



Plant extracts and compounds for combating schistosomiasis

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Abstract Schistosomiasis is a vector-borne water-based disease caused by *Schistosoma* blood flukes. It mostly affects people in low-income regions, 90% of reported cases being in developing countries. *Schistosoma* has a complex lifecycle, alternately infecting mammalian hosts and snails. The snails hosting the parasite are the most viable targets. Selective preparations for reducing the parasite pool in snails and infected water are required as current molluscicides are also nontoxic to other organisms, including fish, and thus affect food supplies in infected areas. Plants (e.g. *Annona crassiflora* Mart., *A. muricata* L., and *A. montana* Macfad.) are attractive potential sources as

alternative molluscicides and novel entity to treat the disease owned to their diverse biologically potent compounds including; saponins, alkaloids, terpenoids, and tannins. Additionally, they can be locally cultivated, providing income for farmers and reducing treatment costs. Here, we review plants, plant extracts and isolated compounds that have shown activities against the host snails or *Schistosoma* in various parts of its life cycle. Plants have a lot of potential and will continue to contribute feasible, effective medicines and/or pesticides; more research is warranted to fully explore their future applications.

Dedicated with great pleasure and honor to Prof. Lars Bohlin, Uppsala University, Sweden and Prof. Rob Verpoorte, Leiden University, Netherland on the occasion of their 75th birthday.

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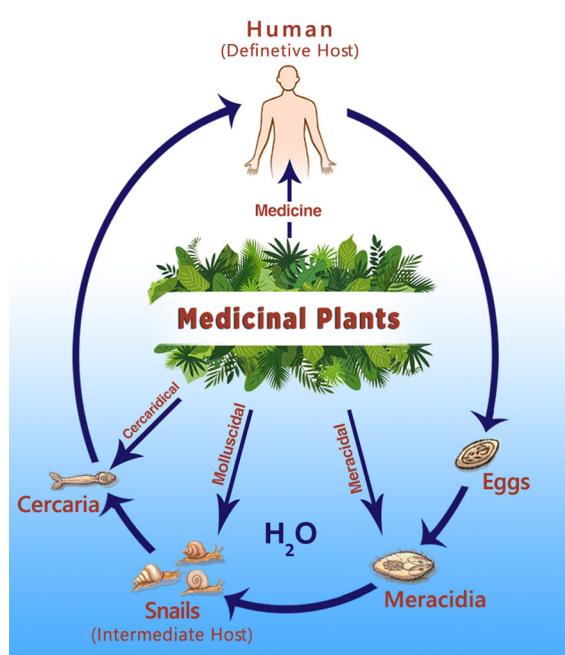
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Graphical abstract



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Introduction

Schistosomiasis, a vector-borne water-based disease caused by *Schistosoma* blood flukes, is one of the most devastating parasitic diseases in tropical and subtropical regions (Walz et al. 2015). It is regarded by World Health Organization (WHO) as the third most common tropical disease, following malaria and intestinal helminthiasis (Murungi et al. 2021). It is also recognized as one of the neglected tropical diseases with high morbidity and mortality rates, affecting a billion of the world's poorest people, mainly in the developing countries (Pereira et al. 2011). The disease is an underappreciated global burden and linked geographically to poverty, and poor health services (Payne and Fitchett 2010). It mostly affects young adults in their productive years, and children younger than 10 years. Untreated infections in young people impair their immunity, increasing their vulnerability to growth retardation, malnutrition and cognitive disorders (Dabo et al. 2011). More than 500,000 people die every year from this parasitic disease (Botelho et al. 2009).

Schistosoma has a complex lifecycle, alternately infecting mammalian hosts and snails especially *S. japonicum* which spreads via all kinds of mammals (Nelwan 2019). One in 30 of the total human

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population surveyed in endemic areas were apparently infected by *Schistosoma* flukes via contaminated water or direct contact with mammals hosting the parasite (Sheir et al. 2001). Molluscicides have a long history of both failures and successes in the control of schistosomiasis (Fenwick et al. 2006). Their application is now considered a key element that has significantly contributed to the decline in schistosomiasis infection and morbidity rates in a number of African countries during the last decades (Brackenbury and Appleton 1997). Chemotherapy and chemoprophylaxis with, praziquantel (PZQ), currently play a crucial role in curing, controlling and preventing the disease (Yousif et al. 2007). PZQ is highly valuable, due to its broad activity spectrum, for instance, it has a potent effect against *S. mansoni* especially at 6–7 weeks old infections. PZQ is thought to target the β subunits of voltage-gated Ca^{2+} channels in *Schistosoma* (Doenhoff et al. 2008). Despite the success, resistance to PZQ is apparently emerging and exacerbating challenges in the management of schistosomiasis globally (Melman et al. 2009). Pinto-Almeida and his colleagues analyze the proteome of *S. mansoni* PZQ-resistant adult worms and compare it with its parental fully PZQ-susceptible strain, using a high throughput LC–MS/MS identification. The results revealed that different proteins of the *S. mansoni* proteome in worms, were downregulated after the exposure to PZQ (Pinto-Almeida et al. 2018). Cotton and Doyle recently identified a gene responsible for PZQ resistance in experimentally selected resistant *S. mansoni*. This study shows that variation at or near Sm. (*S. mansoni*) TRPMPZQ is associated with the resistance (Cotton and Doyle 2022).

Nature has always been a valuable source of drugs and continues to deliver important drug leads (Tulp and Bohlin 2004). WHO estimates that 65–80% of the developing world's population relies on traditional medicine to meet primary health care needs (Kumar et al. 2021) and traditional medicinal plants have

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supplied numerous pharmacologically active compounds (Yousif et al. 2007). Thus, considerable efforts have been made to identify plants products that are environmentally safe, non-toxic and selectively active for the integrated control of schistosomiasis, and several of them have shown promising results. For example, we have recently shown that the Egyptian medicinal plant *Asparagus stipularis* Forssk. has anti-schistosomal activity (El-Seedi et al. 2012). Recent reports documented the potential of some natural products such as *Pulsatilla chinensis* (Bunge) Regel extracts (Table 1) and its molluscicidal activity against *O. hupensis*. They are less harmful to non-target aquatic organisms than the reference molluscicide niclosamide (Chen et al. 2012). In addition, *Agave attenuata* Salm-Dyck is toxic to the target snail *Bulinus africanus* but has fewer (or no) adverse effects on fish and mammals (Brackenbury and Appleton 1997).

Both synthetic molluscicides and molluscicides derived from plants have made progress in the field of host snails eradication. The rising expense and toxicity to non-target organisms of synthetic molluscicides has rekindled interest in plant-based molluscicides (Clark et al. 1997; Zheng et al. 2021).

This review highlights the use of medicinal plants as new alternatives that could either complement or replace conventional control approaches. Potential molluscicidal plant species and their bioactive constituent based on traditional uses (Tables 2 and 3), are listed, in accordance to the literature survey done on the biological control of *Schistosoma* in the field using natural products, at all life stages.

Life cycle of *Schistosoma* species

There are three main species of human helminth parasites of the genus *Schistosoma*: *S. haematobium*, found in Africa and Asia; *S. mansoni*, present in Africa, the Middle-East and South America; and *S. japonicum*, endemic in China and Southwest Asia (Colley et al. 2014). There are also two minor species, *S. intercalatum* and *S. mekongi*, found in West and central Africa (Colley et al. 2014). The adult worms are live in mesenteric vessels (*S. mansoni* and *S. japonicum*) or the vessels surrounding the bladder (*S. haematobium*). The females produce large numbers of

Table 1 Bioactive plant extracts reported to have molluscidal activity against intermediate vector snails

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Acanthaceae</i>						
<i>Acanthus mollis</i> L.-leaves	Bear's Breeches leaves	Petroleum ether	ND	LC ₅₀ = 6.92 µg/ml LC ₉₀ = 15.5 µg/ml after 24 h	+++ ++	Ahmed and Rifaat (2004)
<i>Acanthaceae + Iridaceae</i>						
Binary combination (1:1) of <i>A. mollis</i> <i>L.</i> + <i>Iris pseudacorus</i> L.	Bear's Breeches + Yellow iris L. + Iris pseudacorus L.	Petroleum ether	ND	LC ₅₀ = 3.8 µg/ml after 24 h	+++	Ahmed and Rifaat (2004)
<i>Acanthaceae + Solanaceae</i>						
Binary combination (1:1) of <i>A. mollis</i> <i>L.</i> + <i>Solanum nigrum</i> L.	Bear's Breeches + Black nightshade L. + Solanum nigrum L.	Petroleum ether	ND	LC ₅₀ = 5.1 µg/ml LC ₉₀ = 10.3 µg/ml after 24 h	+++ ++	Ahmed and Rifaat (2004)
<i>Agavaceae</i>						
<i>Agave filifera</i> Salm-Dyck-whole plant	Thread-leaf Agave, Thread Agave whole plant	Aqueous-suspension	Saponins, catechins, tannins and carbohydrates	LC ₅₀ = 42.3 µg/ml LC ₉₀ = 85.6 µg/ml after 24 h	++ ++	Rawi et al. (2011)
<i>(B. alexandrina)</i>						
Cold-aqueous		Saponins, catechins, tannins and carbohydrates		LC ₅₀ = 463.0 µg/ml LC ₉₀ = 692.7 µg/ml after 24 h	+ +	
Boiled-aqueous		Saponins, catechins, tannins and carbohydrates		LC ₅₀ = 257.0 µg/ml LC ₉₀ = 328.9 µg/ml after 24 h	+ +	(<i>B. alexandrina</i>)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Amaryllidaceae</i>						
<i>Crinum zeylanicum</i> L.-whole parts	Crinum Lily	Aqueous	Steroids, alkaloids, catechic tannins, flavonoids, albuminoids, phenols, lipoids, aldehydes, mucilages and terpenoids	LC ₀ = 10 mg/ml LC ₅₀ = 50 mg/ml LC ₁₀₀ = 100 mg/ml (<i>B. pfeifferi</i>) after 24 h	— — — —	Chifundira et al. (1993)
		Ethanol	Steroids, alkaloids, catechic tannins, flavonoids, albuminoids, phenols, lipoids, aldehydes, mucilages and terpenoids	LC ₀ = 5 mg/ml LC ₅₀ = 50 mg/ml LC ₁₀₀ = 100 µg/ml (<i>B. pfeifferi</i>) after 24 h	— — — —	
<i>Annonaceae</i>						
<i>Annona crassiflora</i> Mart.-pulp and seeds	Araticum & marolo	Ethanol	Acetogenins	LC ₁₀ = 0.2 µg/ml LC ₅₀ = 1.67 µg/ml LC ₉₀ = 13.2 µg/ml (<i>B. glaberrata</i> -adult) after 24 h	+++ +++ ++ +++ —	dos Santos and Sant'Ana (2001)
		Ethanol	Acetogenins	LC ₁₀ = 1.0 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 1.0 µg/ml (<i>B. glaberrata</i> -egg)	+++ +++ ++ —	dos Santos and Sant'Ana (2001)
<i>A. crassiflora</i> Mart.-stems	Araticum & marolo	Ethanol	Acetogenins	LC ₁₀ = 0.4 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 2.3 µg/ml (<i>B. glaberrata</i> -adult) after 24 h	+++ +++ ++ +++ —	dos Santos and Sant'Ana (2001)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>A. crassiflora</i> Mart.-roots wood	Araticum & marolo	Ethanol	Acetogenins	LC ₁₀ = 8.3 µg/ml LC ₅₀ = 16.9 µg/ml LC ₉₀ = 34.5 µg/ml (<i>B. glabra</i> -adult)	++ ++ ++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1.0 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 1.0 µg/ml	+++ +++ +++	
				(<i>B. glabra</i> -egg)		
				after 24 h		
				LC ₁₀ = 0.3 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 3.8 µg/ml	+++ +++ +++	
				(<i>B. glabra</i> -adult)	+++	
				LC ₁₀ = 1.0 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 1.0 µg/ml	+++ +++ +++	
				(<i>B. glabra</i> -egg)		
				after 24 h		
				LC ₁₀ = 5.0 µg/ml LC ₅₀ = 9.2 µg/ml LC ₉₀ = 17.0 µg/ml	++ ++ ++	
				(<i>B. glabra</i> -adult)	+++	
				LC ₁₀ = 1.0 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 1.0 µg/ml	+++ +++ +++	
				(<i>B. glabra</i> -egg)		
				after 24 h		
<i>Annona glabra</i> L.- seeds	Pond-apple	Ethanol	Acetogenins	LC ₁₀ = 5.0 µg/ml LC ₅₀ = 9.2 µg/ml LC ₉₀ = 17.0 µg/ml	++ ++ ++	dos Santos and Sant'Ana (2001)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>A. glabra</i> L.-leaves	Pond-apple	Ethanol	Acetogenins	LC ₁₀ = 61.7 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 3.8 µg/ml (<i>B. glaberrata</i> -adult)	++ +++ +++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1.0 µg/ml LC ₅₀ = 1.0 µg/ml LC ₉₀ = 1.0 µg/ml	+++ +++ +++	
				(<i>B. glaberrata</i> -egg)		
				LC ₁₀ = 0.3 µg/ml LC ₅₀ = 1.6 µg/ml LC ₉₀ = 8.8 µg/ml (<i>B. glaberrata</i> -adult)	+++ +++ ++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 9.2 µg/ml LC ₅₀ = 20.3 µg/ml LC ₉₀ = 44.8 µg/ml (<i>B. glaberrata</i> -egg)	++ ++ ++	
				after 24 h LC ₈₀ ≥ 100 µg/ml (<i>B. glaberrata</i>)	+	de S. Luna et al. (2005)
				LC ₁₀ = 0.25 µg/ml LC ₅₀ = 0.97 µg/ml LC ₉₀ = 3.8 µg/ml (<i>B. glaberrata</i> -adult)	+++ +++ +++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1 µg/ml LC ₅₀ = 1 µg/ml LC ₉₀ = 1 µg/ml (<i>B. glaberrata</i> -egg)	+++ +++ +++	
				after 24 h		
<i>Annona muricata</i> L.	Graviola,soursop and gunbanana -leaves	Ethanol	Acetogenins	Flavonoids, xanthones and steroids		
<i>A. muricata</i> L.- stems bark	Graviola,soursop and gunbanana	Ethanol				

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Annona montana</i> Macfad Mart.-leaves	Sinini	Ethanol	Acetogenins	LC ₁₀ = 22.6 µg/ml LC ₅₀ = 62.7 µg/ml LC ₉₀ = 173.9 µg/ml (<i>B. glabra</i> -adult)	++ ++ + ++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 61.6 µg/ml LC ₅₀ = 91.6 µg/ml LC ₉₀ = 136.2 µg/ml (<i>B. glabra</i> -egg)	++ + +	
<i>A. montana</i> -stems bark	Sinini	Ethanol	Acetogenins	LC ₁₀ = 0.55 µg/ml LC ₅₀ = 1.9 µg/ml LC ₉₀ = 6.2 µg/ml (<i>B. glabra</i> -adult)	+++ +++ ++ +++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1 µg/ml LC ₅₀ = 1 µg/ml LC ₉₀ = 1 µg/ml (<i>B. glabra</i> -egg)	+++ +++ ++ after 24 h	
<i>Annona salzmannii</i> A.DC.-leaves	Araticum-da-mata" and "araticumapé	Ethanol	Acetogenins	LC ₁₀ = 0.66 µg/ml LC ₅₀ = 0.95 µg/ml LC ₉₀ = 1.37 µg/ml (<i>B. glabra</i> -adult)	+++ +++ ++ +++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1 µg/ml LC ₅₀ = 1 µg/ml LC ₉₀ = 1 µg/ml (<i>B. glabra</i> -egg)	+++ ++ + after 24 h	

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Annona squamosa</i> L.-seeds	Custard apple	Ethanol	Acetogenins	LC ₁₀ = 0.3 µg/ml LC ₅₀ = 7.5 µg/ml LC ₉₀ = 164.02 µg/ml (<i>B. glabra</i> -adult)	+++ ++ + +++	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1 µg/ml LC ₅₀ = 1 µg/ml LC ₉₀ = 1 µg/ml (<i>B. glabra</i> -egg)	+++ +++ +++	
				LC ₁₀ = 5.8 µg/ml LC ₅₀ = 14.0 µg/ml LC ₉₀ = 34.4 µg/ml (<i>B. glabra</i> -adult)	++ ++ ++	dos Santos and Sant'Ana (2001)
<i>A. squamosa</i> L.-stems/park	Custard apple	Ethanol	Acetogenins	LC ₁₀ = 26.4 µg/ml LC ₅₀ = 51.7 µg/ml LC ₉₀ = 101.4 µg/ml (<i>B. glabra</i> -egg)	++ + +	
				LC ₁₀ = 1.7 µg/ml LC ₅₀ = 3.8 µg/ml LC ₉₀ = 8.55 µg/ml (<i>B. glabra</i> -adult)	+++ +++ ++	dos Santos and Sant'Ana (2001)
<i>A. squamosa</i> L.-roots	Custard apple	Ethanol	Acetogenins	LC ₁₀ = 16.8 µg/ml LC ₅₀ = 28.8 µg/ml LC ₉₀ = 49.2 µg/ml (<i>B. glabra</i> -egg)	++ ++ +	
				after 24 h		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>A. squamosa</i> L.-leaves	Custard apple	Ethanol	Acetogenins	LC ₁₀ = 5.4 µg/ml LC ₅₀ = 44.6 µg/ml LC ₉₀ = 368.0 µg/ml (<i>B. glabrat-a-adult</i>)	++ ++ +	dos Santos and Sant'Ana (2001)
				LC ₁₀ = 1 µg/ml LC ₅₀ = 1 µg/ml LC ₉₀ = 1 µg/ml (<i>B. glabrat-a-egg</i>)	+++ +++ +++	
				after 24 h		
<i>Apiaceae (Umbelliferae)</i>						
<i>Anni mequs</i> L.-flowers	Khella shytani or bullwort	Aqueous-suspension	Saponins, catechins, tannins and carbohydrates	LC ₅₀ = 747.7 µg/ml LC ₉₀ = 1370.3 µg/ml after 24 h	+	Rawi et al. (2011)
				(<i>B. alexandrina</i>)		
				LC ₅₀ = 2126.5 µg/ml LC ₉₀ = 6775.2 pm after 24 h	—	Rawi et al. (2011)
				(<i>B. alexandrina</i>)		
<i>A. majus</i> L.-leaves	Khella shytani or bullwort	Boiled-aqueous	Saponins, catechins, tannins and carbohydrates	LC ₅₀ = 944.9 µg/ml LC ₉₀ = 1394. µg/ml after 24 h	—	Rawi et al. (2011)
				(<i>B. alexandrina</i>)		
				LC ₅₀ = 738.3 µg/ml LC ₉₀ = 1157.2 µg/ml after 24 h	—	Rawi et al. (2011)
				(<i>B. alexandrina</i>)		
				LC ₅₀ = 4731.7 µg/ml LC ₉₀ = 10,377.1 µg/ml after 24 h	—	Rawi et al. (2011)
				(<i>B. alexandrina</i>)		

Table 1 continued

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>A. majus</i> L.-roots	Khella shytani or bullwort	Aqueous	Saponins	LC ₅₀ (N) = 22.500 mg/ml (<i>B. alexandrina</i>)	—	Rawi et al. (1996)
				LC ₅₀ (N) = 8.400 mg/ml (<i>Bu. truncatus</i>)	—	
				LC ₅₀ (T) = 10.600 mg/ml (<i>B. alexandrina</i>)	+	
				LC ₅₀ (T) = 5.900 mg/ml (<i>Bu. truncatus</i>)	+	
				after 4 w	+	
				LC ₅₀ (N) = 1 mg/ml (<i>B. alexandrina</i>)	+	Rawi et al. (1996)
				LC ₅₀ (N) = 720 µg/ml (<i>Bu. truncatus</i>)	+	
				LC ₅₀ (T) = 815 µg/ml (<i>B. alexandrina</i>)	+	
				LC ₅₀ (T) = 390 µg/ml (<i>Bu. truncatus</i>)	+	
				afier 4 w	+	
				LC ₅₀ > 500 µg/ml after 24 h	+	Mahmoud et al. (2011)
				(<i>B. alexandrina</i> -adult)		
				LC ₅₀ = 19.45.2 mg/ml LC ₉₀ = 7.706 mg/ml after 24 h	—	Clark and Appleton (1997)
				(<i>Bu. aficanus</i>)		
<i>Apocynaceae</i>						
<i>Carissa carandas</i>	Karonda	Dry leaves	ND			
L.-leaves						
<i>Rauvolfia caffra</i>	Quinine tree	Aqueous	ND			
Sond.-leaves						

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Arecales</i>						
<i>Areca catechu</i> L.- seeds	Betel nut	Ethanol	Alkaloids	LC ₅₀ = 17.2 µg/ml after 24 h LC ₅₀ = 14.1 µg/ml after 48 h LC ₅₀ = 11.3 µg/ml after 72 h LC ₅₀ = 8.1 µg/ml after 96 h	++ ++ ++ ++	Jaiswal and Singh (2008)
<i>Lymnaea acuminata</i>						
Powder		Alkaloids		LC ₅₀ = 27.2 µg/ml after 24 h LC ₅₀ = 23.3 µg/ml after 48 h LC ₅₀ = 17.5 µg/ml after 72 h LC ₅₀ = 12.3 µg/ml after 96 h	++ ++ ++ ++	Jaiswal and Singh (2008)
Chloroform		Alkaloids		LC ₅₀ = 37.2 µg/ml after 24 h LC ₅₀ = 28.5 µg/ml after 48 h LC ₅₀ = 25.5 µg/ml after 72 h LC ₅₀ = 14.7 µg/ml after 96 h	++ ++ ++	Jaiswal and Singh (2008)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
		Ether	Alkaloids	LC ₅₀ = 17.7 µg/ml after 24 h	++	Jaiswal and Singh (2008)
				LC ₅₀ = 15.9 µg/ml after 48 h	++	
				LC ₅₀ = 13.6 µg/ml after 72 h	++	
				LC ₅₀ = 11 µg/ml after 96 h		
		(<i>L. acuminata</i>)				
		Acetone	Alkaloids	LC ₅₀ = 36 µg/ml after 24 h	++	Jaiswal and Singh (2008)
				LC ₅₀ = 28 µg/ml after 48 h	++	
				LC ₅₀ = 20.3 µg/ml after 72 h	++	
				LC ₅₀ = 13 µg/ml after 96 h		
		(<i>L. acuminata</i>)				
		Column purified	Alkaloids	LC ₅₀ = 11.3 µg/ml after 24 h	++	Jaiswal and Singh (2008)
				LC ₅₀ = 8.4 µg/ml after 48 h	++	
				LC ₅₀ = 6.7 µg/ml after 72 h	++	
				LC ₅₀ = 4 µg/ml after 96 h	+++	
		(<i>L. acuminata</i>)				
Asclepiadaceae						
<i>Marsdenia</i>	Cipó-Seda	Ethanol	Flavonoids, flavonones, flavones, xanthones, pentacyclic triterpenes and leucoanthocyni-dins	LC ₅₀ ≥ 100 µg/ml (<i>B. glabrata</i> -adult)	+	de S. Luna et al. (2005)
<i>alitissima</i> (Jacq.)						
Dugand-						
woods bark						

