *World J Microbiol Biotechnol* 23:301-304. doi: 10.1007/s11274-006-9214-5
- [97] Pavlov AI, Georgiev MI, Panchev IN, Ilieva MP (2005b) Optimization of rosmarinic acid production by *Lavandula vera* MM plant cell suspension in a laboratory bioreactor. *Biotechnol Prog* 21:394-396. doi: 10.1021/bp049678z
- [98] Espinosa-Leal C, Treviño-Neávez JF, Garza-Padrón RA, Verde-Star MJ, Rivas-Morales C, Morales-Rubio ME (2015) Contenido de fenoles totales y actividad anti-radical de extractos metanólicos de la planta silvestre y cultivada *in vitro* de *Leucophyllum frutescens*. *Rev Mex Cienc Farm* 46:52-56. <https://www.redalyc.org/articulo.oa?id=57945705005>
- [99] Yamamoto H, Inoue K, Yazaki K (2000) Caffeic acid oligomers in *Lithospermum erythrorhizon* cell suspension cultures. *Phytochemistry* 53: 651-657. [https://doi.org/10.1016/S0031-9422\(99\)00623-8](https://doi.org/10.1016/S0031-9422(99)00623-8)
- [100] Salinas R, Arellano-García J, Perea-Arango I, Álvarez L, Garduño-Ramírez ML, Marquina S, Zamilpa, A, Castillo-España P (2014) Production of the anti-Inflammatory compound 6-O-Palmitoyl-3-O-β-D-glucopyranosylcampesterol by callus cultures of *Lopezia racemosa* Cav. (Onagraceae) *Molecules* 19:8679-8690
- [101] Jang YP, Lee YJ, Kim YC, Huh H (1998) Production of a hepatoprotective cerebroside from suspension cultures of *Lycium chinense*. *Plant Cell Rep* 18: 252-254. doi: 10.1007/s002990050566
- [102] Chiang L, Abdullah MA (2007) Enhanced anthraquinones production from adsorbent-treated *Morinda elliptica* cell suspension cultures in production

medium strategy. *Process Biochem* 42: 757-763. doi: 10.1016/j.procbio.2007.01.005

[103] Pras N, Woerdenbag HJ, Batterman S, Visser JF, Van Uden W (1993) *Mucuna pruriens*: improvement of the biotechnological production of the anti-Parkinson drug L-dopa by plant cell selection. *Pharm World Sci* 15:263-268. doi: 10.1007/BF01871128

[104] Kouamo K, Creche J, Chénieux JC, Rideau M, Viel C (1985) Alkaloid production by *Ochrosia elliptica* cell suspension cultures. *J Plant Physiol* 118: 277-283. doi: 10.1016/S0176-1617(85)80229-7

[105] Kintzios S, Kollias H, Straitouris E, Makri O (2004) Scale-up micropropagation of sweet basil (*Ocimum basilicum* L.) in an airlift bioreactor and accumulation of rosmarinic acid. *Biotechnol Lett* 26:521-523. doi: 10.1023/b:bile.0000019561.89044.30

[106] Kang TH, Park HM, Kim YB, Kim H, Kim N, Do JH, Kang C, Cho Y, Kim SY (2009) Effects of red ginseng extract on UVB irradiation-induced skin aging in hairless mice. *J Ethnopharmacol* 123: 446-451. doi: 10.1016/j.jep.2009.03.022

[107] Zhong JJ, Bai Y, Wang SJ (1996) Effects of plant growth regulators on cell growth and ginsenoside saponin production by suspension cultures of *Panax quinquefolium*. *J Biotechnol* 45: 227-234. [https://doi.org/10.1016/0168-1656\(95\)00170-0](https://doi.org/10.1016/0168-1656(95)00170-0)

[108] Tam WJ, Constabel F, Kurz WG (1980) Codeine from cell suspension cultures of *Papaver somniferum*. *Phytochemistry* 19:486-487. doi: 10.1016/0031-9422(80)83215-8

[109] Huang FC, Kutchan TM (2000) Distribution of morphinan and benzo [c]

phenanthridine alkaloid gene transcript accumulation in *Papaver somniferum*. *Phytochemistry* 53:555-564. [https://doi.org/10.1016/S0031-9422\(99\)00600-7](https://doi.org/10.1016/S0031-9422(99)00600-7)

[110] Archambault J, Williams RD, Bédard C, Chavarie C (1996) Production of sanguinarine by elicited plant cell culture I. Shake flask suspension cultures. *J Biotech* 46:95-105. doi: 10.1016/0168-1656(95)00184-0

