Saponin production from *Quillaja* genus species. An insight into its applications and biology

Fernando Guerra1*, Sebastián Sepúlveda2

1Universidad de Talca, Instituto de Ciencias Biológicas, Av. Lircay s/n, Casilla 747, Talca, Chile.  
2Université de Lorraine, Boulevard des Aiguillettes, BP 70239, 54506 Vandœuvre-les-Nancy, France.  
*Corresponding author <fguerra@utalca.cl>

ABSTRACT: *Quillaja* genus (*Quillajaceae* family) is endemic to South America, where is represented by two species, *Quillaja saponaria* and *Quillaja brasiliensis*. One outstanding characteristic of these forest tree species is their production of saponins, a family of amphipathic glycosides, involved in the defensive response of plants against biotic and abiotic factors. Saponins are metabolites of economic importance due to their chemical and physical properties. Basic and applied research efforts performed during the last decades, mainly on *Q. saponaria*, have placed these compounds as an important raw material in industrial areas, such as food and beverage, cosmetics, vaccine production, biopesticides, among others. In this review, we summarize information on saponins from *Quillaja* species during the last years, analyzing current developments by application areas, as well as their chemical composition and properties. We also describe the general advances in revealing saponin biosynthesis pathways, related genes and *Quillaja* genomes, as well as the conservation status, domestication processes, and perspectives in the context of implementing genetic improvement programs.

Keywords: triterpenes, secondary metabolism, chemical structure, biosynthesis pathway, genes

**Introduction**

*Quillaja* is a genus of the *Quillajaceae* family of the Fabales order (*Magnoliopsida*) and comprises only two species: *Quillaja saponaria* Molina (*QS*) and *Quillaja brasiliensis* (St. A.-Hil. & Tul.) Mart. (*QB*) (Mello and Cantos, 2014). Distribution of both evergreen forest tree species includes specific areas in South America. QB is endemic to southern Brazil and northeastern Argentina, eastern Paraguay and northern Uruguay, (between 21° S and 31° S and from 46° W to 58° W; ranging up to 1350 m.a.s.l) (Mello et al., 2014). QS is endemic to the Mediterranean regions of Chile, occurring in mixed forests, in the so-called sclerophyllous forest, and growing abundantly between 30° S and 38° S, reaching up to 2000 m.a.s.l (Schlotterbeck et al., 2015). The traditional interest on this genus is because of saponins in the bark, wood, and leaves of trees (Figure 1). Both species, notably QS, represent important commercial sources of raw saponins used as a foaming, wetting, and emulsifying agent, and, in a more purified form, in preparing photographic emulsions, cosmetics, vaccine adjuvants and other medical products (Rodríguez-Díaz et al., 2011). In particular, QS is well known for its relatively high content of triterpene saponins (between 1.2 % in branches and near to 16-20 % of dry weight at the bark), which are specially destined to personal care products and used as vaccine adjuvants (Copaja et al., 2003; Arrau et al., 2011; Magedans et al., 2019). In Chile, annual exportation of QS extracts, bark, and wood powder registered 20.6 million U.S. dollars in 2018 (INFOR, 2019). Along with the increasing demand for QS extract, pressure on natural forests has increased in the last decades, with a prediction of an overexploitation trend for the next years (Schlotterbeck et al., 2015). Since the identification of saponins as the main metabolites in *Quillaja* species, many studies have investigated their composition and applications, mainly on QS and, to a lesser extent, on QB. In the latter, characterization of saponins and the analyses of their potential uses are relatively recent (Costa et al., 2014; Cibulski et al., 2018; Wallace et al., 2017, 2019). In this review, we analyze historical aspects of the utilization of *Quillaja* species, and the progress in the characterization and applications of their saponins, incorporating general advances in the identification of saponin biosynthetic pathways, and related genes. In addition, we discuss considerations

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**Figure 1** – Images of *Quillaja saponaria* (*QS*) and *Quillaja brasiliensis* (*QB*). A) QS bark; B) QS leaves; C) QB bark; D) QB leaves. Photography credits: A-B: Fernando Guerra and Sebastián Sepúlveda; C: Marcela Huertas Suárez; D: Jair Gilberto Kray.
and perspectives in the framework of domestication and development of genetic improvement programs for *Quillaja* species.