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Calotropis procera</i> -Dry and-whole overground parts	Apple of Sodom	Cold-aqueous suspension	ND	LC ₀ = 24.3 µg/ml LC ₁₀ = 98 µg/ml LC ₂₅ = 134 µg/ml LC ₅₀ = 243 µg/ml LC ₉₀ = 360 µg/ml after 24 h	++ ++ + + +	Bakry (2009)
				(<i>B. alexandrina</i>)		
<i>Asparagaceae</i>						
<i>Anemarrhena asphodeloides</i> Bunge-rhizomes	Anemarrhena rhizome, Zhi mu	<i>n</i> -Butanol	Saponins and steroid saponins	LD ₉₈ = 5 µg/ml after 72 h	++	Cui et al. (1999)
				(<i>O. hupehensis</i>)		
<i>Cratylia floribunda</i> Benth.-seeds	Cratylia	<i>n</i> -Hexane	Lectins	LC ₁₀ = 12.9 µg/ml LC ₅₀ = 25.5 µg/ml LC ₉₀ = 50.30 µg/ml	++ ++ ++	dos Santos et al. (2010)
				(<i>B. glabra</i>)		
<i>Dioclea guianensis</i> Benth.-seeds		<i>n</i> -Hexane	Lectins	LC ₁₀ = 13 µg/ml LC ₅₀ = 23.1 µg/ml LC ₉₀ = 41 µg/ml	++ ++ ++	dos Santos et al. (2010)
				(<i>B. glabra</i>)		
<i>Aspidiaceae</i>						
<i>Elaphoglossum piloselloides</i> (C. Presl) T. Moore-	—	Diethyl ether	Bicyclic phloroglucinols	LC ₅₀ = 19.6 µg/ml LC ₁₀₀ ≤ 50.0 µg/ml	+ +	Socolsky et al. (2009)
rhizomes, roots and sterile fronds				(<i>B. peregrine</i>)		
<i>Asteraceae (Compositae)</i>						
<i>Achyrocline satureoides</i> (Lam.) DC.-aerial parts	Macela or marcela	Ethanol	Terpenoids, phenylpropanoids, flavonoids	LC ₉₀ = 43 µg/ml after 24 h	++	Mendes et al. (1999)
				(<i>B. glabra</i>)		
<i>Actinoeris angustifolia</i> (Gardner) Cabral-aerial parts	—	Ethanol	ND	LC ₉₀ = 33 µg/ml after 24 h	++	Mendes et al. (1999)
				(<i>B. glabra</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Alomia myriadenia</i> Sch.Bip. ex Baker-aerial parts	—	Ethanol	ND	LC ₉₀ = 33 µg/ml after 24 h <i>(B. glabra)</i>	++	Mendes et al. (1999)
<i>Ambrosia maritima</i> L. ^c aerial parts	Damasissa or sea ragweed	Chloroform	Sesquiterpene lactone	LC ₉₀ = 3,000 mg/ml <i>(L. cailliaudi)</i> LC ₁₀₀ = 200 µg/ml <i>(B. glabra)</i>	+ —	Slacanin et al. (1988), Abou Basha et al. (1994)
<i>Artemisia herba-alba</i> Asso-aerial parts	Shih or desert wormwood	Petrol	Phenols, flavonoids, alkaloids, sterols, terpenoids and essential oils	LC ₅₀ = 62.9 µg/ml LC ₉₀ = 92 µg/ml after 24 h <i>(Bu. truncatus)</i>	++ ++	Hmamouchi et al. (2000)
<i>n</i> -Hexane			Phenols, flavonoids, alkaloids, sterols, terpenoids and essential oils	LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h <i>(Bu. truncatus)</i>	+ +	Hmamouchi et al. (2000)
Dichloromethane		Phenols, flavonoids, alkaloids, sterols, terpenoids and essential oils		LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h <i>(Bu. truncatus)</i>	++ +	Hmamouchi et al. (2000)
Ethyl acetate		Phenols, flavonoids, alkaloids, sterols, terpenoids and essential oils		LC ₅₀ = 62.9 µg/ml LC ₉₀ = 92 µg/ml after 24 h <i>(Bu. truncatus)</i>	++ ++	Hmamouchi et al. (2000)
Methanol		Phenols, flavonoids, alkaloids, sterols, terpenoids and essential oils		LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h <i>(Bu. truncatus)</i>	++ +	Hmamouchi et al. (2000)
Aqueous		Phenols, flavonoids, alkaloids, sterols, terpenoids and essential oils		LC ₅₀ = 2,598 mg/ml LC ₉₀ = 2,922 mg/ml after 24 h <i>(Bu. truncatus)</i>	— —	Hmamouchi et al. (2000)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Calendula officinalis</i> L.- flowers	African Marigold, Calendula, Common Marigold, Garden Marigold, Marigold, Pot Marigold, Zergul in India, Butterblume in Germany, Chin Chan Ts'ao in China, Galbinea in Romania, and Ringblomma in Sweden	Aqueous	ND	LC ₅₀ (N) = 98 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (N) = 81 µg/ml (<i>Bu. truncatus</i>) LC ₅₀ (D) = 50 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (D) = 18 µg/ml (<i>Bu. truncatus</i>)	++ ++ ++ ++	Rawi et al. (1996)
<i>C. officinalis</i> L.- leaves		Aqueous	ND	LC ₅₀ (N) = 10 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (N) = 81 µg/ml (<i>Bu. truncatus</i>) LC ₅₀ (D) = 67 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (D) = 40 µg/ml (<i>Bu. truncatus</i>)	+++ ++ ++ ++	Rawi et al. (1996)
<i>C. officinalis</i> L.- roots		Aqueous	ND	LC ₅₀ (N) = 205 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (N) = 183 µg/ml (<i>Bu. truncatus</i>) LC ₅₀ (D) = 137 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (D) = 109 µg/ml (<i>Bu. truncatus</i>)	++ + + +	Rawi et al. (1996)
<i>C. officinalis</i> L.- stems		Aqueous	ND	LC ₅₀ (N) = 250 µg/ml (<i>B. alexandrina</i>) LC ₅₀ (N) = 210 µg/ml (<i>Bu. truncatus</i>) LC ₅₀ (D) = 98 µg/ml (<i>B. alexandrina</i>)	++ ++ ++	Rawi et al. (1996)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results LC_{50} (T) = 86 $\mu\text{g/ml}$	Activity	References
<i>(Bu. truncatus)</i>						
<i>Chronolaena odorata</i> (L.) R.M.King & H.Rob.-leaves	Siam weed, devil weed, French weed, communist weed, hagonoy, co ho	Ethanol	ND	after 4 w $LC_{50} = 44 \mu\text{g/ml}$ $LC_{90} = 119.3 \mu\text{g/ml}$	++	Benson (2012)
		Aqueous	ND	after 24 h $LC_{50} = 65.8 \mu\text{g/ml}$ $LC_{90} = 139.5 \mu\text{g/ml}$	++	Benson (2012)
<i>Chrysanthemum viscidolehmannii</i> (Schott) Theil.-aerial parts	Uqhuwan	Petrol	Flavonoids and terpenoids	after 24 h $LC_{50} = 4.4 \mu\text{g/ml}$ $LC_{90} = 8.4 \mu\text{g/ml}$	+++	Hmamouchi et al. (2000)
		<i>n</i> -Hexane	Flavonoids and terpenoids	after 24 h $LC_{50} = 14.1 \mu\text{g/ml}$ $LC_{90} = 28 \mu\text{g/ml}$	++	Hmamouchi et al. (2000)
Dichloromethane	Flavonoids and terpenoids	<i>(Bu. truncatus)</i>	$LC_{50} = 5.9 \mu\text{g/ml}$ $LC_{90} = 14.1 \mu\text{g/ml}$	after 24 h $LC_{50} = 891 \mu\text{g/ml}$ $LC_{90} = 1342 \mu\text{g/ml}$	+++	Hmamouchi et al. (2000)
		Aqueous	Flavonoids and terpenoids	after 24 h $LC_{50} = 300 \mu\text{g/ml}$ $LC_{10} = 2.300 \text{ mg/ml}$ $LC_{25} = 2.600 \text{ mg/ml}$ $LC_{50} = 3.000 \mu\text{g/ml}$ $LC_{90} = 4100 \text{ mg/ml}$	+	Bakry (2009)
<i>Conyza dioscoridis</i> (L.) Desf.-whole overground parts	—	Cold-aqueous suspension	ND	after 24 h $LC_{50} = 3.000 \mu\text{g/ml}$ $LC_{90} = 4100 \text{ mg/ml}$	—	
<i>(B. alexandrina)</i>						

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Echinops ellenbeckii</i> O. Hoffm.-roots	Kebercho	Methanol	Polyphenols and carotenoids	LC ₁₀₀ = 20.25 µg/ml after 24 h (<i>B. pfeifferi</i>)	++	Hymete et al. (2005)
<i>Echinops longistylus</i> A. Rich.-roots	-	Methanol	Alkaloids, saponins, phytosterols, polyphenols and carotenoids	LC ₁₀₀ = 4 µg/ml after 24 h (<i>B. pfeifferi</i>)	+++	Hymete et al. (2005)
<i>Piptocarpha rotundifolia</i> (Less.) Baker-aerial parts	Ashdaisy	Ethanol	ND	LC ₉₀ = 99 µg/ml after 24 h (<i>B. glabratia</i>)	++	Mendes et al. (1999)
<i>Vanillosmopsis erythronappa</i> (DC.) Sch.Bip.-aerial parts	Candeia	Ethanol	ND	LC ₉₀ = 99 µg/ml after 24 h (<i>B. glabratia</i>)	++	Mendes et al. (1999), Silvério et al. (2013)
<i>Verbesina clausenii</i> Sch.Bip. ex Baker-aerial parts	-	Ethanol	ND	LC ₉₀ = 78 µg/ml after 24 h (<i>B. glabratia</i>)	++	Mendes et al. (1999)
<i>Buddlejaceae</i>						
<i>Buddleja lindleyana</i> Fortune- Leaves	-	Ethanol	ND	LC ₅₀ = 39.9 µg/ml after 48 h (<i>O. hupensis</i>)	++	Han et al. (2010)
				LC ₉₀ = 59.3 µg/ml after 48 h (<i>O. hupensis</i>)	++	
<i>Burseraceae</i>						
<i>Commiphora molmol</i> (Engl.) Engl. ex Tschirch-stems	Myrrh	Oil	Essential oil	LC ₅₀ = 5 µg/ml (<i>L. natalensis</i>)	++	Masoud et al. (2000)
				LC ₅₀ = 1.5 µg/ml (<i>B. alexandrina</i>)	++	
				LC ₅₀ = 8.5 µg/ml (<i>Bu. truncatus</i>)	++	
				after 24 h LC ₅₀ = 4 µg/ml (<i>L. natalensis</i>)	+++	
				LC ₅₀ = 6 µg/ml (<i>B. alexandrina</i>)	++	

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Cannaceae</i>				$LC_{50} = 7 \mu\text{g/ml}$		
<i>Canna indica</i> L.- flowers	Indian shot, African arrowroot, flowers	Aqueous-suspension	Saponins, catechins, tannins and carbohydrates	$LC_{50} = 7.6001 \text{ mg/ml}$ $LC_{90} = 13.891 \text{ mg/ml}$	—	Rawi et al. (2011)
		Cold-aqueous	Saponins, catechins, tannins and carbohydrates	$LC_{50} > 10,000 \mu\text{g/ml}$ after 24 h	—	Rawi et al. (2011)
		Boiled-aqueous	Saponins, catechins, tannins and carbohydrates	$LC_{50} = 4.172 \text{ mg/ml}$ $LC_{90} = 5.160 \mu\text{g/ml}$	—	Rawi et al. (2011)
				after 24 h		
<i>C. indica</i> L.-leaves	Indian shot, African arrowroot	Aqueous-suspension	Saponins, catechins, tannins and carbohydrates	$LC_{50} = 7.220 \text{ mg/ml}$ $LC_{90} = 14.105 \text{ g/ml}$	—	Rawi et al. (2011)
		Cold-aqueous	Saponins, catechins, tannins and carbohydrates	$LC_{50} > 10 \text{ mg/ml}$ after 24 h	—	Rawi et al. (2011)
		Boiled-aqueous	Saponins, catechins, tannins and carbohydrates	$LC_{50} = 7.435 \text{ mg/ml}$ $LC_{90} = 9.072 \text{ mg/ml}$	—	Rawi et al. (2011)
				after 24 h		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Cannellaceae</i>						
<i>Warburgia salutaris</i> (G.Bertol.) Chiiov.-leaves	Pepper-bark tree	Aqueous	ND	LC ₅₀ = 2.486 mg/ml LC ₉₀ = 17.370 mg/ml after 24 h	—	Clark and Appleton (1997)
<i>Caricaceae</i>				(<i>Bu. qfricanus</i>)		
<i>Carica papaya</i> L. - seeds and latex	Pappali or papaw or pawpaw	Ethanol	Proteins	LC ₅₀ = 53.4 mg/ml after 24 h LC ₅₀ = 43 µg/ml after 48 h LC ₅₀ = 23 µg/ml after 72 h LC ₅₀ = 15.9 µg/ml after 96 h	++ ++ ++ ++	Jaiswal and Singh (2008)
				(<i>L. acuminata</i>)		
		Seed powder	Proteins	LC ₅₀ = 127.3 µg/ml after 24 h LC ₅₀ = 102.9 µg/ml after 48 h LC ₅₀ = 80.9 µg/ml after 72 h LC ₅₀ = 61.6 µg/ml after 96 h	+ ++ ++ ++	Jaiswal and Singh (2008)
		Chloroform	Proteins	(<i>L. acuminata</i>)		
				LC ₅₀ = 111.9 µg/ml after 24 h LC ₅₀ = 103.9 µg/ml after 48 h LC ₅₀ = 65.7 µg/ml after 72 h LC ₅₀ = 41.2 µg/ml after 96 h	++ ++ ++ ++	Jaiswal and Singh (2008)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results (<i>L. acuminata</i>)	Activity	References
	Carbon tetrachloride	Proteins		LC ₅₀ = 81.2 µg/ml after 24 h	++	Jaiswal and Singh (2008)
				LC ₅₀ = 58.6 µg/ml after 48 h	++	
				LC ₅₀ = 42.2 µg/ml after 72 h	++	
				LC ₅₀ = 30.8 µg/ml after 96 h		
	Ether	Proteins		LC ₅₀ = 111.9 µg/ml after 24 h	+	Jaiswal and Singh (2008)
				LC ₅₀ = 93.5 µg/ml after 48 h	++	
				LC ₅₀ = 67.6 µg/ml after 72 h	++	
				LC ₅₀ = 13.1 µg/ml after 96 h		
	Acetone	Proteins		LC ₅₀ = 97.8 µg/ml after 24 h	++	Jaiswal and Singh (2008)
				LC ₅₀ = 66.3 µg/ml after 48 h	++	
				LC ₅₀ = 41.8 µg/ml after 72 h	++	
				LC ₅₀ = 31.7 µg/ml after 96 h		
	Column purified	Proteins		LC ₅₀ = 27.3 µg/ml after 24 h	++	Jaiswal and Singh (2008)
				LC ₅₀ = 20.1 µg/ml after 48 h	++	
				LC ₅₀ = 12.5 µg/ml		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results after 72 h	Activity	References
Lyophilized latex powder	Proteins			LC ₅₀ = 7.1 µg/ml after 96 h	++	Jaiswal and Singh (2008)
	(<i>L. acuminata</i>)			LC ₅₀ = 19.9 µg/ml after 24 h	++	
				LC ₅₀ = 16.5 µg/ml after 48 h	++	
				LC ₅₀ = 11.9 µg/ml after 72 h	++	
				LC ₅₀ = 8.4 µg/ml after 96 h		
	(<i>L. acuminata</i>)					
<i>C. papaya</i> L.-Leaves	Pappali or papaw or pawpaw Leaves	Ethanol	Polysaccharides, minerals, vitamins, proteins, enzymes, alkaloids, glycosides, fats, oils, lectins, saponins, flavonoids and sterols	LC ₅₀ = 2716.3 µg/ml LC ₉₀ = 4.516 mg/ml (<i>B. pfeifferi</i> -adult) LC ₅₀ = 619.1 µg/ml LC ₉₀ = 1.181 mg/ml (<i>Bu. globosus</i> -adult)	— — — —	Adetunji and Salawu (2010)
<i>Olea europaea</i> L - waste	Mission olive mill waste	<i>n</i> -hexane	Biophenols	LC ₅₀ = 424 µg/ml (<i>Isidorella newcombi</i>)	+	Obied et al. (2007)
<i>Fraxno olive mill waste</i>		<i>n</i> -hexane	Biophenols	LC ₅₀ = 541 µg/ml (<i>I. newcombi</i>)	+	Obied et al. (2007)
<i>Caryocaraceae</i>						
<i>Caryocar brasiliense</i> A.St.-Hil.-leaves	Pequi, souari nut	Ethanol	Tannins, flavanoids and terpenoids	LC ₅₀ = 20 µg/ml LC ₁₀₀ = 90 µg/ml after 48 h	++ ++	Bezerra et al. (2002)
<i>C. brasiliense</i> A.St.-Hil.-barks	Pequi, souari nut	Ethanol	Tannins, hydrolysable tannins and flavanoids	LC ₅₀ = 10 µg/ml LC ₁₀₀ = 90 µg/ml after 48 h	++ ++	Bezerra et al. (2002)
	(<i>B. glabrata</i>)					

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Capparidaceae</i>						
<i>Capparis spinosa</i> L.-	Flinders rose or caper bush aerial parts	Petrol	Coumarins, alkaloids, sterols and essential oils	LC ₅₀ = 70.6 µg/ml LC ₉₀ = 122 µg/ml after 24 h	++ + Hmamouchi et al. (2000)	
			(<i>Bu. truncatus</i>)	LC ₅₀ = 62.9 µg/ml LC ₉₀ = 135.5 µg/ml after 24 h	++ + Hmamouchi et al. (2000)	
				LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h	++ + Hmamouchi et al. (2000)	
				(<i>Bu. truncatus</i>)	LC ₅₀ = 79.2 µg/ml LC ₉₀ = 135.5 µg/ml after 24 h	++ + Hmamouchi et al. (2000)
					LC ₅₀ = 170.6 µg/ml LC ₉₀ = 194 µg/ml after 24 h	++ + Hmamouchi et al. (2000)
					(<i>Bu. truncatus</i>)	
					LC ₅₀ = 2.236 mg/ml LC ₉₀ = 2.846 mg/ml after 24 h	— — Hmamouchi et al. (2000)
						(<i>Bu. Truncates</i>)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Chenopodiaceae</i>						
<i>Atriplex stylosa</i> Viv.-whole overground parts	Atriplex	Methanol		LC ₀ = 3.1 µg/ml LC ₁₀ = 14 µg/ml LC ₂₅ = 21 µg/ml LC ₅₀ = 31 µg/ml LC ₉₀ = 54 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
		Cold-aqueous	ND	(<i>B. alexandrina</i>) LC ₀ = 5.3 µg/ml LC ₁₀ = 32 µg/ml LC ₂₅ = 39 µg/ml LC ₅₀ = 53 µg/ml LC ₉₀ = 88 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
		Cold-aqueous suspension	ND	(<i>B. alexandrina</i>) LC ₀ = 9.2 µg/ml LC ₁₀ = 38 µg/ml LC ₂₅ = 46 µg/ml LC ₅₀ = 94 µg/ml LC ₉₀ = 180 µg/ml after 24 h	+++ ++ ++ ++ +	Bakry (2009)
		Boiled-aqueous	ND	(<i>B. alexandrina</i>) LC ₀ = 4.2 µg/ml LC ₁₀ = 22 µg/ml LC ₂₅ = 30 µg/ml LC ₅₀ = 42 µg/ml LC ₉₀ = 56 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
	Ethanol			(<i>B. alexandrina</i>) LC ₀ = 9.8 µg/ml LC ₁₀ = 42 µg/ml LC ₂₅ = 49 µg/ml LC ₅₀ = 62 µg/ml	++ ++ ++ ++	Bakry (2009)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
				LC ₉₀ = 98 µg/ml after 24 h	++	
				(<i>B. alexandrina</i>)		
				LC ₀ = 6.6 µg/ml	++	Bakry (2009)
				LC ₁₀ = 42 µg/ml	++	
				LC ₂₅ = 53 µg/ml	++	
				LC ₅₀ = 66 µg/ml	++	
				LC ₉₀ = 110 µg/ml	+	
				after 24 h		
				(<i>B. alexandrina</i>)		
				LC ₀ = 7.4 µg/ml	++	Bakry (2009)
				LC ₁₀ = 54 µg/ml	++	
				LC ₂₅ = 62 µg/ml	++	
				LC ₅₀ = 74 µg/ml	++	
				LC ₉₀ = 130 µg/ml	+	
				after 24 h		
				(<i>B. alexandrina</i>)		
				LC ₀ = 1.99 µg/ml	+++	
				LC ₅₀ = 13.51 µg/ml	++	dos Santos et al. (2012)
				LC ₉₀ = 91.57 pm	++	
				after 96 h		
				(<i>B. glabra</i>)		
				LC ₅₀ = 7.7 µg/ml	++	
				LC ₉₀ = 14.1 µg/ml	++	
				after 24 h		
				(<i>Bu. truncatus</i>)		
				LC ₅₀ = 1.4 µg/ml	+++	
				LC ₉₀ = 2.2 µg/ml	+++	Hmamouchi et al. (2000)
				after 24 h		
				(<i>Bu. truncatus</i>)		
				LC ₅₀ = 6.5 µg/ml	++	
				LC ₉₀ = 9.2 µg/ml	++	
				after 24 h		
<i>Chenopodium ambrosioides</i> L.-leaves	Mexican tea	Ethanol	Caryophyllene oxide			
<i>C. ambrosioides</i> L.-aerial parts	Mexican tea	Petrol	Flavonoids, saponins, terpenoids and essential oils			
Hexane			Flavonoids, saponins, terpenoids and essential oils			
Dichloromethane			Flavonoids, saponins, terpenoids and essential oils			