[111] Antognoni F, Zheng S, Pagnucco C, Baraldi R, Poli F, Biondi S (2007) Induction of flavonoid production by UV-B radiation in *Passiflora quadrangularis* callus cultures. *Fitoterapia* 78: 345-352. doi: 10.1016/j.fitote.2007.02.001

[112] Hempel J, Pforte H, Raab B, Engst W, Böhm H, Jacobasch G (1999) Flavonols and flavones of parsley cell suspension culture change the antioxidative capacity of plasma in rats. *Mol Nutr Food Res* 43:201-204 doi: 10.1002/(SICI)1521-3803(19990601)43:3<201::AID-FOOD201>3.0.CO;2-1

[113] Scragg AH, Allan EJ (1986) Production of the triterpenoid quassin in callus and cell suspension cultures of *Picrasma quassioides* Bennett. *Plant Cell Rep* 5:356-359. doi: 10.1007/BF00268601

[114] Saad I, Díaz E, Chávez I, Reyes-Chilpa R, Rubluo A, Jiménez-Estrada M (2000) Antifungal monoterpene production in elicited cell suspension cultures of *Piqueria trinervia*. *Phytochemistry* 55:51-57. [https://doi.org/10.1016/S0031-9422\(00\)00211-9](https://doi.org/10.1016/S0031-9422(00)00211-9) doi: 10.3390/molecules19068679

[115] Chattopadhyay S, Srivastava AK, Bhojwani SS, Bisaria VS (2002) Production of podophyllotoxin by plant cell cultures of *Podophyllum hexandrum* in bioreactor. *J Biosc Bioeng* 93:215-220. doi: 10.1016/S1389-1723(02)80017-2

- [116] Shinde AN, Malpathak N, Fulzele DP (2009) Studied enhancement strategies for phytoestrogens production in shake flasks by suspension culture of *Psoralea corylifolia*. *Bioresour Technol* 100:1833-1839. <https://doi.org/10.1016/j.biortech.2008.09.028>
- [117] Yamamoto O, Yamada Y (1986) Production of reserpine and its optimization in cultured *Rauwolfia serpentina* Benth. *Cells. Plant Cell Rep* 5: 50-53. doi: 10.1007/BF00269717
- [118] Perassolo M, Quevedo CV, Giulietti AM, Talou JR (2011) Stimulation of the proline cycle and anthraquinone accumulation in *Rubia tinctorum* cell suspension cultures in the presence of glutamate and two proline analogs. *Plant Cell Tiss Organ Cult* 106: 153-159. <https://doi.org/10.1007/s11240-010-9903-5>
- [119] Tsutomu N, Hitoshi M, Masao N, Hideko H, Kaisuke Y (1983) Production of cryptotanshinone and ferruginol in cultured cells of *Salvia miltiorrhiza*. *Phytochemistry* 22:721-722. [https://doi.org/10.1016/S0031-9422\(00\)86969-1](https://doi.org/10.1016/S0031-9422(00)86969-1)
- [120] Chen H, Yuan JP, Chen F, Zhang YL, Song JY (1997) Tanshinone production in Ti-transformed *Salvia miltiorrhiza* cell suspension cultures. *J Biotechnol* 58:147-156. doi: 10.1016/S0168-1656(97)00144-2
- [121] Zhao DX, Fu CX, Han YS, Lu DP (2005) Effects of elicitation on jaceosidin and hispidulin production in cell suspension cultures of *Saussurea medusa*. *Process Biochem* 40:739-745. <https://doi.org/10.1016/j.procbio.2004.01.040>
- [122] Sánchez-Sampedro MA, Fernández-Tárrago J, Corchete P (2005) Yeast extract and methyl jasmonate-induced silymarin production in cell cultures of *Silybum marianum* (L.) Gaertn. *J Biotechnol* 119:60-69. doi: 10.1016/j.jbiotec.2005.06.012
- [123] Aburjai T, Bernasconi S, Manocchi LA, Pelizzoni F (1997) Effect of calcium and cell immobilization on the production of cholecalciferol and its derivatives by *Solanum malacoxylon* cell cultures. *Phytochemistry* 46:1015-1018. doi: 10.1016/S0031-9422(97)00408-1
- [124] Pérez-Hernández J, González-Cortázar M, Marquina S, Herrera-Ruiz M, Meckes-Fischer M, Tortoriello J, Cruz-Sosa F, Nicasio-Torres MP (2014) Sphaeralcic acid and Tomentin, Anti-inflammatory compounds produced in cell suspension cultures of *Sphaeralcea angustifolia*. *Planta Med* 80:209-214. doi: 10.1055/s-0033-1360302
- [125] Rico-Rodríguez L, Ortiz-Butrón R, Silva-Torres R, Díaz-Cedillo F, Franco-Colín M (2015) Diseño bifactorial de reguladores de crecimiento para la obtención de un alcaloide a partir del cultivo celular de *Swietenia humilis* Zucc. *Rev Mex Cienc Farm* 46:57-61. <https://www.redalyc.org/articulo.oa?id=57945705006>
- [126] Nieto-Trujillo A, Buendía-González L, García-Morales, C, Román-Guerrero, A, Cruz-Sosa F, Estrada-Zúñiga ME (2017) Phenolic compounds and parthenolide production from *in vitro* cultures of *Tanacetum parthenium*. *Rev Mex Ing Qum* 16: 371-387. <https://www.redalyc.org/pdf/620/62052087004.pdf>
- [127] Yukimune Y, Tabata H, Higashi Y, Hara Y (1996) Methyl jasmonate-induced overproduction of Paclitaxel and Baccatin III in *Taxus* cell suspension cultures. *Nat Biotechnol* 14:1129-1132. <https://doi.org/10.1038/nbt0996-1129>
- [128] Wang Q, Mawassi M, Li P, Gafny R, Sela I, Tanne E (2003b)