**History and management of *Quillaja* species**

Mapuche people, the major ethnic group of southern-central Chile, have historically used the QS tree, "quillay" [a word derived from Mapuche’s language "küllay" or "cúllcan", which means "to wash"], for their personal cleaning, and treating toothaches and respiratory diseases [Arrau et al., 2011; Rodríguez-Díaz et al., 2011]. In 1944, its intensive exploitation encouraged the Chilean government to regulate bark extraction, promoting natural regeneration and sustainable utilization of this species. Nevertheless, illegal exploitations have continued [San Martin, 2000].

Currently, natural forests are the principal source (98 %) of raw material. The total area in Chile comprising QS (within the sclerophyllous forest) has been estimated at 230,000 ha with a biomass production of 2 t ha⁻¹ every 15 years [San Martin, 2000; Schlotterbeck et al., 2015]. The sustainable limit of natural forests has been estimated at 27,000 t yr⁻¹. Nevertheless, it is estimated that 48,000 tons are required to sustain market demands by 2019-2020, challenging its self-renewal ability [Schlotterbeck et al., 2015]. The International Union for Conservation of Nature (IUCN) qualifies this species under no risk of conservation [LC] [BGCI, 2018]. The diversity of QS communities has been analyzed across 39 sites within their distribution range by Letelier et al. [2017], who estimated a Shannon-Wiener’s index of 0.56, associating their presence with geographic and climate variables, and anthropogenic disturbance. Efforts for domesticating this species have been made since the middle of the last century in Chile. Important knowledge has been generated about silvicultural management focused on propagation, productive improvement of natural forests, and establishment of plantations [San Martin et al., 2000; Prehn et al., 2003; Jordan and Roveraro, 2004; Schlotterbeck et al., 2015; Vidal et al., 2016].

Production of QS derived products starts with the logging and subsequent extraction of bark pieces from 25 years-old individuals or older [Magedans et al., 2019]. Then, the bark is air-dried, ground, treated with boiled water or water-alcohol mixtures, stabilized, filtered and evaporated [San Martin, 2000]. Natural regeneration produces new individuals with a bush-like architecture [multiple stems from the stump], which hinders extraction of bark from second-rotation trees. To counteract this effect, sustainable management of QS forests includes the application of thinning on new stems and the use of alternative raw material sources, such as branches from pruning and leaves [containing up to 2.58 % w/w of saponins] [Schlotterbeck et al., 2015]. However, current industrial protocols are not designed to utilize leaves in the extraction process [Magedans et al., 2019]. Starting from seeds, the germination rate depends on parental tree genetics, ranging from 61 to 92 % of success [Prehn et al., 2006]. Then, seedlings are planted in the field after one growing season in a nursery, following 3 × 3 or 3 × 2 m plantation frames [Benedetti et al., 2000]. Moreover, direct organogenesis [Vidal et al., 2016] and in-vitro micropropagation [Prehn et al., 2003; Jordan and Roveraro, 2004] have been suggested as methods to optimize the vegetative multiplication of selected clones with high saponin production rates. During the 2000s, genetic improvement of QS started in Chile with mass selection of parental trees, according to their growth and industrial yield of extracts, breeding and seed orchard establishment [Velozo, 2008]. A set of eight "plus" clones was generated, which was incorporated into operational plantation programs.

The increasing demand for QS saponins in the vaccine market has driven additional research to characterize equivalent saponin fractions in QB in the last two decades. The first publication on this subject was in 2004 [according to search performed at the Web of Science platform, in September 2019]. Unlike QS, QB leaves contain more saponins than bark [0.085 % vs 0.022 % of dry weight] [Magedans et al., 2019]. The renewable use of bioactive saponins from QB leaves has gained more importance in recent years [Yendo et al., 2014; Costa et al., 2013, 2014; Velazco et al., 2018, 2019; Cibulski et al., 2017-2018; Fleck et al., 2009, 2019; Wallace et al., 2017, 2019]. According to Fleck et al. [2009], which could be further increased when aseptic culture media are used. Additionally, the use of seeds to obtain rooted microcuttings was an efficient alternative to produce plants with saponin contents comparable to those of adult trees in field conditions [Fleck et al., 2009].

**Main applications of Quillaja saponins**

Traditional utilization of *Quillaja* saponins in the food and cosmetic industries has been associated with their foaming and emulsifying properties. However, during the last decades, two prominent areas have been developed considering their multiple inhibitory abilities and efficiency as vaccine adjuvants.