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results (<i>Bu. truncatus</i>)	Activity	References
<i>C. murale</i> L.-whole overground parts	Ethyl acetate	Flavonoids, saponins, terpenoids and essential oils	LC ₅₀ = 14.1 µg/ml LC ₉₀ = 28.0 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)	
	Methanol	Flavonoids, saponins, terpenoids and essential oils	LC ₅₀ = 6.9 µg/ml LC ₉₀ = 16.8 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)	
<i>C. ugandae</i> (Aellen) Aellen-whole parts	Cold-aqueous suspension	ND	(<i>Bu. truncatus</i>)	LC ₂₅ = 1100 µg/ml LC ₅₀ = 2450 µg/ml LC ₉₀ = 5200 µg/ml after 24 h	— — —	Bakry (2009)
	Ethanol	Terpenoids, lipoids, alkaloids, glucides-glycosides, flavonoids, albuminoids, saponins and essential oils	(<i>B. alexandrina</i>)	LC ₀ = 1 mg/ml LC ₅₀ = 5 mg/ml LC ₁₀₀ = 10 mg/ml after 24 h	++ — —	Chifundera et al. (1993)
	Aqueous	Terpenoids, lipoids, alkaloids, glucides-glycosides, flavonoids, albuminoids, saponins and essential oils	(<i>B. pfeifferi</i>)	LC ₀ = 1 mg/ml LC ₅₀ = 5 m/ml LC ₁₀₀ = 10 m/ml (<i>L. natalensis</i>)	— — — — after 24 h	Chifundera et al. (1993)
				LC ₀ = 1 mg/ml LC ₅₀ = 5 mg/ml LC ₁₀₀ = 10 mg/ml (<i>B. pfeifferi</i>)	— — — —	
				LC ₀ = 1 mg/ml LC ₅₀ = 5 × 10 ³ mg/ml LC ₁₀₀ = 10 × 10 ³ mg/ml (<i>L. natalensis</i>)	— — — after 24 h	

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Chrysobalanaceae</i>						
<i>Licania cariae</i> Cardozo-leaves	–	Methanol	Catechins, flavonoid glycosides and sugars	LC ₅₀ = 25 µg/ml LC ₁₀₀ = 400 µg/ml after 24 h	++ +	Bilia et al. (2000a)
<i>L. densiflora</i> Kleinii.-leaves	–	Methanol	Catechins, flavonoids, flavonoid glycosides, sugars and tannins	(<i>B. glabra</i>) LC ₀ = 200 µg/ml LC ₁₀₀ = 400 µg/ml after 24 h	+ +	Bilia et al. (2000a)
<i>L. heteromorpha</i> Benth.-leaves	–	Methanol	Catechins, flavonoids, flavonoid glycosides, sugars and tannins	(<i>B. glabra</i>) LC ₀ = 100 µg/ml LC ₁₀₀ = 400 µg/ml after 24 h	– +	Bilia et al. (2000a)
<i>L. licanijfflora</i> (Sagot) S.F.Blake-leaves	–	Methanol	Flavonoids, flavonoid glycosides Triterpenes and triterpene glycosides	(<i>B. glabra</i>) LC ₀ = 400 µg/ml after 24 h	–	Bilia et al. (2000a)
<i>L. pittieri</i> Prance- leaves	–	Methanol	Catechins, flavonoids and flavonoid glycosides	(<i>B. glabra</i>) LC ₅₀ = 25 µg/ml LC ₁₀₀ = 400 µg/ml after 24 h	++ +	Bilia et al. (2000a)
<i>L. pyrifolia</i> Griseb.-leaves	Mercurie	Methanol	Catechins, flavonoids, flavonoid glycosides and carbohydrates	(<i>B. glabra</i>) LC ₅₀ = 25 µg/ml LC ₁₀₀ = 400 µg/ml after 24 h	++ +	Bilia et al. (2000a)
		Chloroform/ Methanol	Triterpenes, triterpene glycosides, catechins, flavonoids and flavonoid glycosides	LC ₀ = 50 µg/ml LC ₅₀ = 100 µg/ml LC ₁₀₀ = 400 µg/ml after 24 h	++ + +	Bilia et al. (2000a)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Combretaceae</i>						
<i>Commiphora molmol</i> (Engl.) Engl. ex Tschirch	Myrrh	Emulsified oil	Terpenes	LC ₅₀ = 155 µg/ml LC ₉₀ = 195 µg/ml <i>(B. alexandrina)</i>	+	Allam et al. (2001)
				LC ₅₀ = 50 µg/ml LC ₉₀ = 95 µg/ml <i>(B. truncata)</i>	++	
				LC ₅₀ = 50 µg/ml LC ₉₀ = 85 µg/ml <i>(L. caillioudii)</i>	++	
				after 24 h LC ₅₀ = 72.4 µg/ml LC ₉₀ = 125 µg/ml <i>(L. caillioudii)</i>	++	Hassan et al. (2010)
<i>Meryta denhamii</i> Seem.-flowers	Pacific puka	Methanol	ND	LC ₅₀ = 85 µg/ml LC ₉₀ = 155 µg/ml <i>(B. alexandrina)</i>	+	
				Butanol	ND	
				LC ₅₀ = 26.4 µg/ml LC ₉₀ = 70.8 µg/ml <i>(L. caillioudii)</i>	++	Hassan et al. (2010)
				LC ₅₀ = 39.8 µg/ml LC ₉₀ = 79.4 µg/ml <i>(B. alexandrina)</i>	++	
				Chloroform	ND	
				LC ₅₀ = 306 µg/ml LC ₉₀ = 125.8 µg/ml <i>(L. caillioudii)</i>	+	Hassan et al. (2010)
				Petroleum ether	ND	
				LC ₅₀ = 236.7 µg/ml LC ₉₀ = 931.8 µg/ml <i>(L. caillioudii)</i>	+	Hassan et al. (2010)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>M. denhamii</i> Seem.-fruits	Pacific puka	Methanol	ND	LC ₅₀ = 100 µg/ml LC ₉₀ = 165 µg/ml (<i>L. cailliaudi</i>)	+	Hassan et al. (2010)
				LC ₅₀ = 85 µg/ml LC ₉₀ = 138 µg/ml	++	
				(<i>B. alexandrina</i>)		
				LC ₅₀ = 864.1 µg/ml LC ₉₀ = 1222.8 µg/ml (<i>B. pfeifferi</i> -adult)	+	Adetunji and Salawu (2010)
				LC ₅₀ = 1095.7 µg/ml LC ₉₀ = 1874.9 µg/ml (<i>Bu. globosus</i> -adult) after 24 h	—	
<i>Cucurbitaceae</i>						
<i>Citrullus colocynthis</i> (L.) Schrad.- fruits and seeds	Handal	Petrol	Sterols and terpenoids	LC ₅₀ = 6.5 µg/ml LC ₉₀ = 9.2 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		
				LC ₅₀ = 4.4 µg/ml LC ₉₀ = 8.4 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		
Dichloromethane		Sterols and terpenoids		LC ₅₀ = 13.0 µg/ml LC ₉₀ = 18.3 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		
Ethyl acetate		Sterols and terpenoids		LC ₅₀ = 135.5 µg/ml LC ₉₀ = 186 µg/ml after 24 h	+	Hmamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		
Methanol		Sterols and terpenoids		LC ₅₀ = 70.6 µg/ml LC ₉₀ = 122 µg/ml after 24 h	++ +	Hmamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Dioscoreaceae</i>						
<i>Dioscorea zingiberensis</i> C.H.Wright- rhizomes	Yams	<i>n</i> -Butanol	Saponins and steroid saponins (<i>O. hupensis</i>)	LC ₉₈ = 5 µg/ml after 72 h	++	Cui et al. (1999)
<i>Dipsacaceae</i>						
<i>Cephaelaria ambrosioides</i> (Sm.) Roem. & Schult.-	Mexican tea	Acetone/ Methanol	Terpenoids and saponins (<i>B. glabrata</i>)	LC ₁₀₀ > 50 µg/ml after 24 h	+++	Pasi et al. (2009)
<i>Euphorbiaceae</i>						
<i>Acalypha indica</i> L.-	Indian Nettle	Ethanol	Glycosides, steroids and terpenoids	LC ₅₀ = 30 µg/ml after 24 h	++	Sharma et al. (2009)
Leaves				LC ₅₀ = 26 µg/ml after 48 h	++	
roots						
<i>Euphorbiaceae</i>						
<i>Acalypha indica</i> L.-	Indian Nettle	Distilled water	Glycosides, steroids and terpenoids	LC ₅₀ = 50 µg/ml after 24 h	++	Sharma et al. (2009)
Leaves				LC ₅₀ = 45 µg/ml after 48 h	++	
Chlorinated water						
<i>Euphorbiaceae</i>						
<i>Acalypha indica</i> L.-	Indian Nettle	Chlorinated water	Glycosides, steroids and terpenoids	LC ₅₀ = 80 µg/ml after 24 h	++	Sharma et al. (2009)
Leaves				LC ₅₀ = 70 µg/ml after 48 h	++	
Tap water						
<i>Euphorbiaceae</i>						
<i>Acalypha indica</i> L.-	Indian Nettle	Tap water	Glycosides, steroids and terpenoids	LC ₅₀ = 80 µg/ml after 24 h	++	Sharma et al. (2009)
Leaves				LC ₅₀ = 70 µg/ml after 48 h	++	
Chlorinated water						

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Euphorbia cphylla</i> Brouss. ex Wild.-aerial parts	Leafless Spurge	Ethanol	ND	LC ₅₀ = 0.66 µg/ml LC ₉₀ = 0.88 µg/ml (<i>L. cailliaudi</i>)	+++ +++ ++	Hassan et al. (2012)
				LC ₅₀ = 87.6 µg/ml LC ₉₀ = 142.5 µg/ml	+	
				(<i>B. alexandrina</i>) after 24 h		
		Cold-aqueous	ND	LC ₅₀ = 80 µg/ml LC ₉₀ = 109.6 µg/ml after 24 h	++ + +	Al-Zanbagi (2005)
<i>Euphorbia</i> <i>helioscopia</i> L.- fresh leaves	Sun spurge			(<i>Bu. wrightii</i>)		
		Hot-aqueous	ND	LC ₅₀ = 96.6 µg/ml LC ₉₀ = 117.7 µg/ml after 24 h	++ + +	Al-Zanbagi (2005)
		Methanol	ND	(<i>Bu. wrightii</i>)		
				LC ₅₀ = 11.3 µg/ml LC ₉₀ = 18.3 µg/ml after 24 h	++ ++ +	Al-Zanbagi (2005)
		Chloroform	ND	(<i>Bu. wrightii</i>)		
				LC ₅₀ = 80.5 µg/ml LC ₉₀ = 173.4 µg/ml after 24 h	++ + +	Al-Zanbagi (2005)
		Acetone	ND	(<i>Bu. wrightii</i>)		
				LC ₅₀ = 8.9 µg/ml LC ₉₀ = 17.5 µg/ml after 24 h	++ ++ +	Al-Zanbagi (2005)
		<i>n</i> -Hexane	ND	(<i>Bu. wrightii</i>)		
				LC ₅₀ = 99.0 µg/ml LC ₉₀ = 127.7 µg/ml after 24 h	++ + +	Al-Zanbagi (2005)
		Methanol	ND	(<i>Bu. wrightii</i>)		
				LC ₅₀ = 44.3 µg/ml	++ ++	Al-Zanbagi et al. (2000)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Euphorbia helioscopia</i> L.- dry leaves	Sun spurge	Chloroform	ND	LC ₅₀ = 65.5 µg/ml after 24 h		
				(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
		Acetone	ND	LC ₅₀ = 76.9 µg/ml	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 114.6 µg/ml after 24 h	+	Al-Zanbagi et al. (2000)
				(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
	fresh stems	Chloroform	ND	LC ₅₀ = 91 µg/ml	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 110 µg/ml after 24 h	+	Al-Zanbagi et al. (2000)
		Acetone	ND	(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 50.8 µg/ml	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 68.2 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
		Methanol	ND	(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 85.3 µg/ml	++	Al-Zanbagi et al. (2000)
		Chloroform	ND	LC ₅₀ = 101.4 µg/ml after 24 h	+	Al-Zanbagi et al. (2000)
				(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
		Acetone	ND	LC ₅₀ = 95 µg/ml	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 111.7 µg/ml after 24 h	+	Al-Zanbagi et al. (2000)
		Methanol	ND	(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 55.9 µg/ml	++	Al-Zanbagi et al. (2000)
		n-Hexane	ND	LC ₅₀ = 73 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
				(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
		Chloroform	ND	LC ₅₀ = 50 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 57.4 µg/ml	++	Al-Zanbagi et al. (2000)
		Acetone	ND	(<i>B. pfeifferi</i>)	++	Al-Zanbagi et al. (2000)
				LC ₅₀ = 23 µg/ml	++	Al-Zanbagi et al. (2000)
		Methanol	ND	LC ₅₀ = 31.7 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results (<i>B. pfeifferi</i>)	Activity	References
<i>Euphorbia helioscopia</i> L.- dry stems	Sun spurge	Chloroform	ND	LC ₅₀ = 75 µg/ml LC ₉₀ = 102 µg/ml after 24 h	++ +	Al-Zanbagi et al. (2000)
		Acetone	ND	LC ₅₀ = 69.3 µg/ml LC ₉₀ = 81.1 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
<i>Euphorbia hirta</i> L.- leaves	Ammampachcharisi Garden Spurge	Ethanol	Glycosides, steroids and terpenoids	LC ₅₀ = 20 µg/ml after 24 h	++ ++	Sharma et al. (2009)
		Distilled water	Glycosides, steroids and terpenoids	LC ₅₀ = 17 µg/ml after 48 h		
		Chlorinated water	Glycosides, steroids and terpenoids	LC ₅₀ = 40 µg/ml after 24 h	++ ++	Sharma et al. (2009)
		Tap water	Glycosides, steroids and terpenoids	LC ₅₀ = 60 µg/ml after 24 h	++ ++	Sharma et al. (2009)
				LC ₅₀ = 55 µg/ml after 48 h		
				(<i>L. acuminata</i>)		
				(<i>L. acuminata</i>)		
				(<i>L. acuminata</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>E. hirta</i> L.-latex	Ammampachcharisi Garden Spurge	Powder	ND	LC ₅₀ = 7.3 µg/ml after 24 h LC ₅₀ = 5.5 µg/ml after 48 h LC ₅₀ = 4.9 µg/ml after 72 h LC ₅₀ = 3.9 µg/ml after 96 h	+++ +++ +++ +++ +++	Yadav and Singh (2011)
<i>Euphorbia milii</i> Des Moul	Christplant	ND	ND	LC ₅₀ = 19 µg/ml LC ₉₀ = 38 µg/ml (<i>B. alexandrina</i>)	++ ++	Bakry and Mohamed (2011)
<i>Euphorbia schimperi</i> Scheele-fresh leaves	—	Hot-aqueous	ND	LC ₅₀ = 72.8 µg/ml LC ₉₀ = 140.6 µg/ml after 24 h	++ +	Al-Zanbagi (2005)
		Cold-aqueous	ND	(<i>Bu. wrightii</i>) LC ₅₀ = 81.8 µg/ml LC ₉₀ = 106 µg/ml after 24 h	++ +	Al-Zanbagi (2005)
		Methanol	ND	(<i>Bu. wrightii</i>) LC ₅₀ = 2.3 µg/ml LC ₉₀ = 6.9 µg/ml after 24 h	+++ ++	Al-Zanbagi (2005)
		Chloroform	ND	(<i>Bu. wrightii</i>) LC ₅₀ = 3 µg/ml LC ₉₀ = 7.7 µg/ml after 24 h	+++ ++	Al-Zanbagi (2005)
		Acetone	ND	(<i>Bu. wrightii</i>) LC ₅₀ = 10.1 µg/ml LC ₉₀ = 13.99 pm after 24 h	++ ++	Al-Zanbagi (2005)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
	<i>n</i> -Hexane	ND		LC ₅₀ = 18 µg/ml LC ₉₀ = 38.6 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2005)
Methanol		ND		LC ₅₀ = 18.6 µg/ml LC ₉₀ = 23.8 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
	(<i>Bu. wrightii</i>)					
Chloroform		ND		LC ₅₀ = 2.2 µg/ml LC ₉₀ = 5.6 µg/ml after 24 h	+++ +++	Al-Zanbagi et al. (2000)
	(<i>B. pfeifferi</i>)					
Acetone		ND		LC ₅₀ = 17.7 µg/ml LC ₉₀ = 24.4 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
	(<i>B. pfeifferi</i>)					
<i>n</i> -Hexane		ND		LC ₅₀ = 11.2 µg/ml LC ₉₀ = 20.8 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
	(<i>B. pfeifferi</i>)					
Methanol		ND		LC ₅₀ = 73.3 µg/ml LC ₉₀ = 95.9 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
	(<i>B. pfeifferi</i>)					
<i>E. schimperi</i> ana Scheele-dry leaves	—			LC ₅₀ = 34.8 µg/ml LC ₉₀ = 56.9 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
	(<i>B. pfeifferi</i>)					
Acetone		ND		LC ₅₀ = 73.6 µg/ml LC ₉₀ = 79.9 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
	(<i>B. pfeifferi</i>)					

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>E. schimperiana</i> Scheele-fresh stems	<i>n</i> -Hexane	ND		LC ₅₀ = 49.0 µg/ml	++	Al-Zanbagi et al. (2000)
				LC ₉₀ = 71.6 µg/ml after 24 h	++	
	Acetone	ND		(<i>B. pfeifferi</i>)		
				LC ₅₀ = 19.5 µg/ml LC ₉₀ = 26 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
	Methanol	ND		(<i>B. pfeifferi</i>)		
				LC ₅₀ = 53.8 µg/ml LC ₉₀ = 86 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
	Chloroform	ND		(<i>B. pfeifferi</i>)		
				LC ₅₀ = 10.5 µg/ml LC ₉₀ = 19.2 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
	<i>n</i> -Hexane	ND		(<i>B. pfeifferi</i>)		
				LC ₅₀ = 1.5 µg/ml LC ₉₀ = 28.4 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
<i>E. schimperiana</i> Scheele-dry stems	Methanol	ND		(<i>B. pfeifferi</i>)		
				LC ₅₀ = 7.6 µg/ml LC ₉₀ = 11.8 µg/ml after 24 h	+++	Al-Zanbagi et al. (2000)
	Chloroform	ND		(<i>B. pfeifferi</i>)		
				LC ₅₀ = 40.5 µg/ml LC ₉₀ = 47.5 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)
Acetone	ND			(<i>B. pfeifferi</i>)		
				LC ₅₀ = 20.9 µg/ml LC ₉₀ = 33.3 µg/ml after 24 h	++	Al-Zanbagi et al. (2000)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
	<i>n</i> -Hexane	ND		LC ₅₀ = 27 µg/ml LC ₉₀ = 38.8 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
<i>E. splendens</i> Bojer – ex Hook.-whole overground parts	Cold-aqueous suspension	ND		(<i>B. pfeifferi</i>) LC ₀ = 4 µg/ml LC ₁₀ = 22 µg/ml LC ₂₅ = 29 µg/ml LC ₅₀ = 40 µg/ml LC ₉₀ = 73 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
	Cold-aqueous	ND		(<i>B. alexandrina</i>) LC ₀ = 3.2 µg/ml LC ₁₀ = 13 µg/ml LC ₂₅ = 21 µg/ml LC ₅₀ = 32 µg/ml LC ₉₀ = 54 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
	Boiled-aqueous	ND		(<i>B. alexandrina</i>) LC ₀ = 2.1 µg/ml LC ₁₀ = 11 µg/ml LC ₂₅ = 15 µg/ml LC ₅₀ = 21 µg/ml LC ₉₀ = 28 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
	Methanol	ND		(<i>B. alexandrina</i>) LC ₀ = 1.1 µg/ml LC ₁₀ = 6.2 µg/ml LC ₂₅ = 8.4 µg/ml LC ₅₀ = 11 µg/ml LC ₉₀ = 27 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
	Ethanol	ND		LC ₀ = 3 µg/ml LC ₁₀ = 18 µg/ml LC ₂₅ = 23 µg/ml LC ₅₀ = 30 µg/ml LC ₉₀ = 48 µg/ml after 24 h	+++	Bakry (2009)
	Acetone	ND		LC ₀ = 4.4 µg/ml LC ₁₀ = 22 µg/ml LC ₂₅ = 31 µg/ml LC ₅₀ = 44 µg/ml LC ₉₀ = 62 µg/ml after 24 h	+++	Bakry (2009)
	Chloroform	ND		LC ₀ = 5.2 µg/ml LC ₁₀ = 30 µg/ml LC ₂₅ = 41 µg/ml LC ₅₀ = 52 µg/ml LC ₉₀ = 78 µg/ml after 24 h	++	Bakry (2009)
<i>Euphorbia splendens</i> Bojer ex Hook.- latex	Aqueous	—		LC ₉₀ (S.) = 1.5 µg/ml LC ₉₀ (Su.) = 0.55 µg/ml LC ₉₀ (F.) = 0.7 µg/ml LC ₉₀ (W.) = 0.9 µg/ml after 24 h	de +++ +++ +++ +++	Vasconcellos and de Amorim (2003)
<i>Jatropha curcas</i> L. Latex	Physic nut Latex	Acetonitrile	ND	LC ₅₀ = 6 µg/ml LC ₉₀ = 10 µg/ml after 24 h	+++ +++	Abdel-Hamid (2003) (<i>B. alexandrina</i>)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>J. curcas</i> L.-seeds	Physic nut	Methanol	Phorbol esters	LC ₅₀ = 5 µg/ml LC ₁₀₀ = 25 µg/ml after 24 h	+++ ++	Rug and Ruppel (2000)
		Aqueous	Phorbol esters	LC ₅₀ = 5,000 mg/ml after 24 h	–	Rug and Ruppel (2000)
			(<i>B. glabrata</i>)			
		Crude oil	Phorbol esters	LC ₅₀ = 50 µg/ml LC ₁₀₀ = 100 µg/ml after 24 h	++ +	Rug and Ruppel (2000)
			(<i>B. glabrata</i>)			
		Methanol	Phorbol esters	LC ₅₀ = 0.2 µg/ml (<i>Bu. truncatus</i>) LC ₁₀₀ = 1 µg/ml (<i>Bu. natalensis</i>) after 24 h	+++ +++ ++	Rug and Ruppel (2000)
				LC ₅₀ = 73.3 µg/ml LC ₉₀ = 118.8 µg/ml after 24 h	++ +	Al-Zanbagi et al. (2000)
			(<i>B. pfeifferi</i>)			
		Cold-aqueous	ND			
	<i>Jatropha glauca</i> Vahl-dry leaves	Methanol	ND	LC ₅₀ = 84.9 µg/ml LC ₉₀ = 160.7 µg/ml after 24 h	++ +	Al-Zanbagi et al. (2000)
			(<i>B. pfeifferi</i>)			
		Chloroform	ND	LC ₅₀ = 16.5 µg/ml LC ₉₀ = 46.8 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
			(<i>B. pfeifferi</i>)			
		Acetone	ND	LC ₅₀ = 102.6 µg/ml LC ₉₀ = 129 µg/ml after 24 h	+ +	Al-Zanbagi et al. (2000)
			(<i>B. pfeifferi</i>)			