Elimination of grapevine virus A (GVA) by cryopreservation of *in vitro*-grown shoot tips of *Vitis vinifera* L. Plant Sci 165:321-327. [https://doi.org/10.1016/S0168-9452\(03\)00091-8](https://doi.org/10.1016/S0168-9452(03)00091-8)

[129] Yue X, Zhang W, Deng M (2011) Hyper-production of 13 C-labeled trans-resveratrol in *Vitis vinifera* suspension cell culture by elicitation and *in situ* adsorption. Biochem Eng J 53:292-296. <https://doi.org/10.1016/j.bej.2010.12.002>

[130] Nagella P, Murthy HN (2010) Establishment of cell suspension cultures of *Withania somnifera* for the production of withanolide A. Bioresour Technol 101: 6735-6739. <https://doi.org/10.1016/j.biortech.2010.03.078>

[131] Pérez-Hernández J, Nicasio-Torres MP, Sarmiento-López LG, Rodríguez-Monroy M (2019) Production of anti-inflammatory compounds in *Sphaeralcea angustifolia* cell suspension cultivated in stirred tank biorreactor. Eng Life Sci 19: 196-205. doi: 10.1002/elsc.201800134

[132] Nicasio-Torres MP, Pérez-Hernández J, González-Cortázar M, Meckes-Fischer M, Tortoriello J, Cruz-Sosa F (2016) Production of potential anti-inflammatory compounds in cell suspension cultures of *Sphaeralcea angustifolia* (Cav.) G. Don. Acta Physiol Plant 38:209. doi:10.1007/s11738-016-2211-x

[133] Nicasio-Torres MP, Serrano-Román J, Pérez-Hernández J, Jiménez-Ferrer E, Herrera-Ruiz M (2017) Effect of dichloromethane-methanol extract and tomentin obtained from *Sphaeralcea angustifolia* cell suspensions in a model of kaolin/carrageenan-induced arthritis. Planta Med Int Open 4, e35-e42. doi: 10.1055/s-0043-108760

[134] García-Rodríguez RV, Chamorro-Cevallos G, Siordia G, Jiménez-Arellanes

MA, Chávez-Soto MA, Meckes-Fischer M (2012) *Sphaeralcea angustifolia* (Cav.) G. Don extract, a potential phytomedicine to treat chronic inflammation. BLACPMA 11:454-463. <https://www.redalyc.org/articulo.oa?idp=1&id=85624131008&cid=47965>

[135] Romero-Cerecero O, Meckes-Fischer M, Zamilpa A, Jiménez-Ferrer JE, Nicasio-Torres P, Pérez-García D, Tortoriello J (2013) Clinical trial for evaluating the effectiveness and tolerability of topical *Sphaeralcea angustifolia* treatment in hand osteoarthritis. J Ethnopharmacol 147: 467-473. doi: 10.1016/j.jep.2013.03.040

[136] Albarello N, Simotilde C, de Castro TC, Gayer CRM, Coelho MGP, de Moura R.S, Mansur E (2013) Anti-inflammatory and antinociceptive activity of field-growth plants and tissue culture of *Cleome spinosa* (Jacq.) in mice. J Med Plants Res 7:1043-1049. doi: 10.5897/JMPR12.153