**Use as inhibitory agent**

*Quillaja* saponins and their phenolic fractions have shown a wide range of biological activities including antiallergic, antifungal, antimicrobial, anticholesterolomic, antinociceptive, anti-inflammatory, analgesic, antipyretic, antiprotozoal, molluscicide, and antiviral. Table 1 summarizes a list of studies concerning these domains, remarking the model of study used and objectives. Furthermore, QS saponins were utilized as
biopesticide against parasitic nematodes [San Martin et al., 2005], aphids [De Geyter et al., 2012], and as a growth inhibitor of *Aedes aegypti* and *Culex pipiens* larvae [Pelah et al., 2002]. One of the explanations of this inhibitory effect is the lytic activity of saponins on cell membranes [Sparg et al., 2004; de Geyter et al., 2012] as well as their capacity to change the membrane fluidity and allow macromolecules to get absorbed to or through the cells of an organism [Podolak et al., 2010; Antolak et al., 2018]. In other cases, this effect was useful to prevent the attachment or viral infection through cell membranes [Roner et al., 2007; Tam and Roner, 2011] or produce membrane disorders in several organisms at high doses [Sen et al., 1998; Pelah et al., 2002]. On the other hand, negative consequences have been reported during the application of saponins-based biopesticides in aquatic environments [Hassan et al., 2013; Jiang et al., 2018a, b], revealing the importance of dosage control, and on the development of beneficial yeast microflora in vineyards after treatment against *Botrytis cinerea* and *Xiphinema index* [Fischer et al., 2011].

The effects of *Quillaja* saponins on food digestibility have been analyzed in animal models. *In vitro* experiences using rumen fermentation systems have demonstrated a positive result of *Quillaja* saponins on the digestibility efficiency in ruminants [Pen, 2007; Bunthoeun et al., 2008; Patra et al., 2012]. The same models reported the effect of QS extract in combination with Yucca schidigera saponins, reducing CH, and CO emissions [Makkar et al., 1998, Bunthoeun et al., 2008]. In the case of fish species, such as *Nila tilapia* or *Cyprinus carpio*, *Quillaja* saponins regulated female fertility, oxygen consumption [Francis et al., 2002b], and food digestibility [Serrano, 2013]. *Cyprinus carpio*, in turn, improved growth, which was related with the lipophilic interactions of saponins with cell membranes, increasing their permeability [Francis et al., 2002b; Bunthoeun et al., 2008] and improving the nutrient intake [Francis et al., 2002a; Serrano, 2013].

**Use as vaccine adjuvants**

Saponins from both *Quillaja* species and their fractions exhibit outstanding adjuvant qualities for the development of animal and human vaccines. Table 2 summarizes a set of applications in this field. *Quillaja* saponins modulate the immune system of mammals used as adjuvants [Katayama and Mine, 2006; Gilabert-Oriol et al., 2015]. Nevertheless, *Quillaja* saponins could become unstable and induce undesirable hemolytic activity. To counteract those effects, saponins are normally used in combination with liposomes [Brunner et al., 2017] or with cholesterol and phospholipids to form a sort of self-assembled cage called the immune-stimulating complex (ISCOM) [Papenmüller et al., 2014; Cibulski et al., 2018]. Interestingly, the same affinity of *Quillaja* saponins to cholesterol is useful to fight cancer cells due to the higher membrane-cholesterol molecules content [Hu et al., 2010; Hassan et al., 2013]. Marciani (2018) and Fleck et al. (2019) have described detailed information on applications of *Quillaja* saponins in the vaccine industry and recent pre-clinical trials extensively elsewhere, respectively.

**Table 1** - List of selected studies analyzing the use of *Quillaja* saponins and associated extracts as inhibitory agents.