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
		<i>n</i> -Hexane	ND	LC ₅₀ = 96 µg/ml LC ₉₀ = 118 µg/ml after 24 h	++ + ++	Al-Zanbagi et al. (2000)
<i>J. glauca</i> Vahl- fresh leaves		Acetone	ND	LC ₅₀ = 6.8 µg/ml LC ₉₀ = 12.5 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
		Methanol	ND	LC ₅₀ = 21.7 µg/ml LC ₉₀ = 29.8 µg/ml after 24 h	++ ++	Al-Zanbagi et al. (2000)
<i>J. mollissima</i> (Pohl) Baill.- stems	Pinhão-branco	Ethanol		LC ₁₀ = 20 µg/ml LC ₅₀ = 33.6 µg/ml LC ₅₀ = 56.3 µg/ml after 96 h	++ ++ ++	dos Santos et al. (2012)
<i>Phyllanthus</i> <i>nummularifolius</i> Poir.-whole plants		Ethanol		Saponins, alkaloids, carotenoids, catechic tannins, glucides-glycosides, flavonoids, albuminoids, steroids, terpenoids, lipoids, anthocyanins, starch and phenols	LC ₀ = 1 mg/ml LC ₅₀ = 10 mg/ml LC ₁₀₀ = 100 mg/ml (<i>B. pfeifferi</i>)	Chifundera et al. (1993)
		Aqueous		Saponins, alkaloids, carotenoids, catechic tannins, glucides-glycosides, flavonoids, albuminoids, steroids, terpenoids, lipoids, anthocyanins, starch and phenols	LC ₀ = 10 mg/ml LC ₅₀ = 50 × 10 ³ µg/ml LC ₁₀₀ = 100 mg/ml (<i>B. pfeifferi</i>)	Chifundera et al. (1993)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results (<i>L. natalensis</i>)	Activity	References
<i>Ricinus communis</i> L.-	Castor oil	Ethanol	Glycosides, steroids and terpenoids	LC ₅₀ = 12 µg/ml after 24 h	++	Sharma et al. (2009)
	Leaves			LC ₅₀ = 10 µg/ml after 48 h	++	
			(<i>L. acuminata</i>)			
Distilled water		Glycosides, steroids and terpenoids	LC ₅₀ = 60 µg/ml after 24 h	++	Sharma et al. (2009)	
			LC ₅₀ = 50 µg/ml after 48 h	++		
		(<i>L. acuminata</i>)				
Chlorinated water		Glycosides, steroids and terpenoids	LC ₅₀ = 80 µg/ml after 24 h	++	Sharma et al. (2009)	
			LC ₅₀ = 70 µg/ml after 48 h	++		
		(<i>L. acuminata</i>)				
Tap water		Glycosides, steroids and terpenoids	LC ₅₀ = 90 µg/ml after 24 h	++	Sharma et al. (2009)	
			LC ₅₀ = 85 µg/ml after 48 h	++		
		(<i>L. acuminata</i>)				
<i>Fabaceae (Leguminosae)</i>						
	<i>Acacia nilotica</i> (L.)	Gum arabic tree Deliile-leaves	Aqueous	ND	LC ₅₀ = 3616.1 µg/ml LC ₉₀ = 15,587.8 µg/ml after 24 h	Clark and Appleton (1997)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Bauhinia variegata</i> L.-leaves	Mountain ebony	Powder	Saponins and quercetins	LC ₅₀ = 244.7 µg/ml after 24 h LC ₅₀ = 203.5 µg/ml after 48 h LC ₅₀ = 155.9 µg/ml after 72 h LC ₅₀ = 126.7 µg/ml after 96 h	++ + + + + ++	Singh et al. (2012)
		Ethanol	Saponins and quercetins	(<i>L. acuminata</i>) LC ₅₀ = 38.4 µg/ml after 24 h LC ₅₀ = 28.7 µg/ml after 48 h LC ₅₀ = 20.9 µg/ml after 72 h LC ₅₀ = 14.4 µg/ml after 96 h	++ ++ ++ ++ ++	Singh et al. (2012)
		Ether	Saponins and quercetins	(<i>L. acuminata</i>) LC ₅₀ = 57.2 µg/ml after 24 h LC ₅₀ = 34.8 µg/ml after 48 h LC ₅₀ = 20.7 µg/ml after 72 h LC ₅₀ = 15.03 µg/ml after 96 h	++ ++ ++ ++ ++	Singh et al. (2012)
		Acetone	Saponins and quercetins	(<i>L. acuminata</i>) LC ₅₀ = 38.6 µg/ml after 24 h LC ₅₀ = 30.1 µg/ml after 48 h	++ ++ ++ ++	Singh et al. (2012)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Mimosa tenuiflora</i> Benth.:stems	Jurema preta	Ethanol	Flavonoids	LC ₁₀ = 6.6 µg/ml LC ₅₀ = 20.2 µg/ml LC ₉₀ = 62.1 µg/ml after 96 h	+++ ++ ++	dos Santos et al. (2012)
<i>Poincianella pyramidalis</i> (Tul.) L.P.Queiroz-leaves	Catingueira, pau-de-rato, catinga-de-porco,catingueira leaves	Ethanol	Flavonoids	LC ₁₀ = 0.04 µg/ml LC ₅₀ = 0.94 µg/ml LC ₉₀ = 20.03 µg/ml after 96 h	+++ +++ ++	dos Santos et al. (2012)
<i>Sesbania sesban</i> (L.) Merr.-leaves	Egyptian sesban	Methanol	ND	LC ₀ = 1.8 µg/ml LC ₁₀ = 8 µg/ml LC ₂₅ = 14 µg/ml LC ₅₀ = 18.0 µg/ml	+++ +++ ++	Hasheesh et al. (2011)
<i>L. acuminata</i>						
				LC ₅₀ = 21.5 µg/ml after 72 h	++	Singh et al. (2012)
				LC ₅₀ = 15.5 µg/ml after 96 h	++	
				LC ₅₀ = 43.3 µg/ml after 24 h	++	
				LC ₅₀ = 35.4 µg/ml after 48 h	++	
				LC ₅₀ = 22.5 µg/ml after 72 h		
				LC ₅₀ = 15.2 µg/ml after 96 h		
<i>B. glabrata</i>						
				LC ₁₀ = 0.04 µg/ml LC ₅₀ = 20.03 µg/ml LC ₉₀ = 14 µg/ml after 96 h	+++ +++ ++	

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
				LC ₉₀ = 31.0 µg/ml after 24 h	++	
<i>S. sesban</i> (L.) Merr. -leaves	Egyptian sesban	Dry powder	ND	(<i>Bu. truncatus</i>) LC ₀ = 5.1 µg/ml LC ₁₀ = 39.5 µg/ml LC ₂₅ = 44.9 µg/ml LC ₅₀ = 51.0 µg/ml LC ₉₀ = 62.4 µg/ml after 24 h	+++ ++ ++ ++ ++	Mahmoud et al. (2011)
<i>Ginkgoaceae</i>						
<i>Ginkgo biloba</i> L.- leaves	Ginkgo	Petroleum ether	Terpene triactones, flavonol glycosides, alkylphenols, phenolic acids, polysaccharides and amino acids	LC ₅₀ = 7.8 µg/ml LC ₉₀ = 32.7 µg/ml after 24 h	++ ++	Yang et al. (2008)
		Ethyl acetate	Terpene triactones, flavonol glycosides, alkylphenols, phenolic acids, polysaccharides and amino acids	(<i>O. hupehensis</i>) LC ₅₀ = 27.3 µg/ml LC ₉₀ = 54.9 µg/ml after 24 h	++ ++	Yang et al. (2008)
		Ethanol	Terpene triactones, flavonol glycosides, alkylphenols, phenolic acids, polysaccharides and amino acids	(<i>O. hupehensis</i>) LC ₅₀ = 64.1 µg/ml LC ₉₀ = 170.1 µg/ml after 24 h	++ +	Yang et al. (2008)

eggs

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Ginkgo biloba</i> L.	—	Petroleum ether	Ginkgolic acids	LD ₅₀ = 13.4 µg/ml after 24 h	++	Yang et al. (2006)
<i>Icacinaeae</i>						
<i>Apodtes dimidiata</i> E. Mey. ex Arn.-leaves	White pear leaves	Aqueous	ND	LC ₅₀ = 1252.4 µg/ml LC ₉₀ = 3407.9 µg/ml after 24 h	—	Clark and Appleton (1997)
<i>Lamiaceae</i>						
<i>Caryopteris clandestina</i> auct.-roots bark	Bluebeard, blue spirea or blue mist	Chloroform	ND	LC ₁₀₀ < 5 µg/ml after 24 h	+++	Hannedouche et al. (2002)
<i>Hyptis pectinate</i> (L.) Poit.-leaves	Comb bushmint	Ethanol	ND	LC ₁₀ = 8.5 µg/ml LC ₅₀ = 25.3 µg/ml LC ₅₀ = 75.7 µg/ml after 96 h	+++ ++	dos Santos et al. (2012)
<i>Lavandula stoechas</i> L.-aerial parts	Spanish lavender, French lavender	Petrol	Phenols, flavonoids, saponins, sterols, terpenoids and essential oils	LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h	+	Hmamouchi et al. (2000)
<i>n</i> -Hexane				LC ₅₀ = 116.4 µg/ml LC ₉₀ = 143.0 µg/ml after 24 h	+	Hmamouchi et al. (2000)
Dichloromethane				LC ₅₀ = 122.5 µg/ml LC ₉₀ = 144 µg/ml after 24 h	+	Hmamouchi et al. (2000)
Ethyl acetate				LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h	+	Hmamouchi et al. (2000)
(<i>Bu. truncatus</i>)						

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
		Methanol	Phenols, flavonoids, saponins, sterols, terpenoids and essential oils	LC ₅₀ = 116.4 µg/ml LC ₉₀ = 143 µg/ml after 24 h	+	Hmamouchi et al. (2000)
		Aqueous	Phenols, flavonoids, saponins, sterols, terpenoids and essential oils	LC ₅₀ = 1682 µg/ml LC ₉₀ = 259 µg/ml after 24 h	—	Hmamouchi et al. (2000)
		Essential oil	Terpenoids	LC ₅₀ = 50 µg/ml LC ₉₀ = 100 µg/ml after 3 h	++ + +	Salama et al. (2012)
<i>Marrubium vulgare</i> L.-aerial parts	White horehound	Petrol	Phenols, flavonoids, saponins, terpenoids and essential oils	LC ₁₀₀ = 200 µg/ml after 24 h	(<i>B. alexandrina</i> -adult)	
<i>Origanum compactum</i> Benth.-aerial parts	Oregano	<i>n</i> -Hexane	Phenols, flavonoids, saponins, terpenoids and essential oils	LC ₅₀ = 85.2 µg/ml LC ₉₀ = 135.5 µg/ml after 24 h	(<i>B. alexandrina</i> -egg)	
		Dichloromethane	Phenols, flavonoids, saponins, terpenoids and essential oils	LC ₅₀ = 17.8 µg/ml LC ₉₀ = 33.6 µg/ml after 24 h	++ ++ +	Hmamouchi et al. (2000)
		Ethyl acetate	Phenols, flavonoids, saponins, terpenoids and essential oils	LC ₅₀ = 8.2 µg/ml LC ₉₀ = 16.8 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)
				(<i>B. truncatus</i>)		
				LC ₅₀ = 1.0 µg/ml LC ₉₀ = 2.0 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
				(<i>B. truncatus</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Thymus capitatus</i> Hoff. et Link - aerial parts		Methanol	Phenols, flavonoids, saponins, terpenoids and essential oils	LC ₅₀ = 33.5 µg/ml LC ₉₀ = 67.0 µg/ml after 24 h	++ ++	Hnamouchi et al. (2000)
		Essential oil	Terpenoids	(<i>Bu. truncatus</i>) LC ₅₀ = 200 µg/ml LC ₉₀ = 400 µg/ml LC ₁₀₀ = no effect after 3 h	+	Salama et al. (2012)
			(<i>B. alexandrina</i> -adult)			
<i>Liliaceae</i>						
<i>Asparagus racemosus</i> Wild.-whole parts	Satavar, Satamuli, Satavari	Ethanol	Steroids, alkaloids, carotenoids, catechic tannins, glucosides- glycosides, albuminoids, coumarins, phenols, aldehydes, lipoids, saponins and flavonoids	LC ₀ = 1 mg/ml LC ₅₀ = 5 × 10 ³ µg/ml LC ₁₀₀ = 10 mg/ml (<i>B. pfeifferi</i>) LC ₀ = 0.1 mg/ml LC ₅₀ = 1 mg/ml LC ₁₀₀ = 10 mg/ml (<i>L. natalensis</i>) after 24 h	— — — — — — — —	Chitfundera et al. (1993)
		Aqueous				
<i>Liriop graminifolia</i> (L.) Baker-flowers			Saponins and steroid saponins	n-Butanol LC ₉₈ = 5 µg/ml after 72 h	+++	Cui et al. (1999)
<i>Meliaceae</i>						

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Melia azedarach</i> L.- fruits	Mallayvembu	Hexane	Flavans, flavonoids and alkaloids	LC ₅₀ = 4.44 µg/ml LC ₉₀ = 14.14 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
		Petrol	Flavans, flavonoids, terpenoids and alkaloids	LC ₅₀ = 5.93 µg/ml LC ₉₀ = 14.14 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)
		Dichloromethane	Flavans, flavonoids, terpenoids and alkaloids	(<i>Bu. truncatus</i>) LC ₅₀ = 11.11 µg/ml LC ₉₀ = 17.77 µg/ml after 24 h	+++ ++	Hmamouchi et al. (2000)
		Ethyl acetate	Flavans, flavonoids, terpenoids and alkaloids	(<i>Bu. truncatus</i>) LC ₅₀ = 122.4 µg/ml LC ₉₀ = 144 µg/ml after 24 h	+	Hmamouchi et al. (2000)
		Methanol	Flavans, flavonoids, terpenoids and alkaloids	(<i>Bu. truncatus</i>) LC ₅₀ = 164.6 µg/ml LC ₉₀ = 193 µg/ml after 24 h	+	Hmamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		
				Tannins, coumarins and flavonoids	LC ₁₀₀ = 10 µg/ml after 48 h	++
					(<i>B. glabrata</i>)	Bezerra et al. (2002)
					LC ₂₀ = 40 µg/ml LC ₅₀ = 70 µg/ml LC ₁₀₀ = 100 µg/ml after 48 h	++ ++ +
					(<i>B. glabrata</i>)	Bezerra et al. (2002)
Mimosoideae						
<i>Dimorphandra mollis</i> Benth.- barks	Fava-d'anta	Ethanol				
<i>Stryphnodendron polphyllum</i> Mart.-barks	Barbatimão	Ethanol	Tannins			

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>S. polypodium</i> Mart.-leaves	Barbatimão	Ethanol	Tannins and flavonoids	LC ₂₀ = 10 µg/ml LC ₅₀ = 70 µg/ml LC ₁₀₀ = 100 µg/ml after 48 h	+++ ++ ++	Bezerra et al. (2002)
<i>S. adstringens</i> (Mart.) Coville-leaves	Barbatimão	Ethanol	Tannins and flavonoids	LC ₂₀ = 10 µg/ml LC ₅₀ = 90 µg/ml LC ₁₀₀ = 100 µg/ml after 48 h	+++ ++ +	Bezerra et al. (2002)
<i>S. adstringens</i> (Mart.) Coville-barks	Barbatimão	Ethanol	Tannins	LC ₅₀ = 60 µg/ml LC ₁₀₀ = 100 µg/ml after 48 h	++ ++	Bezerra et al. (2002)
Myristicaceae						
<i>Moesa lanceolata</i> Forssk.-whole parts	False assegai	Ethanol	Saponins, alkaloids, carotenoids, catechic tannins, glucosides, flavonoids, albuminoids, phenols, terpenoids, lipoids and anthocyanins	LC ₀ = 10 µg/ml LC ₅₀ = 100 mg/ml LC ₁₀₀ = 10 mg/ml (<i>B. pfeifferi</i>)	++ ++ + +	Chifundera et al. (1993)
<i>L. natalensis</i>						
Aqueous				LC ₀ = 100 µg/ml LC ₅₀ = 0.500 mg/ml LC ₁₀₀ = 1 mg/ml (<i>L. natalensis</i>)	+ + +	Chifundera et al. (1993)
				after 24 h		
				LC ₀ = 0.5 mg/ml LC ₅₀ = 1 mg/ml LC ₁₀₀ = 10 mg/ml (<i>B. pfeifferi</i>)	+ + +	
				LC ₀₀ = 0.5 mg/ml LC ₅₀ = 1 mg/ml LC ₁₀₀ = 10 mg/ml (<i>L. natalensis</i>)	+ + +	
				after 24 h		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
Myrtaceae						
<i>Eugenia dysenterica</i> DC.-leaves	Cagaiteira	Ethanol	Tannins and flavonoids	LC ₅₀ = 10 µg/ml LC ₁₀₀ = 100 µg/ml after 48 h	++ + (<i>B. glabra</i> -adult)	Bezerra et al. (2002)
<i>Piperaceae</i>						
<i>Piper aduncum</i> L.-stems	Pimenta-de-macaco		Dichloromethane: Methanol (2:1)	ND		
++	Rapado et al. (2011)	Jaborandi	Dichloromethane: Methanol (2:1)	ND		
<i>Piper crassimervium</i>						
Kunth-Stems				Rapado et al. (2011)		
LC ₉₀ = 38.2 µg/ml after 24 h						
(<i>B. glabra</i> -adult)						
<i>Piper cuyabatum</i>			Pimenta do mato			
C. DC.-stems				Rapado et al. (2011)		
LC ₉₀ = 12.9 µg/ml after 24 h						
(<i>B. glabra</i> -adult)						
<i>Piper diolympifolium</i>				Dichloromethane: Methanol (2:1)	ND	
Kunth-stems						
LC ₉₀ = 23.3 µg/ml after 24 h				Rapado et al. (2011)		
(<i>B. glabra</i> -adult)						
<i>Piper hostmannianum</i>				Dichloromethane: Methanol (2:1)	ND	
(Miq.) C.DC.-stems						
LC ₉₀ = 34.3 µg/ml						

Table 1 continued

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results after 24 h	Activity	References
Dichloromethane	Coumarins, alkaloids, saponins, sterols, terpenoids and essential oils	(<i>Bu. truncatus</i>)				
Ethyl acetate	Coumarins, alkaloids, saponins, sterols, terpenoids and essential oils	(<i>Bu. truncatus</i>)	LC ₅₀ = 128.8 µg/ml LC ₉₀ = 173 µg/ml after 24 h	+	Hmamouchi et al. (2000)	
Methanol	Coumarins, alkaloids, saponins, sterols, terpenoids and essential oils	(<i>Bu. truncatus</i>)	LC ₅₀ = 122.5 µg/ml LC ₉₀ = 144 µg/ml after 24 h	+	Hmamouchi et al. (2000)	
Aqueous	Coumarins, alkaloids, saponins, sterols, terpenoids and essential oils	(<i>Bu. truncatus</i>)	LC ₅₀ = 128.8 µg/ml LC ₉₀ = 173 µg/ml after 24 h	+	Hmamouchi et al. (2000)	
<i>Pulsatilla chinensis</i> (Bunge) Regel	Baitouweng or white-haired old man	Alcohol	Triterpenoidal saponins			
<i>Ziziphus spinosa-christii</i> (L.) Desf.-aerial parts	Pulsatillae radix, Baitouweng	Ethanol	ND			
<i>Ziziphus vulgaris</i> Lamk. leaves	Chinesedate	Petrol	Phenols, flavans, flavonoids, alkaloids, saponins, sterols and terpenoids			
Dichloromethane		(<i>Bu. truncatus</i>)	LC ₅₀ = 11.2 µg/ml LC ₉₀ = 17.8 µg/ml after 24 h	++	Hmamouchi et al. (2000)	
		(<i>O. hupensis</i>)	LC ₅₀ = 0.48 µg/ml LC ₉₀ = 2.6 µg/ml after 24 h	+++	Chen et al. (2012)	
		(<i>L. caerulea</i>)	LC ₅₀ = 311 µg/ml LC ₉₀ = 500 µg/ml after 24 h	+	Hassan et al. (2012)	
		(<i>Bu. truncatus</i>)	LC ₅₀ = 8.2 µg/ml LC ₉₀ = 16.8 µg/ml after 24 h	++	Hmamouchi et al. (2000)	