[137] Pérez-González MZ, Gutiérrez-Rebolledo GA, Yépez-Mulia L, Rojas-Tomé IS, Luna-Herrera J. Jiménez-Arellanes MA (2107) Antiprotozoal, antimycobacterial and anti-inflammatory evaluation of *Cnidoscolus chayamansa* (Mc Vaugh) extract and the isolated compounds. Biomed Pharmacother 89:89-97. doi: 10.1016/j.biopha.2017.02.021

[138] Pérez-González MZ, Siordia-Reyes AG, Damián-Nava P, Hernández-Ortega S, Macías-Rubalcava ML, Jiménez-Arellanes MA (2018) Hepatoprotective and anti-inflammatory activities of the *Cnidoscolus chayamansa* (Mc Vaugh) leaf extract in chronic models. Evid Based Complement Alternat Med 2018: 3896517. doi: 10.1155/2018/3896517

[139] Pérez-González MZ, Macías-Rubalcava ML, Hernández-Ortega S,

Siordia-Reyes AG, Jiménez-Arellanes MA (2019) Additional compounds and the therapeutic potential of *Cnidoscolus chayamansa* (McVaugh) against hepatotoxicity induced by antitubercular drugs. *Biomed Pharmacother* 117: 1091140, 1-10. doi: 10.1016/j.biopha.2019.109140

[140] Pérez-González MZ, Nieto-Trujillo A, García-Martínez I, Estrada-Zuñiga ME, Bernabe-Antonio A, Jiménez-Arellanes MA, Cruz-Sosa F (2018) Establishing a callus culture of *Cnidoscolus chayamansa* Mc Vaugh: A species with ethnopharmacological value. *Adv Biochem Biotech* 2018:1-5. https://gavinpublishers.com/admin/assets/articles_pdf/1544189353article_pdf1396193972.pdf

[141] Pérez-González MZ, Nieto-Trujillo A, García-Martínez I, Estrada-Zuñiga ME, Bernabé-Antonio A, Jiménez-Arellanes MA Cruz-Sosa F (2019) Lupeol acetate production and antioxidant activity of a cell suspension culture from *Cnidoscolus chayamansa* leaves. *S Afr J Bot* 125:30-38. <https://doi.org/10.1016/j.sajb.2019.06.030>

[142] Taniguchi S, Imayoshi Y, Kobayashi E, Takamatsu Y, Ito H, Hatano T, Harukuni S, Tokudac H, Nishinoc H, Sugitad D, Shimurad S, Yoshida T (2002) Production of bioactive triterpenes by *Eriobotrya japonica* calli. *Phytochemistry* 59: 315-323. [https://doi.org/10.1016/S0031-9422\(01\)00455-1](https://doi.org/10.1016/S0031-9422(01)00455-1)

[143] Tortoriello J, Ortega A (1993) Sedative effect of galphimine B, a nor-seco-triterpenoid from *Galphimia glauca*. *Planta Med* 59:398-400. doi: 10.1055/s-2006-959717

[144] Tortoriello J, Ortega A, Herrera-Ruiz M, Trujillo J, Reyes-Vázquez C (1998) Galphimine-B modifies electrical

activity of ventral tegmental area neurons in rats. *Planta Med* 64:309-313. doi: 10.1055/s-2006-957440

[145] Osuna L, Pereda-Miranda R, Tortoriello J, Villarreal ML (1999) Production of the sedative triterpene galphimine B in *Galphimia glauca* tissue culture. *Planta Med* 65:149-152. doi: 10.1055/s-1999-14057

[146] Osuna L, Moyano E, Mangas S, Bonfill M, Cusidó RM, Piñol MT, Zamilpa A, Tortoriello L, Palazón J (2008) Immobilization of *Galphimia glauca* plant cell suspensions for the production of enhanced amounts of Galphimine-B. *Planta Med* 74:94-99. doi: 10.1055/s-2007-993763

[147] Aguilar-Santamaría L, Ramírez G, Herrera-Arellano A, Zamilpa A, Jiménez JE, Alonso-Cortés D, Cortés-Gutiérrez EI, Ledesma N, Tortoriello J (2007) Toxicological and cytotoxic evaluation of standardized extracts of *Galphimia glauca*. *J Ethnopharmacol* 109: 35-40. doi: 10.1016/j.jep.2006.06.013

[148] Herrera-Ruiz M, Jiménez-Ferrer JE, De Lima TCM, Avilés-Montes D, Pérez-García D, González-Cortázar M, Tortoriello J (2006) Anxiolytic and antidepressant-like activity of a standardized extract from *Galphimia glauca*. *Phytomedicine* 13:23-28. doi: 10.1016/j.phymed.2005.03.003