| Model          | Description                                                                 | Source                        |
|----------------|------------------------------------------------------------------------------|-------------------------------|
| Bacteria       | Use of QS saponins as an agent to enhance the disinfection process in the beverage industry, inhibiting the growth of Asaia spp in vitro. | Antolak et al. (2018)         |
|                | Use of QS and Yucca saponins to reduce E. coli growth.                      | Sen et al. (1998)             |
| Fungi          | Antifungal activity of QS saponins used as a component of cellulose nanofibrous membrane against *Penicillium roqueforti* and *Aspergillus ochraceus*. | Dietl et al. (2010)           |
|                | Activity of QS saponins and phenolic compounds against *Botrytis cinerea* in fresh strawberries coating. | Zúñiga et al. (2012)          |
| Protozoa       | Evaluation of antiprotozoal activity of QS saponins against *Trichomonas vaginalis*. | Rocha et al. (2012)           |
| Virus          | Application and determination of optimal concentration of QS saponins against viruses (vaccinia virus, herpes simplex virus 1, HIV-1, HIV-2, varicella-zoster virus and retrovirus). | Roner et al. (2007)           |
| Murine         | Analysis of QS and Yucca saponin effects on diabetic-induced rats, reporting hypocholesterolemic, hypoglycemic and antioxidant qualities of saponins. | Fidan and Dündar (2008)       |
|                | Analgesic/antinociceptive effects of QS saponins in two murine thermal models. | Arrau et al. (2011)           |
| Others         | Assessment of antiviral activity of QS saponins against Rhesus rotavirus, by inhibition of virus-host attachment. | Tam et al. (2011)             |
|                | Testing of topical anti-inflammatory activity of quillaic acid in an inflammatory-induced mice model. | Rodriguez-Díaz et al. (2011)  |
|                | Analysis of QS dry extract and their polyphenol molecules on cholesterol solubilization. | Vinarova et al., (2015)       |
|                | Evaluation of a polyphenol-rich QS extract to retard oxidation of chicken meat lipids. | Fellenberg et al., (2011)     |
|                | Analysis of anti-molluscicide activity of QS saponins in grey field slug.   | González-Cruz and San Martin (2013) |
|                | Study of antimicrobial activity of QS saponins in the leather soaking process. | Zengin (2013)                |
Table 2 – List of selected studies on utilization of Quillaja saponins as vaccine adjuvants.

| Focus/application | Description | Source |
|-------------------|-------------|--------|
| Polio vaccine     | Assessment of QB aqueous extract, QB-90 saponin fraction and Quil-A as adjuvants in a polio vaccine. | Costa et al. (2014) |
| Bovine viral diarrhea virus | Study of an aqueous extract and QB-80 saponin fraction from QB as vaccine adjuvant using bovine viral diarrhea virus antigen in a murine model. | Cibulski et al. (2017) |
| Porcine reproductive and respiratory syndrome virus | Evaluation of QS Quil-A saponin fraction in the up-regulation of immune genes against porcine reproductive and respiratory syndrome virus. | Charrentantanakul and Fabros (2018) |
| Immuno-response activation | Study of the activation of triterpenoid saponins in the activation of dendritic cells to induce immune responses. | Marciani (2018) |
| Anti-cancer | Evaluation of the potential of anticancer cholesterol-QS saponin nanoparticles against lymphoma cell lines. | Hu et al. (2010) |
| | Assessment of Blocking and Balancing particles (BBE) based in the QS saponin fraction Quil-A, against tumor cells of renal cell carcinoma. | Hassan et al. (2013) |
| Synthetic derived adjuvants | Analysis of Quillaja saponins limitations as vaccine adjuvant and evaluation of synthetic saponins. | Adams et al. (2010) |
| ISCOMs | Study of the Quil-A saponin fraction effects on liposomal phosphatidylcholine/cholesterol to form ISCOM matrices. | Paepenmüller and Müller-Goymann (2014) |
| Liposomes | Description of the mechanism of action of an ISCOM matrix using QB-90 saponin fraction. | Cibulski et al. (2018) |

Different saponin fractions have been characterized since 1970s, with emphasis on QS, and more recently on QB. *Quillaja saponaria* saponin fractions, such as QS21 or their commercial presentations (e.g. Quil-A adjuvant), are widely used for the development of animal vaccines. Their use in human vaccines is under evaluation (Cibulski et al., 2015; Gilabert-Oriol et al., 2015). QS21, the most purified fraction of the aqueous extract of QS, shows some problems, such as its low concentration, difficulty in extraction and purification and/or chemical instability (Adams et al., 2010; Brunner et al., 2017). These drawbacks have promoted the development of synthetic adjuvants for the QS-21 saponin fraction. Research on chemical synthesis is still in progress; however, remarkable progress about the structure and activity of the synthetic saponins has been achieved (Fernández-Tejada et al., 2016). In parallel, QB saponins have emerged as a powerful alternative to QS saponin fractions. Indeed, QB-90, a QB fraction, has shown similar activity in comparison to Quil-A, when it is used in ISCOMs, promoting a local and transient “immunocompetent environment” (Cibulski et al., 2018). These findings are especially important for conservation and management of *Quillaja* species since alternative fractions from both species could be used for the development of new vaccines (Fleck et al., 2009; Costa et al., 2014; Cibulski et al., 2018).