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results (<i>Bu. truncatus</i>)	Activity	References	
	Ethyl acetate	Phenols, flavans, flavonoids, alkaloids, saponins, sterols and terpenoids	LC ₅₀ = 23.8 µg/ml LC ₉₀ = 56.3 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)		
Methanol	Phenols, flavans, flavonoids, alkaloids, saponins, sterols and terpenoids	(<i>Bu. truncatus</i>)	LC ₅₀ = 4.1 µg/ml LC ₉₀ = 8.4 µg/ml after 24 h	+++ ++	Hmamouchi et al. (2000)		
Aqueous	Phenols, flavans, flavonoids, alkaloids, saponins, sterols and terpenoids	(<i>Bu. truncatus</i>)	LC ₅₀ = 424 µg/ml LC ₉₀ = 548 µg/ml after 24 h	+ +	Hmamouchi et al. (2000)		
<i>Rosaceae</i>							
<i>Agrimonia eupatoria</i> auct. fl. amer.- aerial parts	Agrimony	Chloroform	Triterpenes, sterols and sesquiterpenes	LC ₀ = 100 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	++ +	Bilia et al. (2000b)	
<i>Methanol</i>		(<i>B. glabra</i>)	Triterpene glycosides, flavonoid glycosides and tannins	LC ₀ = 50.0 µg/ml LC ₁₀₀ = 500.0 µg/ml after 24 h	++ +	Bilia et al. (2000b)	
<i>Cotoneaster nebrodensis</i> (Guss.) K Koch- stems and leaves	-	Methanol	Flavonoid glycosides, tannins and sugars	(<i>B. glabra</i>)	LC ₀ = 50 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	++ +	Bilia et al. (2000b)
<i>Prunus africana</i> (Hook.f.) Kalkman-stem barks	African cherry, pygeum	Methanol	Catechins, sugars, A-type and B-type proanthocyanidins, lignans and aromatic derivatives	(<i>B. glabra</i>)	LC ₀ = 50 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	++ +	Bilia et al. (2000b)
<i>Prunus prostrata</i> Labl.-aerial parts	Rock cherry, Mountain cherry, Prostrate cherry	Methanol	Coumarin glycosides, flavanoid glycosides, sugars and A-type and B-type proanthocyanidins	(<i>B. glabra</i>)	LC ₀ = 50 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	++ +	Bilia et al. (2000b)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Potentilla montana</i> Brot.-ipogea parts	—	Methanol	Catechin glycosides, A-type and B-type proanthocyanidins and sugars	LC ₀ = 100 µg/ml LC ₅₀ = 250 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	+	Bilia et al. (2000b)
<i>Potentilla tormentilla</i> Stokes-aerial parts	Tormentil	Methanol	Triterpene glycosides, sugars and tannins	LC ₀ = 100 µg/ml LC ₅₀ = 250 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	+	Bilia et al. (2000b)
<i>Pyracantha coccinea</i> M.Roem.-roots	Scarlet firethorn	Methanol	Flavonoid glycosides, sugars and aromatic derivatives glycosides and lignin glycosides	LC ₁₀₀ = 500 µg/ml after 24 h	+	Bilia et al. (2000b)
<i>P. coccinea</i> M.Roem.-leaves	Scarlet firethorn	Methanol	Flavonoid, glycosides and sugars	LC ₀ = 100 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	+	Bilia et al. (2000b)
<i>Pyrus bourgaeana</i> Decne.-leaves and stems	Iberian pears	Methanol	Triterpene glycosides, flavonoid glycosides and aromatic derivatives glycosides	LC ₀ = 100 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	+	Bilia et al. (2000b)
<i>Quillaja saponaria</i> Molina-leaves	Soapbark	Chloroform/ Methanol	Coumarins, aromatic derivative glycosides and flavonoid glycosides	LC ₀ = 100 µg/ml LC ₅₀ = 250 µg/ml LC ₁₀₀ = 500 µg/ml after 24 h	+	Bilia et al. (2000b)
<i>Q. saponaria</i> Molina-bark	Soapbark	Aqueous	ND	LC ₁₀₀ = 2 mg/ml after 24 h	—	Bilia et al. (2000b)
				(<i>B. glabra</i>)		
				(<i>B. glabra</i>)		
				(<i>B. glabra</i>)		

Table 1 continued

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>C. bigardia</i> Duhamel-fruits	Bittersweet orange	Petrol	Phenols, coumarins, flavonoids, alkaloids, saponins, terpenoids and essential oils	LC ₅₀ = 106.7 µg/ml LC ₉₀ = 140 µg/ml after 24 h	+	Hmamouchi et al. (2000)
		<i>n</i> -Hexane	Phenols, coumarins, flavonoids, alkaloids, saponins, terpenoids and essential oils	LC ₅₀ = 3.5 µg/ml LC ₉₀ = 7.1 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
Dichloromethane			Phenols, coumarins, flavonoids, alkaloids, saponins, terpenoids and essential oils	(<i>Bu. truncatus</i>) LC ₅₀ = 2.1 µg/ml LC ₉₀ = 4.2 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
Ethyl acetate			Phenols, coumarins, flavonoids, alkaloids, saponins, terpenoids and essential oils	(<i>Bu. truncatus</i>) LC ₅₀ = 4.1 µg/ml LC ₉₀ = 8.4 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
Methanol			Phenols, coumarins, flavonoids, alkaloids, saponins, terpenoids and essential oils	(<i>Bu. truncatus</i>) LC ₅₀ = 128.8 µg/ml LC ₉₀ = 173 µg/ml after 24 h	++ +	Hmamouchi et al. (2000)
<i>Ruta chalepensis</i> L.-aerial parts	Fringed rue	Petrol	Phenols, coumarins, flavonoids, alkaloids, sterols and essential oils	(<i>Bu. truncatus</i>) LC ₅₀ = 2.1 µg/ml LC ₉₀ = 4.2 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)
Dichloromethane			Phenols, coumarins, flavonoids, alkaloids, sterols and essential oils	(<i>Bu. truncatus</i>) LC ₅₀ = 1.4 µg/ml LC ₉₀ = 2.2 µg/ml after 24 h	+++ +++	Hmamouchi et al. (2000)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
		Ethyl acetate	Phenols, coumarins, flavonoids, alkaloids, sterols and essential oils	LC ₅₀ = 70.6 µg/ml LC ₉₀ = 122 µg/ml after 24 h	++	Hmamouchi et al. (2000)
Methanol			Phenols, coumarins, flavonoids, alkaloids, sterols and essential oils	LC ₅₀ = 154.8 µg/ml LC ₉₀ = 190 µg/ml after 24 h	+	Hmamouchi et al. (2000)
Aqueous		Powder	Phenols, coumarins, flavonoids, alkaloids, sterols and essential oils	LC ₅₀ = 1092 µg/ml LC ₉₀ = 1556 µg/ml after 24 h	—	Hmamouchi et al. (2000)
<i>Sapotaceae</i>	<i>Manisops elengi</i> L. ^a barks		Saponins and quercetins	LC ₅₀ = 91.2 µg/ml after 24 h	++	Singh et al. (2012)
				LC ₅₀ = 69.7 µg/ml after 48 h	++	
				LC ₅₀ = 47.2 µg/ml after 72 h	++	
				LC ₅₀ = 36.4 µg/ml after 96 h		
				(<i>L. acuminata</i>)		
				LC ₅₀ = 44.6 µg/ml after 24 h	++	
Ethanol			Saponins and quercetins	LC ₅₀ = 31.4 µg/ml after 48 h	++	Singh et al. (2012)
				LC ₅₀ = 21.7 µg/ml after 72 h	++	
				LC ₅₀ = 15.1 µg/ml after 96 h		
				(<i>L. acuminata</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
		Ether	Saponins and quercetins	LC ₅₀ = 45.1 µg/ml after 24 h	++	Singh et al. (2012)
				LC ₅₀ = 36.9 µg/ml after 48 h	++	
				LC ₅₀ = 29.4 µg/ml after 72 h	++	
				LC ₅₀ = 23.5 µg/ml after 96 h		
			(<i>L. acuminata</i>)			
				LC ₅₀ = 58.3 µg/ml after 24 h	++	Singh et al. (2012)
				LC ₅₀ = 46.3 µg/ml after 48 h	++	
				LC ₅₀ = 34.1 µg/ml after 72 h	++	
				LC ₅₀ = 25.8 µg/ml after 96 h		
			(<i>L. acuminata</i>)			
				LC ₅₀ = 52.8 µg/ml after 24 h	++	Singh et al. (2012)
				LC ₅₀ = 46.5 µg/ml after 48 h	++	
				LC ₅₀ = 34.25 µg/ml after 72 h	++	
				LC ₅₀ = 26.02 µg/ml after 96 h		
			(<i>L. acuminata</i>)			
		Chloroform	Saponins and quercetin			

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Solanaceae</i>						
<i>Cestrum parqui</i> (Lam.) L'Hér.-whole overground parts	Willow-leaved jessamine, green cestrum, palqui	Cold-aqueous suspension	ND	LC ₀ = 86 µg/ml LC ₁₀ = 422 µg/ml LC ₂₅ = 620 µg/ml LC ₅₀ = 860 µg/ml LC ₉₀ = 1,600 mg/ml after 24 h	++ + + + -	Bakry (2009)
<i>Cestrum diurnum</i> L.-leaves	Day-blooming jasmine	Dry powder	ND	LC ₀ = 189.3 µg/ml LC ₁₀ = 207.3 µg/ml LC ₂₅ = 217.7 µg/ml LC ₅₀ = 229.3 µg/ml LC ₉₀ = 251.4 µg/ml after 24 h	++ + + + +	Mahmoud et al. (2011)
<i>Datura stramonium</i> L.-leaves	Thorn-apple, Jamestown	Dry powder	ND	LC ₀ = 2.5 µg/ml LC ₁₀ = 11.3 µg/ml LC ₂₅ = 18.2 µg/ml LC ₅₀ = 25.8 µg/ml LC ₉₀ = 40.4 µg/ml after 24 h	+++ ++ ++ ++ ++	Mahmoud et al. (2011)
<i>Physalis angulata</i> L.-whole plant	Angular winter cherry, balloon cherry, camapu, cutleaf groundcherry	Acetone	ND	(<i>B. alexandrina</i> -adult)		
		Ethyl acetate	ND	LC ₀ = 2.5 µg/ml LC ₁₀ = 11.3 µg/ml LC ₂₅ = 18.2 µg/ml LC ₅₀ = 25.8 µg/ml LC ₉₀ = 40.4 µg/ml after 24 h	+++ ++ ++ ++ ++	(<i>B. alexandrina</i> -adult) dos Santos et al. (2003)
				(<i>B. tenagophila</i>)		
				LC ₅₀ = 55.3 µg/ml LC ₉₀ = 139.4 µg/ml	++ +	(<i>B. tenagophila</i>) dos Santos et al. (2003)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Solanum aculeastrum</i> Dunal-berries	Soda apple, goat bitter apple or poison apple	Methanol	Steroidal glucosaponins and steroidal glucoalkalooids	LC ₁₀₀ = 50 µg/ml after 24 h (<i>B. pfeifferi</i>)	++	Wanyonyi et al. (2002, 2003)
<i>S. aculeastrum</i> Dunal-root barks	Soda apple, goat bitter apple or poison apple	DCCC fractions	Steroidal glucosaponins and steroidal glucoalkalooids	LC ₁₀₀ = 20.0 µg/ml after 24 h (<i>B. pfeifferi</i>)	++	Wanyonyi et al. (2002, 2003)
<i>Solanum elaeagnifolium</i> Cav.-berries	Silverleaf nightshade	Methanol	ND	LC ₃₀ = 13 µg/ml LC ₅₀ = 37 µg/ml LC ₁₀₀ = 53 µg/ml after 24 h (<i>B. pfeifferi</i>)	+++ ++ ++	Wanyonyi et al. (2003)
<i>S. elaeagnifolium</i> Cav.-roots	Silverleaf nightshade	Aqueous	Glycoalkalooids	LC ₅₀ = 9.5 µg/ml LC ₉₀ = 13 µg/ml after 24 h (<i>Bu. truncatus</i>)	+++ ++	Bekkouche et al. (2000)
<i>S. elaeagnifolium</i> Cav.-fruits	Silverleaf nightshade	<i>n</i> -Butanol	Alkaloids, tannins and saponins	LC ₅₀ = 79.9 µg/ml after 24 h (<i>Bu. truncatus</i>)	++	Larhsini et al. (2010)
		Glycoalkalooids extract	Alkaloids, tannins and saponins	LC ₅₀ = 12.2 µg/ml after 24 h (<i>Bu. truncatus</i>)	++	Larhsini et al. (2010)
		Aqueous	Alkaloids, tannins and saponins	LC ₅₀ = 9.5 µg/ml after 24 h (<i>Bu. truncatus</i>)	++	Larhsini et al. (2010)
				LC ₅₀ = 81.1 µg/ml after 24 h (<i>Bu. truncatus</i>)	++	Larhsini et al. (2010)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Solanum nigrum</i> L.	Black nightshade	Aqueous	ND	LC ₅₀ = 18.6 µg/ml LC ₉₀ = 37.7 µg/ml (<i>B. alexandrina</i>)		Ahmed and Ramzy (1997)
Leaves				LC ₅₀ = 14.5 µg/ml LC ₉₀ = 32.2 µg/ml (<i>Bu. truncatus</i>)		
				LC ₅₀ = 17.7 µg/ml LC ₉₀ = 34.7 µg/ml after 24 h (<i>L. natalensis</i>)		
Ethanol		ND		LC ₅₀ = 2.98 µg/ml LC ₉₀ = 5.95 µg/ml after 24 h (<i>B. alexandrina</i>)	+++ ++	El-Sherbini et al. (2009)
Methanol		ND		LC ₅₀ = 6.9 µg/ml LC ₉₀ = 16.2 µg/ml after 24 h (<i>B. alexandrina</i>)	++ ++	El-Sherbini et al. (2009)
Acetone		ND		LC ₅₀ = 6.6 µg/ml LC ₉₀ = 11.7 µg/ml after 24 h (<i>B. alexandrina</i>)	++ ++	El-Sherbini et al. (2009)
Chloroform		ND		LC ₅₀ = 70.8 µg/ml LC ₉₀ = 142.7 µg/ml after 24 h (<i>B. alexandrina</i>)	++ +	El-Sherbini et al. (2009)
Petroleum ether		ND		LC ₅₀ = 4.2 µg/ml LC ₉₀ = 8.6 µg/ml after 24 h (<i>B. alexandrina</i>)	+++ +++	El-Sherbini et al. (2009)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Solanum sinicum</i> Boiss.-leaves	—	Ethanol	ND	LC ₅₀ = 3.2 µg/ml LC ₉₀ = 6.04 µg/ml after 24 h	+++ +++	El-Sherbini et al. (2009)
Methanol		ND		(<i>B. alexandrina</i>) LC ₅₀ = 7.8 µg/ml LC ₉₀ = 17.5 µg/ml after 24 h	+++ ++	El-Sherbini et al. (2009)
Acetone		ND		(<i>B. alexandrina</i>) LC ₅₀ = 8.7 µg/ml LC ₉₀ = 11.2 µg/ml after 24 h	+++ ++	El-Sherbini et al. (2009)
Chloroform		ND		(<i>B. alexandrina</i>) LC ₅₀ = 82.5 µg/ml LC ₉₀ = 164.7 µg/ml after 24 h	++ +	El-Sherbini et al. (2009)
Petroleum ether		ND		(<i>B. alexandrina</i>) LC ₅₀ = 5.8 µg/ml LC ₉₀ = 9.9 µg/ml after 24 h	+++ +++	El-Sherbini et al. (2009)
<i>Solanum sodomaeum</i> Dunal-leaves	Dead sea apple	Methanol	Glycoalkaloids	(<i>B. alexandrina</i>) LC ₅₀ = 14.5 µg/ml LC ₉₀ = 21 µg/ml after 24 h	++ ++	Bekkouche et al. (2000)
<i>S. sodomaeum</i> Dunal-seeds	Dead sea apple	Methanol	Glycoalkaloids	(<i>Bu. truncatus</i>) LC ₅₀ = 15.5 µg/ml LC ₉₀ = 27 µg/ml after 24 h	++ ++	Bekkouche et al. (2000)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Solanum villosum</i> Mill.-leaves	Inhab al-deeb or red-fruit nightshade, hairy nightshade, Red nightshade	Ethanol ND		LC ₅₀ = 4.88 µg/ml LC ₉₀ = 8.95 µg/ml after 24 h	+++ +++	El-Sherbini et al. (2009)
Methanol		ND		LC ₅₀ = 8.4 µg/ml LC ₉₀ = 18.3 µg/ml after 24 h	+++ ++	El-Sherbini et al. (2009)
Acetone		ND		LC ₅₀ = 9.67 µg/ml LC ₉₀ = 12.9 µg/ml after 24 h	+++ ++	El-Sherbini et al. (2009)
Chloroform		ND		LC ₅₀ = 90 µg/ml LC ₉₀ = 175.7 µg/ml after 24 h	++ +	El-Sherbini et al. (2009)
Petroleum ether		ND		LC ₅₀ = 6.3 µg/ml LC ₉₀ = 11.02 µg/ml after 24 h	+++ ++	El-Sherbini et al. (2009)
<i>Solanum asperum</i> Rich.-fruits	'jussara' or 'coça-coça'	Methanol	Glycoalkaloid	LC ₁₀ = 7 µg/ml LC ₅₀ = 25.1 µg/ml LC ₉₀ = 44.1 µg/ml after 24 h	+++ ++ ++	Silva et al. (2008)
Alkaloidal fraction				LC ₁₀ = 2.9 µg/ml LC ₅₀ = 9.7 µg/ml LC ₉₀ = 17.3 µg/ml after 24 h	+++ +++ ++	Silva et al. (2008)
				(<i>B. glabra</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>S. S. jaberense</i> Agra & – M. Nee-aerial parts	Methanol	Alkaloids		LC ₅₀ = 56 µg/ml LC ₉₀ = 80.3 µg/ml after 24 h	++ ++	Silva et al. (2006)
	Ethanol	Alkaloids		(<i>B. glabrata</i>) LC ₅₀ = 22.8 µg/ml LC ₉₀ = 39.8 µg/ml after 24 h	++ ++	Silva et al. (2006)
	<i>n</i> -Butanol	Alkaloids		(<i>B. glabrata</i>) LC ₅₀ = 30.5 µg/ml LC ₉₀ = 46.3 µg/ml after 24 h	++ ++	Silva et al. (2006)
<i>Solanum stipulaceum</i> Roem. & Schult.-stems	Alkaloids extract	Alkaloids		(<i>B. glabrata</i>) LC ₅₀ = 45.2 µg/ml LC ₉₀ = 56 µg/ml after 24 h	++ ++	Silva et al. (2006)
<i>Withania somnifera</i> (L.) Dunal-leaves	Ashwagandha or winter cherry	Dichloromethane	Alkaloids, flavonoids, terpenes/sterols, tannins and saponins	LC ₅₀ = 94 µg/ml after 24 h	++	Larhsini et al. (2010)
<i>Verbena</i> ceae	<i>Lantana camara</i> L.-whole overground parts	Cold-aqueous suspension	ND	LC ₀ = 123 µg/ml LC ₁₀ = 520 µg/ml LC ₂₋₅ = 820 µg/ml LC ₅₀ = 1230 µg/ml LC ₉₀ = 2400 µg/ml after 24 h	++ ++ ++ ++ ++	Bakry (2009)

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
<i>Zygophyllaceae</i>						
<i>Guayacum officinale</i> s-whole overground parts	Roughbark lignum-vita	Methanol	ND	LC ₀ = 3.5 µg/ml LC ₁₀ = 17 µg/ml LC ₂₅ = 24 µg/ml LC ₅₀ = 35 µg/ml LC ₉₀ = 62 µg/ml after 24 h	+++ ++ ++ ++ ++	Bakry (2009)
				(<i>B. alexandrina</i>)		
		Cold-aqueous suspension	ND	LC ₀ = 15 µg/ml LC ₁₀ = 62 µg/ml LC ₂₅ = 97 µg/ml LC ₅₀ = 120 µg/ml LC ₉₀ = 210 µg/ml after 24 h	++ ++ ++ + +	Bakry (2009)
				(<i>B. alexandrina</i>)		
		Cold-aqueous	ND	LC ₀ = 11 µg/ml LC ₁₀ = 78 µg/ml LC ₂₅ = 88 µg/ml LC ₅₀ = 110 µg/ml LC ₉₀ = 230 µg/ml after 24 h	++ ++ ++ + +	Bakry (2009)
				(<i>B. alexandrina</i>)		
		Boiled-aqueous	ND	LC ₀ = 9 µg/ml LC ₁₀ = 48 µg/ml LC ₂₅ = 62 µg/ml LC ₅₀ = 90 µg/ml LC ₉₀ = 140 µg/ml after 24 h	++ ++ ++ ++ +	Bakry (2009)
				(<i>B. alexandrina</i>)		

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
		Ethanol	ND	LC ₀ = 7.8 µg/ml LC ₁₀ = 42 µg/ml LC ₂₅ = 57 µg/ml LC ₅₀ = 78 µg/ml LC ₉₀ = 120 µg/ml after 24 h	+++	Bakry (2009)
		Acetone	ND	LC ₀ = 14 µg/ml LC ₁₀ = 52 µg/ml LC ₂₅ = 61 µg/ml LC ₅₀ = 84 µg/ml LC ₉₀ = 140 µg/ml after 24 h	++	Bakry (2009)
		Chloroform	ND	LC ₀ = 10 µg/ml LC ₁₀ = 52 µg/ml LC ₂₅ = 65 µg/ml LC ₅₀ = 100 µg/ml LC ₉₀ = 180 µg/ml after 24 h	+++	Bakry (2009)
Zygophyllum gaenium Emb. Maire-aerial parts	—	Petrol	Flavonoids, terpenoids and saponins	LC ₅₀ = 8.7 µg/ml LC ₉₀ = 16.8 µg/ml after 24 h	+++ ++	Hmamouchi et al. (2000)
		<i>n</i> -Hexane	Flavonoids, terpenoids and saponins	(<i>Bu. truncatus</i>) LC ₅₀ = 13 µg/ml LC ₉₀ = 18.3 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)
Dichloromethane	Flavonoids, terpenoids and saponins	Flavonoids, terpenoids and saponins	(<i>Bu. truncatus</i>) LC ₅₀ = 28.2 µg/ml LC ₉₀ = 67 µg/ml after 24 h	++ ++	Hmamouchi et al. (2000)	

Table 1 continued

Species-part	Vernacular or English name	Extract type	Class of compounds	Results	Activity	References
Ethyl acetate		Flavonoids, terpenoids and saponins		LC ₅₀ = 122.4 µg/ml	+	Himamouchi et al. (2000)
				LC ₉₀ = 186 µg/ml after 24 h	+	
Aqueous		Flavonoids, terpenoids and saponins		LC ₅₀ = 372 µg/ml LC ₉₀ = 437 µg/ml after 24 h	+	Himamouchi et al. (2000)
				(<i>Bu. truncatus</i>)		

(++) means strong activity < 5 µg/mL; (+) means moderate activity < 100 µg/mL; (-) means weak activity 100–1000 µg/mL; (–) means very weak activity > 1000 µg/mL

that migrate through the blood vessel wall or are excreted with urine or feces (Nation et al. 2020).

The lifecycles of all *Schistosoma* species are quite similar, requiring freshwater snails as intermediate hosts, and mammalian hosts infected via water-borne contact with the free-swimming larvae released from the snails, as illustrated in Fig. 1. (Selbach et al. 2016). The lifecycle involves sexual maturation of adult *schistosomes* in man and an asexual multiplicative stage in the molluscan host (Nanes Sarfati et al. 2021). Differences between the species lie mainly in the type of intermediate host snails and the infective sites in the mammalian host, although there are also differences (*inter alia*) in egg morphology, and rates of both egg production and infection of mammalian hosts (Salvador-Recatalà and Greenberg 2012).

Once eggs reach fresh water they hatch and release swimming miracidia (larvae) that infect specific aquatic snails. The genera *Biomphalaria*, *Oncomelania* and *Bullinus* host *S. mansoni*, *S. japonicum*, and *S. haematobium*, respectively (Fig. 1) (Ross et al. 2002). In the snails, the miracidia asexually multiply after a few weeks and form thousands of cercariae that leave the snail and penetrate the skin of humans (e.g. children, fishermen, farmers, and women doing daily domestic tasks in infected water) or other mammals. If the cercariae penetrate the human body, they develop into schistosomules, then mate and migrate to the perivascular or mesenteric veins (*S. mansoni* and *S. japonicum*) or urinary bladder plexus (*S. haematobium*), where they produce eggs and the cycle starts again (Fig. 1) (Gryseels et al. 2006). *Schistosoma* often live 5–10 years, and even lifespans up to 40 years have been reported (Hokke and Deelder 2001).

History of schistosomiasis

Schistosomiasis has a history in Egypt of more than 5000 years (Barakat 2013; Abou-El-Naga 2018), with reports of *S. haematobium* eggs in ancient mummies (Barakat 2013). The disease was described by the ancient Egyptians on medical papyri as “â-a-â” (Abou-El-Naga 2018) and Avicenna (Ibn-Sina) in his famous book “The Canon of Medicine” (Othman and Soliman 2015). Key scientific discoveries regarding the disease have also been made in Egypt. Theodor Bilharz, a German surgeon who became Chief of the

Table 2 Screening of isolated molluscidal bioactive compounds

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
<i>Agavaceae</i>					
3-O- β -D-glucopyranosyl-(1 \rightarrow 3)- β -D-glucopyranosyl-(1 \rightarrow 3)- β -D-xylopyranoside	<i>Furcraea sellowii</i> K.Koch-leaves	SiO ₂ CC, Sephadex LH-20 CC, Pre-TLC	MP, UV, IR, (EI & D/CI) MS, ¹ H- & ¹³ C-NMR	LC ₉₀ = 6 μ g/ml after 24 h	El-Nahas et al. (2005)
Steroidal saponin (77)				(<i>B. alexandrina</i>)	
3-O-[α -L-rhamnopyranosyl (1 \rightarrow 3)- α -L-arabinopyranosyl (1 \rightarrow 3)- β -D-glucopyranosyl]-25R,5 α -spirostan-2 α ,3 β -diol-Spirostanol glycoside (36)	<i>Yucca aloifolia</i> L.-inflorescences	SiO ₂ (CC & TLC)	FAB-MS, ¹ H- & ¹³ C-NMR	LC ₁₀₀ = 10 μ g/ml (<i>B. glaberrata</i>)	Kishor and Sati (1990)
<i>Apocynaceae</i>					
Plumericin-iridoid (60)	<i>Plumeria rubra</i> L.-heartwood	SiO ₂ (CC & TLC)	MP, UV, IR, (EI & D/CI) MS, ¹ H- & ¹³ C-NMR (1D, 2D)	LC ₁₀₀ = 6.3 μ g/ml (<i>B. glaberrata</i>)	Hamburger et al. (1991)
Isoplumericin-iridoid (56)	<i>P. rubra</i> L.-heart wood	SiO ₂ (CC & TLC)	MP, UV, IR, (EI & D/CI) MS, ¹ H- & ¹³ C-NMR (1D, 2D)	LC ₁₀₀ = 6.3 μ g/ml (<i>B. glaberrata</i>)	Hamburger et al. (1991)
<i>Araliaceae</i>					
3-O-[α -L-arabinopyranosyl]-heederagenin-Triterpenoid saponin (6)	<i>Polyscias dichrostachya</i> Baker-leaves	SiO ₂ CC, TLC, Sephadex LH-20	D/CIMS, ¹ H- & ¹³ C-NMR	Active at less than 10 μ g/ml after 24 h (<i>B. glaberrata</i>)	Gopalsamy et al. (1990)
3-O-[α -L-rhamnopyranosyl -(1 \rightarrow 2)- α -L-arabinopyranosyl]-heederagenin-Triterpenoid saponin (8)	<i>P. dichrostachya</i> Baker-leaves	SiO ₂ CC, TLC, Sephadex LH-20	D/CIMS, ¹ H- & ¹³ C-NMR	Active at less than 10 μ g/ml after 24 h (<i>B. glaberrata</i>)	Gopalsamy et al. (1990)
3-O-[β -D-glucopyranosyl-(1 \rightarrow 2)- α -L-hederagenin]-heederagenin-triterpenoid saponin (9)	<i>P. dichrostachya</i> Baker-leaves	SiO ₂ CC, TLC, Sephadex LH-20	D/CIMS, ¹ H- & ¹³ C-NMR	Active at less than 10 μ g/ml after 24 h (<i>B. glaberrata</i>)	Gopalsamy et al. (1990)
<i>Asteraceae</i>					
7 α -hydroxy-3-desoxyzaluzanin C-sesquiterpene lactone (5)	<i>Podachaenium eminens</i> (Lag.) Sch.Bip. ex Sch.Bip	SiO ₂ CC	¹ H- & ¹³ C- NMR and X-ray diffraction	LC ₁₀₀ = 1 μ g/ml after 24 h (<i>B. glaberrata</i>)	Froneczek et al. (1984)

Table 2 continued

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
2-n-pentylquinoline-quinoline alkaloid (40)	<i>Galipea bracteate</i> (Nees & Mart.) Schult.-stems	SiO ₂ CC	IR, EI-MS ¹ H- & ¹³ C-NMR	Active at 20 µg/ml (<i>B. glabrata</i>)	Vieira and Kubo (1990)
2-(1'-pentenyl)-quinoline-quinoline alkaloid (39)	<i>G. bracteate</i> (Nees & Mart.) Schult.-stems	SiO ₂ CC	IR, EI-MS ¹ H- & ¹³ C-NMR	Active at 10 µg/ml (<i>B. glabrata</i>)	Vieira and Kubo (1990)
2-(3', 4'-methylene- dioxyphenylethyl)-quinoline alkaloid (37)	<i>G. bracteate</i> (Nees & Mart.) Schult.-stems	SiO ₂ CC	IR, EI-MS ¹ H- & ¹³ C-NMR	Active at 5 µg/ml (<i>B. glabrata</i>)	Vieira and Kubo (1990)
Rutin (78)	<i>Calendula officinalis</i> L.- flowers	CC, TLC	IR, MS and ¹ H- NMR	LC ₅₀ = 27.2 µg/ml after 72 h (<i>B. alexandrina</i>)	El-Sheikh et al. (2012)
				LC ₅₀ = 34.7 µg/ml after 72 h (<i>B. truncatus</i>)	
<i>Asidiaceae</i>					
Elaphopilosin-A	<i>Elaphoglossum piloselloides</i> (C. Presl) T. Moore- rhizomes, roots and sterile fronds	SiO ₂ CC, RPHPLC	IR, HREIMS, ¹ H- & ¹³ C- NMR	LC ₅₀ = 2.9 µg/ml LC ₉₀ = 4.5 µg/ml after 24 h (<i>B. peregrine</i> -adult)	Socolsky et al. (2009)
2-[{2,4-dihydroxy-6-methoxy-3-(1-oxobutyl)phenyl]methyl]-3,5-dihydroxy-4-methyl-4-(3-methylbut-2-enyl)-6-(1-oxobutyl)-2,5-cyclohexa-dien-1-one-bicyclic phloroglucinol (30)	<i>E. piloselloides</i> (C. Presl) T. Moore- rhizomes, roots and sterile fronds	SiO ₂ CC, RPHPLC	IR, HREIMS, ¹ H- & ¹³ C- NMR	LC ₅₀ = 0.9 µg/ml LC ₉₀ = 1.37 µg/ml after 24 h (<i>B. peregrine</i> -adult)	Socolsky et al. (2009)
Elaphopilosin-B					
2-[{2,4-dihydroxy-6-methoxy-3-(1-oxohexyl)phenyl]methyl]-3,5-dihydroxy-4-methyl-4-(3-methylbut-2-enyl)-6-(1-oxobutyl)-2,5-cyclohexa-dien-1-one-bicyclic phloroglucinol (32)	<i>E. piloselloides</i> (C. Presl) T. Moore- rhizomes, roots and sterile fronds	SiO ₂ CC, HPLC	IR, FABMS, ¹ H- & ¹³ C- NMR	LC ₅₀ = 10.6 µg/ml LC ₉₀ = 14.7 µg/ml after 24 h (<i>B. peregrine</i> -adult)	Socolsky et al. (2009)
Acetylation of elaphopilosin-A (62)					

Table 2 continued

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
Acetylation of elaphopilosin-B (54)	<i>E. piloselloides</i> (C. Presl) T. Moore- rhizomes, roots and sterile fronds	SiO ₂ CC, HPLC	IR, HRFABMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC, NOESY	LC ₅₀ = 30 µg/ml LC ₉₀ = 43.1 µg/ml after 24 h	Socolsky et al. (2009)
<i>Compositae</i>					
Ambrosin- sesquiterpene lactone (71)	<i>Ambrosia maritima</i> L.- aerial parts	CC, HPLC	UV	Molluscicidal (B. glabra)	Slacanin et al. (1988)
Damsin- sesquiterpene lactone (72)	<i>A. maritima</i> L.-aerial parts	CC, HPLC	UV	Molluscicidal (B. glabra)	Slacanin et al. (1988)
<i>Dipsacaceae</i>					
6α-hydroxyhederagenic acid- Terpenoid (52)	<i>Cephalaria ambrosioides</i> (Sm.) Roem. & Schult.- roots	SiO ₂ CC, TLC, MPLC	(HR-FAB & FAB) MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, HMBC, NOESY	LC ₁₀₀ > 50 µg/ml after 24 h	Pasi et al. (2009)
Hederagenic acid- terpenoid (51)	<i>C. ambrosioides</i> (Sm.) Roem. & Schult.- roots	SiO ₂ CC, TLC, MPLC	(HR-FAB & FAB) MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, HMBC, NOESY	LC ₁₀₀ > 50 µg/ml after 24 h	Pasi et al. (2009)
Leontoside A (akeboside Stb)- saponin (50)	<i>C. ambrosioides</i> (Sm.) Roem. & Schult.- roots	SiO ₂ CC, TLC, MPLC	(HR-FAB & FAB) MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, HMBC, NOESY	LC ₁₀₀ = 5.4 µg/ml after 24 h	Pasi et al. (2009)
Kalopanax saponin A (α-hederin)- saponin (75)	<i>C. ambrosioides</i> (Sm.) Roem. & Schult.- roots	SiO ₂ CC, TLC, MPLC	(HR-FAB & FAB) MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, HMBC, NOESY	LC ₁₀₀ = 6.2 µg/ml after 24 h	Pasi et al. (2009)
Saponin PG (sapindoside B)- saponin (47)	<i>C. ambrosioides</i> (Sm.) Roem. & Schult.- roots	SiO ₂ CC, TLC, MPLC	(HR-FAB & FAB) MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, HMBC, NOESY	LC ₁₀₀ = 12.8 ± 0.3 µg/ml after 24 h	Pasi et al. (2009)
Dipsacoside B- saponin (64)	<i>C. ambrosioides</i> (Sm.) Roem. & Schult.- roots	SiO ₂ CC, TLC, MPLC	(HR-FAB & FAB) MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, HMBC, NOESY	LC ₁₀₀ > 50 µg/ml after 24 h	Pasi et al. (2009)
<i>Euphorbiaceae</i>					

Table 2 continued

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
(2S,3S,4R,5R,6R,11S,13R, 14S,15R,17R)-5,6,11,14,17-pentacetoxy-3-benzoyloxy-15-hydroxy-9-oxo-seget-8(12)-ene diterpene (22)	<i>Euphorbia paralias</i> L.- aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₅₀ = 40 µg/ml after 24 h	Abdelgaleil et al. (2002)
(2S,3S,4R,6R,8R,11S,12S,13R, 14R,15R)-5,6,11,14,17-pentacetoxy-3-benzoyloxy-15-hydroxy-9-oxo-segetane-diterpene (23)	<i>E. paralias</i> L.- aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₅₀ = 60 µg/ml after 24 h (<i>B. alexandrina</i>)	Abdelgaleil et al. (2002)
(2S,3S,4R,5R,6R,8R,11S,12S,13R,14R,15R)-6,11,14,17-tetraacetoxyl-5-(2-acetoxyacetoxyl)-3-benzoyloxy-15-hydroxy-9-oxo-segetane diterpene (21)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₂₅ = 40 µg/ml LC ₅₀ = 100 µg/ml after 24 h (<i>B. alexandrina</i>)	Abdelgaleil et al. (2002)
(2S,3S,4R,5R,6R,8R,11S,12S,13R,14R,15R)-6,14,17-triacetoxy-5-(2-hydroxyacetoxyl)-3-benzoyloxy-15-hydroxy-9-oxo-segetane-diterpene (15)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₁₀₀ = 10 µg/ml after 24 h (<i>B. alexandrina</i>)	Abdelgaleil et al. (2002)
(2R,3R,4S,5R,7S,8R,13R,15R)-2,3,5,7,15-pentaacetoxyl-8-isobutyryloxy-9,14-dioxojatroph-6(17),11E-diene- diterpene (18)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₁₅ = 20 µg/ml LC ₂₅ = 60 µg/ml LC ₅₀ = 100 µg/ml LC ₁₀₀ = 100 µg/ml after 24 h (<i>B. alexandrina</i>)	Abdelgaleil et al. (2002)
(2R,3R,4S,5R,7S,8R,13R,15R)-2,3,5,7,15-pentaacetoxyl-8-isobutyryloxy-9,14-dioxojatroph-6(17),11E-diene-diterpene (24)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₂₅ = 40 µg/ml LC ₅₀ = 100 µg/ml after 24 h (<i>B. alexandrina</i>)	Abdelgaleil et al. (2002)
(2R,3R,4S,5R,7S,8R,13R,15R)-3,5,7,15-tetraacetoxyl-8-isobutyryloxy-9,14-dioxojatroph-6(17),11E-diene-diterpene (29)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₂₅ = 60 µg/ml LC ₅₀ = 60 µg/ml after 24 h (<i>B. alexandrina</i>)	Abdelgaleil et al. (2002)

Table 2 continued

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
(2S,3S,4R,5R,6R,8R,12S,13S,14R,15R)-1,5,8,14-tetraacetoxyl-3-benzoyloxy-15-hydroxy-9-oxo-paralane-diterpene (25)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₁₅ = 20 µg/ml LC ₂₅ = 80 µg/ml LC ₅₀ = 80 µg/ml LC ₁₀₀ = 100 µg/ml after 24 h	Abdelgaleil et al. (2002)
(2S,3S,4R,5R,6R,8R,12S,13S,14R,15R)-5,8,14-triacetoxyl-3-benzoyloxy-15-hydroxy-9-oxo-paralane-diterpene (26)	<i>E. paralias</i> L.-aerial parts	SiO ₂ CC, TLC, HPLC	¹ H- & ¹³ C-NMR, COSY, NOESY	LC ₁₀ = 30 µg/ml LC ₁₅ = 100 µg/ml LC ₂₅ = 100 µg/ml LC ₅₀ = 100 µg/ml after 24 h	(<i>B. alexandrina</i>) Abdelgaleil et al. (2002)
Velamolone (ent-15,16-epoxy-2-oxo-3,13(16),14-clerodatrien)-terpene (46)	<i>Croton campestris</i> A.St.-Hil.-roots bark	SiO ₂ CC, Prep TLC	UV, IR, CI-MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, NOESY	LC ₁₀₀ = 3 µg/ml after 24 h	(<i>B. alexandrina</i>) El Babili et al. (1998), El Babili et al. (2006)
Velamolone acetate (ent-15,16-epoxy-20-acetoxy-2-oxo-3,13(16),14-clerodatrien)-terpene (48)	<i>C. campestris</i> A.St.-Hil.-roots bark	SiO ₂ CC, Prep TLC	UV, IR, CI-MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, NOESY	LC ₁₀₀ = 6 µg/ml after 24 h	(<i>Bu. truncatus</i>) El Babili et al. (1998), El Babili et al. (2006)
Velamolone ((ent-15,16-epoxy-20-hydroxy-2-oxo-3,13(16),4-clerodatrien)-terpene (49)	<i>C. campestris</i> A.St.-Hil.-roots bark	SiO ₂ CC, Prep TLC	UV, IR, CI-MS, ¹ H- & ¹³ C-NMR, COSY, HMQC, NOESY	LC ₁₀₀ = 20 µg/ml after 24 h	(<i>Bu. truncatus</i>) El Babili et al. (1998), El Babili et al. (2006)
Milliamine L-terpene (79)	<i>E. miltii</i> Des Moul.- latex	Sephadex LH-20, LPLC, HPTLC, MPLC, semi prep HPLC	UV, IR, (D/Cl & EI) MS, ¹ H- & ¹³ C-NMR, COSY	LC ₁₀₀ = 2.5 × 10 ³ µg/ml after 24 h	(<i>B. glabrata</i>) Zani et al. (1993)
Milliamine M-terpene (28)	<i>E. miltii</i> Des Moul.- latex	Sephadex LH-20, LPLC, HPTLC, MPLC, semi prep HPLC	UV, IR, (D/Cl & EI) MS, ¹ H- & ¹³ C-NMR, COSY	LC ₁₀₀ = 10 × 10 ³ µg/ml after 24 h	(<i>B. glabrata</i>)

Table 2 continued

Chemical name-class	Species/part	Separation tools	Spectral analysis	Results	References
Miliamine D-terpene (27)	<i>E. militii</i> Des Moul.- latex	Sephadex LH-20, LPLC, HPTLC, MPLC, semi prep HPLC	UV, IR, ^1H - & ^{13}C -NMR, COSY	$\text{LC}_{100} = 100 \times 10^3 \mu\text{g}/\text{ml}$ after 24 h	Zani et al. (1993)
<i>Fabaceae</i>				(<i>B. glabra</i>)	
Quercetin (3,3,4,5,7-penta hydroxyflavone) (33)	<i>Bauhinia variegata</i> L.- Leaves	CC, TLC	ND	$\text{LC}_{50} = 12.1 \mu\text{g}/\text{ml}$ after 24 h	Singh et al. (2012)
				$\text{LC}_{50} = 9.9 \mu\text{g}/\text{ml}$ after 48 h	
				$\text{LC}_{50} = 6.8 \mu\text{g}/\text{ml}$ after 72 h	
				$\text{LC}_{50} = 5.4 \mu\text{g}/\text{ml}$ after 96 h	
				(<i>L. acuminata</i>)	
<i>Ginkgoaceae</i>					
Ginkgolic acid C13:0 (80)	<i>Ginkgo biloba</i> L.- sarcotesta granules	SiO ₂ CC, HPLC	LC-MS-ESI, ^1H - & ^{13}C -NMR,	$\text{LC}_{50} = 3.95 \mu\text{g}/\text{ml}$ $\text{LC}_{90} = 5.9 \mu\text{g}/\text{ml}$ after 24 h	Yang et al. (2008)
				$\text{LC}_{50} = 2.7 \mu\text{g}/\text{ml}$ $\text{LC}_{90} = 3.6 \mu\text{g}/\text{ml}$ after 48 h	
				(<i>O. hupehensis</i>)	
Ginkgolic acid C15:1 (81)	<i>G. biloba</i> L.-sarcotesta granules	SiO ₂ CC, HPLC	LC-MS-ESI, ^1H - & ^{13}C -NMR	$\text{LC}_{50} = 1.5 \mu\text{g}/\text{ml}$ $\text{LC}_{90} = 4.7 \mu\text{g}/\text{ml}$ after 24 h	Yang et al. (2008)
				$\text{LC}_{50} = 0.8 \mu\text{g}/\text{ml}$ $\text{LC}_{90} = 2.2 \mu\text{g}/\text{ml}$ after 48 h	
<i>G. sarcotestas</i>	CC, HPLC	LC-MS-ESI, ^1H - & ^{13}C -NMR	(<i>O. hupehensis</i>)	$\text{LD}_{50} = 13.4 \mu\text{g}/\text{ml}$ after 24 h	Yang et al. (2006)

Table 2 continued

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
Ginkgolic acid C17:1 (82)	<i>G. biloba</i> L.-sarcotesta granules	SiO ₂ CC, HPLC	LC-MS-ESI, ¹ H- & ¹³ C-NMR	LC ₅₀ = 11.4 µg/ml LC ₉₀ = 17.8 µg/ml after 24 h LC ₅₀ = 6.25 µg/ml LC ₉₀ = 10.01 µg/ml after 48 h	Yang et al. (2008)
<i>Icacinaeae</i>					
Genipin- iridoid (35)	<i>Apodytes dimidiata</i> E.Mey. ex Arn.- barks	SiO ₂ CC, CC, SiO ₂ TLC	(HR & EI) MS, ¹ H- & ¹³ C-NMR	LC ₅₀ = 25.3 µg/ml LC ₉₀ = 32.6 µg/ml	Drewes et al. (1996)
Genipin 10-acetate-iridoid (34)	<i>A. dimidiata</i> E.Mey. ex Arn.- barks	SiO ₂ CC, CC, SiO ₂ TLC	(HR & EI) MS, ¹ H- & ¹³ C-NMR	LC ₅₀ = 21.7 µg/ml LC ₉₀ = 39.4 µg/ml	Drewes et al. (1996)
<i>Lamiaceae</i>					
15-deoxyfuerstione (10)	<i>Caryopteris clandestinensis</i> auct.- roots bark	ND	ND	LD ₁₀₀ = 3- 4 µg/ml after 24 h	Hannedouche et al. (2002)
Fuerstione (13)	<i>C. clandestinensis</i> auct.- roots bark	ND	ND	LD ₁₀₀ = 3- 4 µg/ml after 24 h	Hannedouche et al. (2002)
α-caryopterone (42)	<i>C. clandestinensis</i> auct.- roots bark	ND	ND	LD ₁₀₀ = 1- 2 µg/ml after 24 h	Hannedouche et al. (2002)
<i>Leguminosae</i>					
2'-hydroxy 3,7,8,4',5'-pentamethoxy flavones (58)	<i>Parkia clappertoniiana</i> Keay- leaves	TLC, CC	UV, IR, EIMS, ¹ H- & ¹³ C-NMR, NOE	LC ₈₀ = 25 µg/ml	Lemnich et al. (1996)
Pachyelaside A- triterpene saponin (57)	<i>Pachyelasma tessmannii</i> (Harms) Harms- roots bark	CC, HPLC	IR, MALDI-TOF-MS ¹ H- & ¹³ C-NMR, DEPT, DQF-COSY, HSQC, HMBG, HOHAHA, NOE, NOESY	LC ₅₀ = 2 µg/ml after 24 h	Nihei et al. (2005)