**Structure of Quillaja saponins**

*Quillaja* saponins are special due to their triterpene nature. They have two sugar moieties attached to a central triterpene glycone structure and are classified as bidesmosidic molecules (Van Setten and van de Werken, 1996). Detailed descriptions of molecular structures of QS and/or QB saponins have already been presented (Van Setten and van de Werken, 1996; Kite et al., 2004; Fleck et al., 2019; Wallace et al., 2017, 2019). The most common triterpene aglycone skeleton is quillaic acid (Figure 2), which is generally glycosylated at C-3 and C-28 (Bankefors et al., 2010; Fleck et al., 2019). The disaccharide residue attached to C-3, β-D-Galp-(1→2)-β-D-GlcAp, is frequently branched at O-3 of the glucuronic acid unit by α-L-Rhap or a β-D-Xylop unit. On the other hand, C-28 is linked to a complex oligosaccharide, generally composed by a disaccharide residue (α-L-Rhap-(1→2)-β-D-Fucp and different ornamentations. Some of these groups, such as 3,5-dihydroxy-6 methylloctanoic acid,
contribute to the adjuvant activity of *Quillaja* saponins (Kite et al., 2004; Fleck et al., 2019). Saponins from leaves or wood differ from those present in bark, in terms of the different terminal attached molecules (see R-groups in Table 3). According to recent studies, 58 saponins in QS bark have been reported (Fleck et al., 2019). For QB, it has been recently described 48 saponin in leaves and 54 saponins in bark (Wallace et al., 2017, 2019). Additionally, aqueous extracts of QS contain a set of other components including calcium, oxalate, proteins, sugars, and tannins (Maier et al., 2015). These components are important in determining physical characteristics, such as capacity to form foam (San Martin and Briones, 2000).

**Advances in the characterization of genes involved in saponin biosynthesis and *Quillaja* genomes**

Triterpene saponins are molecules with a defensive role in plants against biotic and/or abiotic stress factors.

![Figure 2 - General saponin bidesmosidic molecule in QS and QB (adapted from van Setten et al., 1996 and Wallace et al., 2019).](image)

These metabolites promote plant immunity against a wide range of insects, pathogens, and herbivores (Hussain et al., 2019). The level of saponins increases in response to environmental stress [e.g water deficit] and could be involved in adaptive mechanism to survive in adverse soil and climate conditions (Szakiel et al., 2011). Saponin formation is mediated by the jasmonic and salicylic acid signaling pathways and their respective transcription factors, which upregulate the expression of key genes involved in saponin accumulation (Costa et al., 2013; Yendo et al., 2014). Yendo et al. (2014) and Kuwahara et al. (2019) present a complete description of saponin biosynthesis and illustrate the molecular transformation of different metabolites across the pathways. Briefly, saponin biosynthesis takes place in cytosol and starts from isoprenoids through the mevalonate acid pathway. Consecutively, the formed molecules are transformed by enzymes of OSC [oxidosqualene cyclases], cytochrome P450 and UDP [uridine diphosphate glycosyltransferase] families, contributing to triterpenoids molecular diversity (Luo et al., 2011; Yendo et al., 2014; Kuwahara et al., 2019). In the case of quillaic acid, the OSC β-amyrin synthase catalyzes the production of β-amyrin. Finally, enzymes of P450 and UDP families transform the β-amyrin molecule, forming quillaic acid (Meesapyodsuk et al., 2007). The knowledge of the full biosynthesis pathway for quillaic acid is still incomplete and its elucidation is a major challenge for industrial applications of saponins (Shang and Huang, 2019). According to Kamstrup et al. (2000), differences in accumulation of these compounds depends on genetic variation, which affects different enzymatic activities.

Most of the genes implicated in saponin biosynthesis are ubiquitous in plants. According to most references, genes and enzymes of *Quillaja* species are similar to other β-amyrin derived triterpene accumulating species. Nevertheless, characterization of new genes, for

| R0 | R1 | R2 | R3 | R4 | Ra | Rb | Rc | Rd |
|----|----|----|----|----|----|----|----|----|
| Xyl | Api-Xyl | Glu | Rha | Ac | CHO | OH | OH | CH3 |
| Rha | Xyl | Api | MeBu | Fa-Ara-Rha | Fa-Ara | CH2OH | H | H | COOCH3 |
| H | Xyl | Xyl | H | Glu | OHMeHex | CH2OCOCH3 |
| Xyl | GluA | MeBu | CH2 |
| H | Ac | H |