Table 2 continued

Chemical name-class	Species/part	Separation tools	Spectral analysis	Results	References
Pachyelaside B-triterpene saponin (55)	<i>P. tessmannii</i> (Harms) Harms- roots bark	CC, HPLC	IR, MALDI-TOF-MS ^1H - & ^{13}C -NMR, DEPT, DQF-COSY, HSQC, HMBC,	$\text{LC}_{50} = 2 \mu\text{g/ml}$ after 24 h (<i>B. glaberrata</i>)	Nihei et al. (2005)
Pachyelaside C-triterpene saponin (61)	<i>P. tessmannii</i> (Harms) Harms- roots bark	CC, HPLC	HOHAHA, NOE, NOESY IR, MALDI-TOF-MS ^1H - & ^{13}C -NMR, DEPT, DQF-COSY, HSQC, HMBC,	$\text{LC}_{50} = 2 \mu\text{g/ml}$ after 24 h (<i>B. glaberrata</i>)	Nihei et al. (2005)
Pachyelaside D-triterpene saponin (63)	<i>P. tessmannii</i> (Harms) Harms-roots bark	CC, HPLC	HOHAHA, NOE, NOESY IR, MALDI-TOF-MS ^1H - & ^{13}C -NMR, DEPT, DQF-COSY, HSQC, HMBC,	$\text{LC}_{50} = 8 \mu\text{g/ml}$ after 24 h (<i>B. glaberrata</i>)	Nihei et al. (2005)
<i>Myrsinaceae</i>					
Miaesaponin V1 ₂	<i>Maesa lanceolata</i> Forssk.- leaves	Semi-prep HPLC	Comparison with reported data	$\text{LC}_{50} = 0.5 \mu\text{g/ml}$ (<i>B. glaberrata</i>)	Apers et al. (2001)
3- β -O-[(α -L-rhamnopyranosyl-(1 → 2)- β -D-galactopyranosyl-(1 → 3)]- β -D-galactopyranosyl-(1 → 2)- β -D-glucopyranuronyl]-21 β ,22 α -diangeloyloxy-13 β ,28-epoxyolean-16 α ,28 α -diol-triterpenoid saponin (76)					
<i>Papaveraceae</i>					
Sanguinarine-alkaloid (BSA) (70)	<i>Macleaya cordata</i> (Willd.) R.Br.-fruits	SiO ₂ CC, HPLC	ND	$\text{LC}_{50} = 0.59 \mu\text{g/ml}$ $\text{LC}_{90} = 1.38 \mu\text{g/ml}$ after 48 h	Ming et al. (2011)
				$\text{LC}_{50} = 0.19 \mu\text{g/ml}$ $\text{LC}_{90} = 0.54 \mu\text{g/ml}$ after 72 h	
				(<i>O. hupensis</i>)	

Table 2 continued

Chemical name-class	Species/part	Separation tools	Spectral analysis	Results	References
Chelerythrine-alkaloid (BCHE) (73)	<i>M. cordata</i> (Willd.) R.Br.-fruits	SiO ₂ CC, HPLC	ND	LC ₅₀ = 2.7 µg/ml LC ₉₀ = 5.2 µg/ml after 48 h	Ming et al. (2011)
				LC ₅₀ = 2.01 µg/ml LC ₉₀ = 4.86 µg/ml after 72 h	
		(<i>O. hapensis</i>)			
<i>Primulaceae</i>					
Anagallide B- oleanane saponin (74)	<i>Anagallis arvensis</i> L	SiO ₂ CC, prep HPLC	HRFABMS, ¹ H- & ¹³ C-NMR,	LC ₁₀₀ = 5.0 µg/ml (<i>B. glabra</i>)	Abdel-Gawad et al. (2000)
				LC ₁₀₀ = 2.5 µg/ml (<i>O. quadrasi</i>)	
Desglucanagalloloside B oleanane saponin (69)	<i>A. arvensis</i> L	SiO ₂ CC, prep HPLC	HRFABMS, ¹ H- & ¹³ C-NMR,	LC ₁₀₀ = 2.5 µg/ml (<i>B. glabra</i>)	Abdel-Gawad et al. (2000)
				LC ₁₀₀ = 1.25 µg/ml (<i>O. quadrasi</i>)	
<i>Phytolaccaceae</i>					
3-O-[3'- <i>O</i> -β-D-galactopyranosyl]-β-D-glucopyranosyl] 2β-hydroxyoleanolic acid-oleanane saponin (66)	<i>Phytolacca dodecandra</i> L'Hér.-berries	SiO ₂ CC, TLC, HPLC	LSIMS, ¹ H- & ¹³ C-NMR, COSY, TOCSY, HETCOR, NOESY	LC ₅₀ = 5- 10 µg/ml (<i>B. glabra</i>)	Thilborg et al. (1993, 1994)
Mixture (1:1) of 3-O-[<i>O</i> -β-D-galactopyranosyl-(1 → 3)- <i>O</i> -[β-D-glucopyranosyl-(1 → 4)]-β-D-glucopyranosyl] bayogenin- saponin (65) + 3-O-[2',4'-di- <i>O</i> -β-D-glucopyranosyl]- hederagenin-oleane saponin	<i>P. dodecandra</i> L'Hér.-berries	SiO ₂ CC, TLC, HPLC	FABMS, ¹ H- & ¹³ C-NMR, COSY, TOCSY, HETCOR, NOESY	LC ₅₀ = 10-20 µg/ml (<i>B. glabra</i>)	Thilborg et al. (1993, 1994)
3-O-β-D-glucopyranosylserjaniic acid-saponin (14)	<i>P. icosaandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC	LC ₁₀₀ = 3.1 µg/ml after 24 h (<i>B. glabra</i>)	Treyvaud et al. (2000)

Table 2 continued

Chemical name-class	Species/part	Separation tools	Spectral analysis	Results	References
3-O-(β -D-galactopyranosyl)-(1 → 3)- β -D-glucopyranosyl) serjanie acid-saponin (16)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC	LC ₁₀₀ = 3.1 µg/ml after 24 h	Treyvraud et al. (2000)
3-O-(β -D-glucopyranosyl)-(1 → 2)- β -D-glucopyranosyl) serjanie acid-saponin (17)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC	LC ₁₀₀ = 10 µg/ml after 24 h	Treyvraud et al. (2000)
3-O-(β -D-glucopyranosyl)-(1 → 3)- β -D-glucopyranosyl-(1 → 3)- β -D-glucopyranosyl serjanie acid-saponin (19)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC	LC ₁₀₀ = 12.5 µg/ml after 24 h	Treyvraud et al. (2000)
3-O-(α -L-rhamnopyranosyl)-(→)- β -D-glucopyranosyl-(1 → 2)- β -D-glucopyranosyl serjanie acid-saponin (20)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC	LC ₁₀₀ = 50 µg/ml after 24 h	Treyvraud et al. (2000)
3-O-(α -L-rhamnopyranosyl)-(1 → 2)- β -D-glucopyranosyl-(1 → 2)- β -D-glucopyranosyl sphaerulagenic acid-saponin (83)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC	LC ₁₀₀ > 50 µg/ml after 24 h	Treyvraud et al. (2000)
3-O-(β -D-galactopyranosyl)-(1 → 3)- β -D-glucopyranosyl serjanie acid 28-O- β -D-glucopyranoside-saponin (7)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC, chemical transformation	LC ₁₀₀ > 50 µg/ml after 24 h	Treyvraud et al. (2000)
3-O-(β -D-glucopyranosyl)-(1 → 2)- β -D-glucopyranosyl serjanie acid 28-O- β -D-glucopyranoside-saponin (12)	<i>P. icostandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC, chemical transformation	LC ₁₀₀ > 50 µg/ml after 24 h	Treyvraud et al. (2000)

Table 2 continued

Chemical name-class	Species-part	Separation tools	Spectral analysis	Results	References
3-O-(α -L-rhamnopyranosyl-(1 → 2)- β -D-glucopyranosyl-(1 → 2)- β -D-glucopyranosyl-28-O- β -D-glucopyranoside - saponin (11)	<i>P. icoxandra</i> L.-berries	SiO ₂ CC, Sephadex LH-20 SiO ₂ , LPLC on Lobar RP-18 C, CPC, HPLC	ESMS, ¹ H- & ¹³ C-NMR, COSY, HSQC, HMBC, chemical transformation	LC ₁₀₀ > 50 μ g/ml after 24 h	Treyraud et al. (2000)
<i>Rubiaceae</i>					Lemmich et al. (1995)
3-O-[O - α -L-rhamnopyranosyl-(1 → 3)-O- β -D-glucopyranosyl-(1 → 3)]- β -D-glucopyranosyl-oleanolic acid	<i>Cattunaregam nilotica</i> (Stapf) Tirveng.-fruits	SiO ₂ CC, TLC	FABMS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₅₀ = 3 μ g/ml (<i>B. alexandrina</i>)	Lemmich et al. (1995)
oleanane saponin (85)					
28-O- β -D-glucopyranosyl-3- O-[O - α -L-rhamnopyranosyl-(1 → 3)-O- β -D-glucopyranosyl-(1 → 3)]- β -D-glucopyranosyl-oleanolate -oleanane saponin (87)	<i>C. nilotica</i> (Stapf) Tirveng.-fruits	SiO ₂ CC, TLC	FABMS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₅₀ = 3 μ g/ml (<i>B. alexandrina</i>)	Lemmich et al. (1995)
3-O-[2',3'-di- O (β -D-glucopyranosyl)- β -D-glucopyranosyl]oleanolic acid	<i>C. nilotica</i> (Stapf) Tirveng.-fruits	SiO ₂ CC, TLC	FABMS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₅₀ = 26 μ g/ml (<i>B. alexandrina</i>)	Lemmich et al. (1995)
oleanane saponin (86)					
3-O-[O - β -D-glucopyranosyl-(1 → 3)- β -D-glucopyranosyl] oleanolic acid-oleanane saponin (84)	<i>C. nilotica</i> (Stapf) Tirveng.-fruits	SiO ₂ CC, TLC	FABMS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₅₀ = 3 μ g/ml (<i>B. alexandrina</i>)	Lemmich et al. (1995)
<i>Rutaceae</i>					
Cuspane	<i>Angostura paniculata</i> (Engl.) T.S.Elias-leaves	SiO ₂ CC	UV, IR, ¹ H- & ¹³ C-NMR, NOESY, X-ray	LC ₅₀ = 5 μ g/ml (<i>B. glabrata</i>)	Vieira et al. (1992)
(1-hydroxy-2,3,5,6-tetramethoxy-9-acridone) - alkaloid (59)					
Cuscline	<i>A. paniculata</i> (Engl.) T.S.Elias -leaves	SiO ₂ CC	UV, IR, ¹ H- & ¹³ C-NMR, NOESY, X-ray	LC ₅₀ = 20 μ g/ml (<i>B. glabrata</i>)	Vieira et al. (1992)
(1,2,3,5,6-pentamethoxy-9-acridone) - alkaloid (88)					
<i>Sapindaceae</i>					
Pulsatilla saponin D	<i>Serjania salzmanniana</i> Schldl. stems	CC, Prep TLC, SiO ₂ TLC	HRFAB-MS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₁₀ = 90 μ g/ml LC ₂₀ = 100 μ g/ml after 48 h	Ekabo et al. (1996)
(3-O-[[β -D -glucopyranosyl-(1 → 4)]- α -L-rhamnopyranosyl-(1 → 2)]- α -L-arabinopyranosyl] hederaegenin-saponin (31)					

Table 2 continued

Chemical name-class	Species/part	Separation tools	Spectral analysis	Results	References
3-O-[β -D-glucopyranosyl-(1 → 4)]- α -L-rhamnopyranosyl-(1 → 2)- α -L-arabinopyranosyl oleanolic acid-saponin (44)	<i>S. salzmanniana</i> Schidl.-stems	CC, Prep TLC, SiO ₂ TLC	LR-FAB-MS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₁₀ = 100 µg/ml LC ₂₀ = 100 µg/ml after 48 h	Ekabo et al. (1996)
Salzmannianoside A (3-O-[[β -D-glucopyranosyl-(1 → 4)]- α -L-rhamnopyranosyl-(1 → 2)]- α -L-arabinopyranosyl]-gypsogenin)-saponin (45)	<i>S. salzmanniana</i> Schidl.-stems	Prep TLC, SiO ₂ TLC and CC	HRFAB-MS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, NOESY	LC ₂₀ = 100 µg/ml after 48 h	Ekabo et al. (1996)
Salzmannianoside B (3-O-[[β -D-glucopyranosyl-(1 → 4)]- α -L-arabinopyranosyl-(1 → 3)- α -rhamnopyranosyl-(1 → 2)]- α -L-arabinopyranosyl]-hederagenin)-saponin (53)	<i>S. salzmanniana</i> Schidl.-stems	CC, Prep TLC, SiO ₂ TLC	(HRFAB & CI) MS, ¹ H- & ¹³ C-NMR, COSY, HETCOR, HOHAHA, NOESY	LC ₁₀ = 100 µg/ml LC ₂₀ = 70 µg/ml after 48 h	Ekabo et al. (1996)
<i>Sapotaceae</i>					
Sapogenin-saponin (43)	<i>Minuusops elengi</i> L.-barks	TLC, CC	ND		Singh et al. (2012)
<i>Solanaceae</i>					
mixture of solamarginine + solasonine-glucoalkaloid (2) + (68)	<i>Solanum mammosum</i> L.-fruits	SiO ₂ TLC	¹ H- & ¹³ C-NMR, COSY, DQF-COSY, TOCSY, HMQC, NOESY	LC ₁₀₀ = 25.0 µg/ml	Mahato et al. (1980), Alzérecca and Hart (1982), Puri et al. (1994)

Table 2 continued

Chemical name-class	Species/part	Separation tools	Spectral analysis	Results	References
Tomatine-glucoalkaloid (89)	<i>S. mammosum</i> L.-fruits	SiO ₂ CC	Comparison with reported data	LD ₁₀₀ = 25 µg/ml (<i>L. cubensis</i>)	Alzétreca and Hart (1982)
Solanandaine-alkaloid (67)	<i>S. asperum</i> Rich.-fruits	TLC; Sephadex LH-20	IR, HREIMS, LC-MS, ¹ H- & ¹³ C-NMR, HBBD, DEPT, COSY, HSQC, HMBIC TOCSY, NOESY after 24 h	LC ₁₀ = 46.4 µg/ml LC ₅₀ = 73.1 µg/ml LC ₉₀ = 99.7 µg/ml	Silva et al. (2008)
Solasonine-alkaloid (68)	<i>S. asperum</i> Rich.-fruits	TLC; Sephadex LH-20	IR, HREIMS, LC-MS, ¹ H- & ¹³ C-NMR, HBBD, DEPT, COSY, HSQC, HMBIC TOCSY, NOESY after 24 h	LC ₁₀ = 22 µg/ml LC ₅₀ = 47 µg/ml LC ₉₀ = 72 µg/ml	Silva et al. (2008)
Solamargine-Alkaloid (2)	<i>S. asperum</i> Rich.-fruits	TLC; Sephadex LH-20	IR, HREIMS, LC-MS, ¹ H- & ¹³ C-NMR, HBBD, DEPT, COSY, HSQC, HMBIC, TOCSY, NOESY after 24 h	LC ₁₀ = 8.1 µg/ml LC ₅₀ = 26.3 µg/ml LC ₉₀ = 63.6 µg/ml	Silva et al. (2008)

HPLC high performance liquid chromatography, *TLC* thin layer chromatography, *CC* column chromatography, *SiO₂* silica gel, *CPC* centrifugal partition chromatography, *LP LC* low pressure liquid chromatography, *MPLC* medium pressure liquid chromatography, *Prep TLC* preparative thin layer chromatography, *DCCC* droplet counter-current chromatography, *NOESY* nuclear over hauser effect spectroscopy, *COSY* homonuclear correlated spectroscopy, *TOCSY* total correlation spectroscopy, *HMBC* heteronuclear multiple bond connectivity, *EL-MS* electron ionization mass spectrometry, *TOF MS* time-of-flight mass spectrometry, *MALDI* matrix-assisted laser desorption/ionization, *NMR* nuclear magnetic resonance, *HETCOR* heteronuclear correlation spectroscopy, *UV* ultraviolet, *IR* infrared, *MP* melting point

Table 3 Plants species (and their localitis) used to cure schistosomiasis in patients based on traditional uses

Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Acanthaceae</i>					
<i>Peristrophe bicalyculata</i> (Retz.) Nees	Panicled Foldwing	Aerial parts	Mali	ND	Bah et al. (2006), Gaudani et al. (2010)
<i>Anacardiaceae</i>					
<i>Ozona insignis</i> Delile	—	Leaves	Zimbabwe	At 33.8 mg/ml all schistosomes died (<i>Schistosoma mansoni</i>)	Mølgaard et al. (2001)
<i>Lannea bicolor</i>	Dikbas	Roots bark	Zimbabwe	At 25.3 mg/ml all schistosomes died (<i>S. mansoni</i>)	Mølgaard et al. (2001)
<i>Sclerocarya birrea</i> (A.Rich.) Hochst.	Jelly plum	Twigs Roots	Africa Africa	ND ND	He et al. (2002) Sparg et al. (2000)
<i>Rhus geninzi</i>	—	Roots	South Africa	ND	Sparg et al. (2000)
<i>Annonaceae</i>					
<i>Annona senegalensis</i> Pers. <i>A. muricata</i> L.	Wild Custard Apple and Wild Soursop Graviola, Soursop and Gunbanana	Roots Leaves	Mali Brazil	ND See Table 1	Bah et al. (2006) dos Santos and Sant'Ana (2001)
<i>Apocynaceae</i>					
<i>Carrissa edulis</i> (Forssk.) Vahl.	Climbing num-num, simple-spined num-num	Roots	South Africa	ND	Sparg et al. (2000)
<i>Landolphia kirkii</i> Dyer	Sand apicot-vine, rubber vine	Roots	South Africa	ND	Sparg et al. (2000)
<i>Saba senegalensis</i> (A.D.C.) Pichon	Weda	Leaves	Mali	ND	Bah et al. (2006)
<i>Asclepiadaceae</i>					
<i>Calotropis procera</i> (Aiton)	Apple of Sodom	Roots	Mali	See Table 1	Bah et al. (2006)
<i>Dryand.</i>					
<i>Asparagaceae</i>					
<i>Agave attenuate</i> Salm-Dyck	Foxtail agave	Dry leaves	South Africa	ND	Brackenbury and Appleton (1997)

Table 3 continued

Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Asparagus africanus</i> Lam	<i>African asparagus</i>	Roots	South Africa	ND	Sparg et al. (2000)
<i>Asphodelaceae</i>					
<i>Bulbine abyssinica</i> A.Rich	Bushy bulbine	Roots	South Africa	ND	Sparg et al. (2000)
<i>Asteraceae (Compositae)</i>					
<i>Alomia myriadenia</i> Sch.Bip. ex Baker	—	Aerial parts	Brazil	See Table 1	Mendes et al. (1999)
<i>Achyrocline satureoides</i> (Lam.) DC	Macela or marcela	Aerial parts	Brazil	see Table 1	Mendes et al. (1999)
<i>Actinotis</i>	—	Aerial parts	Brazil	See Table 1	Mendes et al. (1999)
<i>angustifolia</i> (Gardner) Cabrala	—	Roots	South Africa	ND	Sparg et al. (2000)
<i>Berkheya speciosa</i> (DC.) O.Hoffm	—	Barks, twigs and leaves	South Africa	ND	Ojewole (2004)
<i>Piptocarpha rotundifolia</i> (Less.) Baker	Ashdaisy	Aerial parts	Brazil	See Table 1	Mendes et al. (1999)
<i>Vanilloopsis erythropappa</i> (DC.) Sch.Bip	Candeia tree	Aerial parts	Brazil	See Table 1	Mendes et al. (1999)
<i>Verbesina clausenii</i> Sch.Bip. ex Baker	—	Aerial parts	Brazil	See Table 1	Mendes et al. (1999)
<i>Balanitaceae</i>					
<i>Balanites maughamii</i> Sprague	- Desert date Or Haglig	Fruits, leaves and barks	South Africa	ND	Ojewole (2004)
<i>Boraginaceae</i>					
<i>B. aegyptica</i> (L.) Delile	Desert date Or Haglig	Fruits, leaves and barks	South Africa	ND	Ojewole (2004)
<i>Trichodesma physaloides</i> (Fenzl) A. DC	Chocolate bells	Tubers	South Africa	ND	Sparg et al. (2000)
<i>Caesalpiniaceae</i>					
<i>Afelia quanzensis</i> Welw	Pod Mahogany	Roots	South Africa	ND	Sparg et al. (2000)
<i>Camelliaceae</i>					