| R0 | R1 | R2 | R3 | R4 | Ra | Rb | Rc | Rd |
|----|----|----|----|----|----|----|----|----|
| Xyl | Xyl | Glu | Glu | Fa-OH |
| Rha | Api | H | Rha | Fa-Ara |
| DeHex | Api-Xyl | Pen | Fa |
| H | Xyl | Xyl | Hex |
| H | H |
example, those belonging to UDP and P450 families, has two principal challenges: 1) identify genes belonging to multigenic families in a non-sequenced organism, and 2) select an affordable number of candidate sequences for studies on functional characterization [Augustin et al., 2011]. Overcome these limits is crucial to get new insights into saponin biosynthesis. Meanwhile, the increasing interest of industrial community on saponin applications and the reduction in genome sequencing costs could improve our current knowledge about the genetic basis underlying qualitative and quantitative traits determining saponin production.

QS and QB genome comprises 28 chromosomes [2n] [Kubitzki, 2007]. For QS, the genome size is estimated at 821.52 Mbp (~ 0.42 pg, 1C) [Garcia et al., 2010]. In 2015, 12 microsatellites were identified in the QS nuclear genome, which were used to analyze the pattern of genetic variation and its relationship with the spatial distribution and anthropogenic influence in Chile [Letelier et al., 2015]. In 2019, a draft of the QS plastome genome was released [Vizoso et al., 2019]. Its sequence length is 0.133 Mbp, harboring 112 putative genes, including those encoding Photosystems, RuBisCO, Cytochrome b/f complex, ribosomal subunits, among others. To the authors’ knowledge, there are no other reports about the QS or QB nuclear genome or plastome sequences. According to Vizoso et al. [2019], the photosynthetic rate and methyl jasmonate-associated defensive mechanisms could be indirectly associated to saponin accumulation. They related the saponin contents with abiotic factors, under which chloroplasts may act as environmental sensors, mediating the signaling required to increase the accumulation of triterpene saponins as a defensive response. Interestingly, Costa et al. [2013] reported the same accumulation pattern for QB and linked it to similar mechanisms, as Vizoso et al. [2019]. Recently, Matveeva and Otten [2019] suggested the ability of *Agrobacterium* to induce horizontal gene transfer in QS. In their study, the authors found four bacterial genes in a QS contig sequence database. Nevertheless, due to the incomplete coverage of the contigs, it is still too early for conclusions about the presence of bacterial genes in QS genome and the possibility of its natural transformation by *Agrobacterium*.

**Final Remarks**

*Quillaja* saponins are well documented from a biological and chemical perspective. The wide range of applications, particularly for QS saponins, has represented several contributions to the knowledge about these metabolites since 1950. Although “omic” technologies have experienced significant advances during the last decades, and important information has been accumulated for model species, there are still important gaps for *Quillaja* species. These include the sequence of nuclear genomes, expression analyses (at transcriptional or protein level) focused on identifying genes and enzymes involved in saponin biosynthesis and accumulation, and the characterization of genetic mechanisms underlying its quantitative and qualitative variation, among others. Considering the increasing demand of saponins by the food, chemical, and pharmacological industries, additional actions are required to support sustainable biomass production from both natural forests and dedicated plantations. Important progress has been reached in the management and domestication of *Quillaja* species during the last 50 years, especially with QS in Chile. This advance has permitted to develop different protocols for biomass harvesting and management from natural forests, plant production [by seed and micropropagation], plantation and selection of high-saponin productive clones. Basic information about genetic variation and heritability regarding traits that determine biomass production, tree architecture, wood properties, concentration and diversity of saponins (and other metabolites), among others, and the identification and conservation of superior germplasm, will also contribute to developing advanced genetic improvement programs. At the same, optimization of silvicultural techniques [in forest and/or agricultural sites], including soil and plant spacing management, use of systems for efficient water use, and the definition of biomass harvest cycles allow realizing the potential generated by breeding programs. These managed systems may take advantage of high biomass production and adaptability of *Quillaja* species and overcome current limitations of some non-tree species producing saponins, which are characterized by low biomass and use in food production. Progress in domestication and industrial uses of *Quillaja* species has been supported for governmental and private initiatives, and funding is still required in the short and mid-term. Policies promoting protection of *Quillaja* diversity, its forestry, as well as the development of new products or industrial processes are necessary for a sustainable use of this resource.

**Authors’ Contributions**

**Conceptualization:** Guerra, F.P.; Sepúlveda, S. **Software development:** Sepúlveda, S. **Writing and editing:** Guerra, F.P.; Sepúlveda, S.

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