Table 3 continued

Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Warburgia ugandensis</i> Sprague	Green heart	Fruits, twigs and leaves	South Africa		Ojewole (2004)
<i>Warburgia salutaris</i> (G.Bertol.) Chiov	Pepper-bark tree	Fruits, twigs and leaves	South Africa		Ojewole (2004)
<i>Capparidaceae</i>					
<i>Cadaba farinosa</i> Forsk	Indian Cadaba	Leaves	Mali	ND	Bah et al. (2006)
<i>Capparis tomentosa</i> Lam	Woolly caper bush	Leaves and roots	Mali	ND	Bah et al. (2006)
<i>Celastraceae</i>					
<i>Maytenus senegalensis</i> (Lam.) Exell	Spike thorn	Roots	South Africa	ND	Sparg et al. (2000)
<i>Chrysobalanaceae</i>					
<i>Licania tomentosa</i> (Benth.) Fritsch	—	Leaves	Venezuela		Bilia et al. (2000a)
<i>Cochlospermaceae</i>					
<i>Cochlospermum tinctorium</i> Perrier ex A.Rich	—	Leaves	Mali	ND	Bah et al. (2006)
<i>Combretaceae</i>					
<i>Anogeissus leiocarpa</i> (DC.) Guill. & Perr	African birch	Leaves	Mali	ND	Bah et al. (2006)
<i>Combretum imberbe</i> Wawra	Leadwood tree	Bark, twigs and leaves	South Africa		Ojewole (2004)
<i>Combretum mollle</i> R.Br. ex G.Don	Velvet bushwillow	Bark, twigs and leaves	South Africa		Ojewole (2004)
<i>Combretum micranthum</i> G.Don	Guinean kinkeliba	Leaves	Mali	ND	Bah et al. (2006)
<i>Combretum imberbe</i>	Leadwood tree	Roots	South Africa	ND	Sparg et al. (2000)
<i>Ebenaceae</i>					
<i>Euclea divinorum</i> Hiern	Magic guarri, diamond leaf, toothbrush tree, or diamond leaved <i>Euclea</i>	Roots	South Africa	ND	Sparg et al. (2000)
<i>Euclea natalensis</i>	-	Barks	South Africa	ND	Sparg et al. (2000)

Table 3 continued

Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Euphorbiaceae</i>					
<i>Antidesma venosum</i> E.Mey. ex Tul	Tassel-berry	Roots	South Africa	ND	Sparg et al. (2000)
<i>Euphorbia splendens</i> Bojer ex Hook	—	Latex	Brazil	See Table 1	Schall et al. (2001)
<i>E. hirta</i> L.	Ammampach charisi	Whole plant	Mali	See Table 1	Bah et al. (2006)
<i>Ricinus communis</i> L.	Castor oil	Roots	South Africa	See Table 1	Sparg et al. (2000)
<i>Securinega virosa</i> (Roxb. ex Willd.) Baill	White berry bush, snowberry tree or simple-leaf bush weed	Roots	Mali	ND	Bah et al. (2006)
<i>Fabaceae (Leguminosae)</i>					
<i>Abrus precatorius</i> L.	Jequirity bean or rosary pea	Roots	South Africa	ND	Sparg et al. (2000)
		Stems	Zimbabwe	At 1.5 mg/ml all schistosomes died	Mølgaard et al. (2001)
		Roots	Zimbabwe	At 0.6 mg/ml all schistosomes died	Mølgaard et al. (2001)
			(<i>S. mansoni</i>)	(<i>S. mansoni</i>)	
<i>Acacia karroo</i> Hayne	Sweet thorn	Leaves	Zimbabwe	At 103 mg/ml no schistosomes died	Mølgaard et al. (2001)
<i>Albizia antunesiana</i> Harms	—	Roots	Zimbabwe	At 100 mg/ml no schistosomes died (<i>S. mansoni</i>)	Mølgaard et al. (2001)
<i>Cassia italica</i> (Mill.) F.W.Andrews	—	Leaves	Mali	ND	Bah et al. (2006)
<i>C. nigricans</i> Vahl	—	Whole plant	Mali	ND	Bah et al. (2006)
<i>Cassia sieberiana</i> DC	Drumstick tree	Leaves	Mali	ND	Bah et al. (2006)
<i>Dalbergia sissoo</i> DC	Indian rosewood	Fruits, leaves, roots and stems bark	Nigeria	ND	Adenusi and Odaibo (2009)
<i>Elephantorrhiza goetzei</i> (Harms) Harms	Goetze's elephantorrhiza	Stems bark	Zimbabwe	At 0.8 mg/ml all schistosomes died (<i>S. mansoni</i>)	Mølgaard et al. (2001)
<i>Lonchocarpus laxiflorus</i> Guill. & Perr	—	Roots	South Africa	ND	Sparg et al. (2000)
		Barks	Mali	ND	Bah et al. (2006)

Table 3 continued

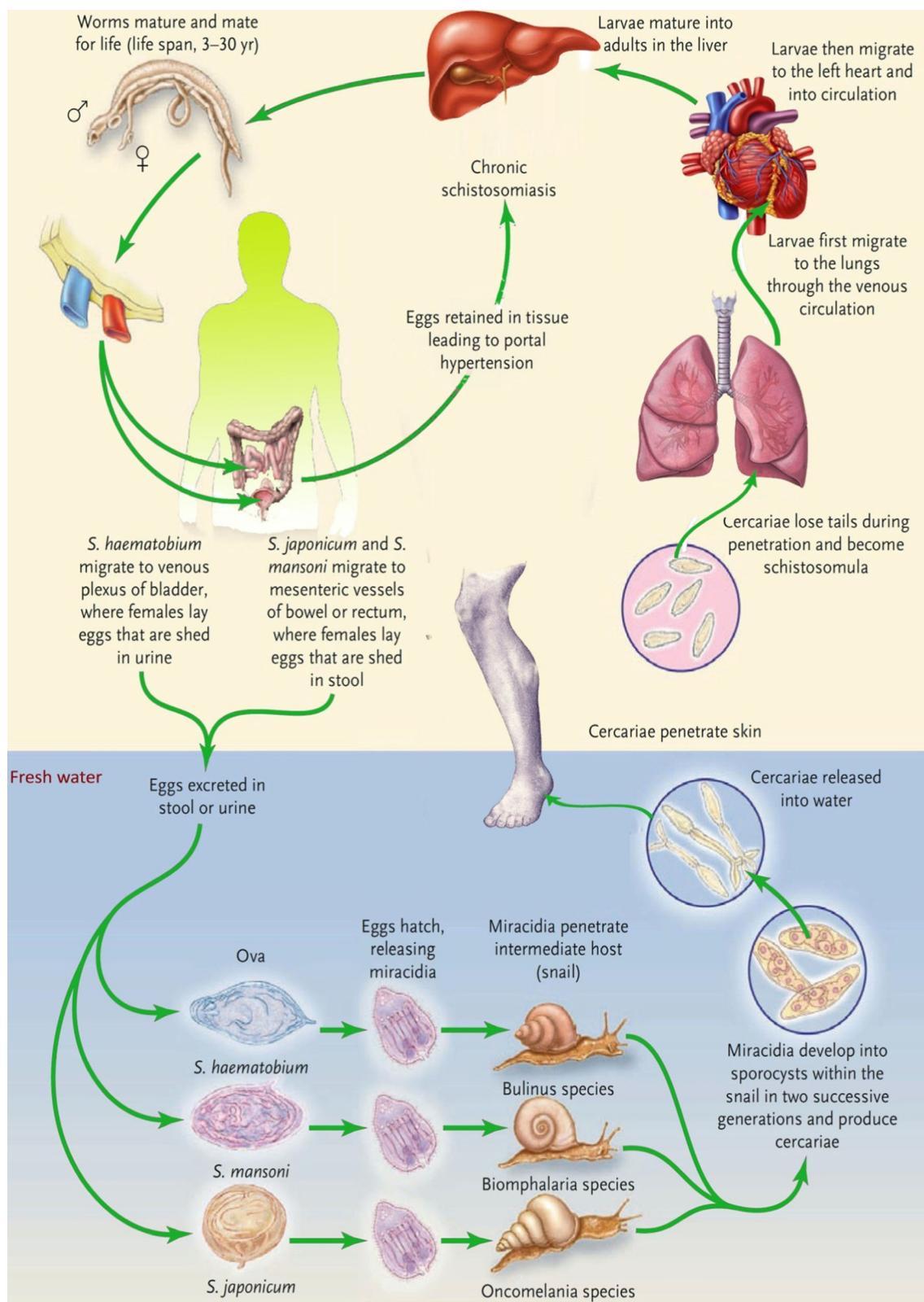
Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Peltophorum africanum</i> Sond	Weeping wattle or African wattie	Leaves + stems	Zimbabwe	At 100 × 10 ³ µg/ml no schistosomes died (<i>S. mansoni</i>)	Mølgård et al. (2001)
<i>Pseudarthria hookeri</i> Wight & Arn	—	Leaves and roots	South Africa	ND	Sparg et al. (2000)
<i>Pterocarpus angolensis</i> DC	Muninga, kiaat	Barks and roots	South Africa	ND	Sparg et al. (2000)
		Leaves	Zimbabwe	At 102 mg/ml no schistosomes died (<i>S. mansoni</i>)	Mølgård et al. (2001)
		Stems	Zimbabwe	At 117 mg/ml no schistosomes died (<i>S. mansoni</i>)	Mølgård et al. (2001)
		Barks	Zimbabwe	At 70 mg/ml for bark some of the schistosomes died (<i>S. mansoni</i>)	Mølgård et al. (2001)
<i>Tetrapleura tetrapetra</i> (Schum. & Thom.) Taub	Aridan	Fruits	Africa	ND	Ojewole (2004)
<i>Stylosanthes erecta</i> P. Beauvois	Egyptian sesban	Aerial parts	Mali	ND	Bah et al. (2006)
<i>Sesbania sesban</i> (L.) Merr	Egypt	Leaves	Egypt	See Table 1	Mahmoud et al. (2011)
<i>Flacourtiaceae</i>					
<i>Flacourtia indica</i> (Burm.f.) Merr	Indian Coffee Plum, Indian plum	Roots	South Africa	ND	Sparg et al. (2000)
<i>Lamiaceae</i>					
<i>Leucas martinicensis</i> (Jacq.) R.Br	—	Whole plant	Mali	ND	Bah et al. (2006)
<i>Liliaceae</i>					
<i>Allium cepa</i> L.	Onion	Bulbs	Mali	See Table 4	Bah et al. (2006)
<i>Aloe buettneri</i> A.Berger	—	Roots	Mali	ND	Bah et al. (2006)
<i>Loganiaceae</i>					
<i>Strychnos henningsii</i> Gilg	—	Barks	South Africa	ND	Sparg et al. (2000)
<i>Meliaceae</i>					
<i>Trichilia emetica</i> Vahl	Natal mahogany	Roots	South Africa	ND	Sparg et al. (2000)

Table 3 continued

Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Menispermaceae</i>					
<i>Cissampelos mucronata</i> A.Rich	—	Roots	South Africa	ND	Sparg et al. (2000)
<i>Mimosaceae</i>					
<i>Entada Africana</i> Guill. & Perr	“Samanére”	Roots	Mali	ND	Bah et al. (2006)
<i>Moraceae</i>					
<i>Ficus thonningii</i> Blume	Strangler figs	Leaves	Mali	ND	Bah et al. (2006)
<i>Nymphaeaceae</i>					
<i>Nymphaea micrantha</i> Guill. & Perr	—	Whole part	Mali	ND	Bah et al. (2006)
<i>Olacaceae</i>					
<i>Ximenia americana</i> L.	Yellow plum	Roots Roots	Mali South Africa	ND ND	Bah et al. (2006) Sparg et al. (2000)
<i>X. caffra</i> Sond	Large sour plum	Roots and leaves	South Africa	ND	Sparg et al. (2000)
<i>Periplacaceae</i>					
<i>Mondia whitei</i> (Hook.f.) Skeels.	“isirigun”	Roots	South Africa	ND	Sparg et al. (2000)
<i>Phytolaccaceae</i>					
<i>Phytolacca dodecandra</i> L'Hér.	Soap berry <i>etihad</i>	Berries	Ethiopia	Investigated in Table 1	Esser et al. (2003)
<i>Poaceae</i>					
<i>Zea mays</i> L.	Maize	Spikes	Mali	ND	Bah et al. (2006)
<i>Polygonaceae</i>					
<i>Securidaca longepedunculata</i> Fresen	Violet tree, fiber tree and Rhodesian violet	Roots	Mali	ND	Bah et al. (2006)
<i>Ranunculaceae</i>					
<i>Clematis brachiata</i> Thunb	Traveler's-joy	Stems and leaves	South Africa	ND	Sparg et al. (2000)
<i>Rhamnaceae</i>					

Table 3 continued

Species	Common name/English Name	Used part	Locality	Activity in lab	References
<i>Ziziphus mucronata</i> Wild	Buffalo thorn	Roots	South Africa	ND	Sparg et al. (2000)
		Roots bark	Zimbabwe	At 101 mg/ml no schistosomes died (<i>S. mansoni</i>)	Mølgård et al. (2001)
<i>Rutaceae</i>					Bah et al. (2006)
<i>Citrus aurantiifolia</i> (Christm.) Swingle	Swingle Lime; Mexican Lime	Leaves and fruits	Mali	ND	Bah et al. (2006)
<i>Sapotaceae</i>					
<i>Vitellaria paradoxa</i> C.F.Gaertn	Shea tree	Roots and leaves	Mali	ND	Bah et al. (2006)
<i>Solanaceae</i>					
<i>Cestrum diurnum</i> L.	Day-blooming jasmine	Leaves	Egypt	See Table 1	Mahmoud et al. (2011)
<i>Datura stramonium</i> L.	Thorn-apple, Jamestown	Leaves	Egypt	See Table 1	Mahmoud et al. (2011)
<i>Physalis angulata</i> L.	Angular winter cherry, balloon cherry, camapu, cutleaf groundcherry	Whole plant	China	See Table 1	dos Santos et al. (2003)
<i>Solanum nigrum</i> L.	Nightshade	Leaves	Egypt	See Table 1	El-Sherbini et al. (2009)
<i>Solanum sinicum</i> Boiss.	—		Egypt	See Table 1	El-Sherbini et al. (2009)
<i>Solanum villosum</i> Mill.	Inab al-deeb or red-fruit nightshade, hairy nightshade, Red nightshade		Egypt	See Table 1	El-Sherbini et al. (2009)
<i>Sterculiaceae</i>					
<i>Melochia arenosa</i> Benth.	—	Aerial parts	Brazil	ND	Truiti et al. (2005)
<i>Vitaceae</i>					
<i>Ampelocissus grantii</i> (Baker)	—	Tuberles	Mali	ND	Bah et al. (2006)
Planch					
<i>Cissus quadrangularis</i> L.	Perandai	Roots, aerial parts and whole part	Mali	ND	Bah et al. (2006)
<i>Zingiberaceae</i>					
<i>Aframomum latifolium</i> K.Schum.	—	Fruits	Mali	ND	Bah et al. (2006)
<i>Zygophyllaceae</i>					
<i>Balanites aegyptiaca</i> (L.) Delile.	Desert date	Roots	Mali	ND	Bah et al. (2006)



►Fig. 1 Life cycle of *Schistosoma*

Surgical Services in Kasr El-Ain Hospital and Medical School, Cairo, discovered *Schistosoma* worms in 1851. Their lifecycle was subsequently described by Robert Leiper, while working in Cairo, in 1915 (Di Bella et al. 2018). The Delta Nile was become a favorite habitat for breeding the host snails of both urinary and intestinal schistosomiasis (Malek 1975).

S. japonicum eggs were discovered in two Chinese ancient bodies dating back roughly to 2180 years ago. Symptoms similar to schistosomiasis were described in the earliest Chinese ancient literature of around 4700 years ago. In the mid-1950s, the first nationwide assessment of the disease indicated that schistosomiasis was endemic in 433 cities across 12 provinces globally, affecting about 11.6 million people (Zhou et al. 2021).

Distribution of schistosomiasis

Schistosomiasis is endemic in 77 countries, and most prevalent in tropical and subtropical regions (Tefera et al. 2020). It is estimated that the severe cases are located in Sub-Saharan Africa; other highly endemic areas are Yemen and the Philippine Island of Mindanao. It is also still endemic, but at lower levels, in North Africa and several Middle East countries. In Asia, it is endemic in parts of the Yangtze River Basin in China, and middle reaches of the Mekong River in Laos and Cambodia (Bergquist and Tanner 2010). In China, a national survey in 2007 revealed that the infection rate among residents in the Fork Beach endemic area was 1.87% (Chen et al. 2012). At least 3 million people in Brazil are thought to be infected by *Schistosoma*, and around 25 million are at risk (Chitsulo et al. 2000). It is also endemic in several Venezuelan states, and several foci in Suriname (Zoni et al. 2016).

Economic implications

Calculating economic effects of schistosomiasis is far from straightforward, but it has been estimated to cause annual losses due to disability (complete or

partial) amounting to US\$ 445, 16, 118, and 60 million in Africa, South-West Asia, South-East Asia and the USA, respectively (at least US\$ 641 million in total). These sums do not include all costs of related public health programs, medical care or compensation for illness, which may also be substantial (Wright 1972). For instance, estimated costs for controlling schistosomiasis morbidity among school-age children by selective and mass chemotherapy are US\$ 0.67 and 0.59 per infected child, respectively (Talaat and Evans 2000). Thus, reviving large-scale school-based health programs to reduce morbidity rates of school-aged children, and to improve their physical growth and cognitive development, would be very costly (Husein et al. 1996).

Globally, the number of people treated for schistosomiasis rose from 12.4 million in 2006 to 33.5 million in 2010 (Carod Artal 2012). US\$150 million have been spent on controlling neglected tropical diseases, including schistosomiasis, in sub-Saharan Africa (Gray et al. 2010). In 2008, a presidential initiative was announced calling for a commitment of US\$ 350 million funding by the USA over five years to combat neglected tropical diseases, and an increase in the number of targeted countries (Liese and Schubert 2009). It has been estimated that 1.2 billion PZQ tablets will be needed annually to treat 400 million people in Africa for at least five years, at an annual cost of US\$100 million (Utzinger et al. 2009). This sum could not be afforded easily within the endemic areas of the world without help from the developed countries. However, much less money would be required if alternative medicine(s) could be produced locally in appropriate financial and ecological frameworks.

Geographical implications

In many areas with high rates of schistosomiasis the local health systems are under-resourced (Amazigo et al. 2012) and subject to diverse disturbances that disrupt ecological equilibria and contexts, within which disease hosts or vectors and parasites breed, develop, and transmit the disease (Cable et al. 2017). Thus, they may strongly influence the emergence and proliferation of parasitic diseases, including schistosomiasis. It has been estimated that 97% of the infections are on the African continent (Steinmann et al. 2006), at least partly due to the lack (or paucity)

of health systems, poverty and general neglect (Utzinger et al. 2011). Furthermore, 95% of all parasitic infections occur in the developing world, since it has the most conducive combinations of human behavior, anthropogenic disturbance and both climatic and physical conditions (Sattenspiel 2000). Notably, in endemic parts of the world schistosomiasis is intimately connected to the construction of infrastructure such as small, multipurpose dams and large hydroelectric dams for power production and irrigation systems (Utzinger et al. 2011). In the modern era, for example, the Aswan High Dam caused ecological changes that stimulated disease transmission (Malek 1975). Schistosomiasis remains the most important water-borne disease, and it was promoted by prevailing socio-ecological systems, since transmission is governed by unsafe human behavior (*e.g.* unprotected direct surface water contacts and open defecation) (Acka et al. 2010).

In Egypt, recent projects have been carried out in many regions to reclaim land from the desert for agriculture using water of the Nile River. This, together with the increased human activities in the reclaimed areas, has stimulated transmission of both *S. mansoni* and *S. haematobium*, manifested by the presence of infected *B. truncatus* and *B. alexandrina*, the intermediate host snails for the respective parasitic species, in these areas. Parts of the desert utilizing Nile water has wider spread of schistosomiasis (El-Kady et al. 2000). Schistosomiasis became a national burden and emerging as a major public health problem in the Egyptian reclaimed areas unless adequate control measures are taken (Abou-El-Naga 2018). There are also cases of human infection in American and European metropolitan areas, due to immigrations (Fuentes et al. 2010; Roure et al. 2017).

Medical implications

Hepatitis has been detected in 70% of surveyed schistosomiasis patients, and *S. mansoni* is one of the two major risk factors, together with viral hepatitis, for chronic liver disease and liver cirrhosis. Thus, millions of the world's population live under constant risk of both schistosomiasis and hepatitis (Halim et al. 1999). Similarly, clinical history of urinary schistosomiasis is associated with significantly increased risk of bladder cancer (Bedwani et al. 1998). Furthermore,

schistosomiasis can infect the children in endemic populations, and disease symptoms in children include anemia, liver fibrosis, immunity impairment, and severe physical and mental disorders (El Baz et al. 2003).

The free-swimming *Schistosoma* larvae penetrate the skin, causing rashes, erythema and itchy skin prior to fever, chills, cough and muscle aches. As the parasite matures in the host veins (mesenteric or vesical), the symptoms change dramatically, and blood becomes visibly present in patients' urine and stool (Kolářová et al. 2013; Nation et al. 2020). Under chronic conditions and if parasite eggs damage the organ in which they were deposited, the carrier can suffer from liver, kidney, and bladder complications. Advanced intestinal schistosomiasis is manifested with enlarged liver and spleen, fibrosis, and portal hypertension while symptoms of advanced uro-genital schistosomiasis include hydronephrosis and calcification of the bladder (Andrade 2009; Salas-Coronas et al. 2020).

Social implications

Advances in understanding *Schistosoma* epidemiology, and the greater availability of effective diagnosis and new tools have improved both current management and prospects for refining control of schistosomiasis. Nevertheless, infection and morbidity rates of the disease are still high, particularly in poor and otherwise disadvantaged populations. This disease of poverty has proved to be difficult to control for centuries (Utzinger et al. 2011). Despite prolonged mass antiparasitic drug therapy programs and other control measures, it has not been eradicated and continues to spread to new geographical areas (Siddiqui et al. 2011). Globally, WHO has estimated that about (600–779) million people are at risk of infection due to their exposure to contaminated water and 200–209 million people are infected with these parasites (Steinmann et al. 2006; WHO 2012). Schistosome infections require direct contact in water between the skin and the infective cercariae, thus sociocultural factors are strongly linked to infection rates, as illustrated in Fig. 2. In addition, recording and reporting of incidences, prevalence and death rates remain a challenge, and further efforts are needed to improve tracking the disease both nationally and



Fig. 2 Infection by schistosomes is strongly associated with social activities (washing clothes, bathing, etc.) in which people come into direct contact with freshwater inhabited by schistosomes' intermediate host snails

globally. Accordingly, schistosomiasis is still considered one of the major health, socio-economic and developmental challenges facing many of the world's poorest countries.

Strategies to eliminate schistosomiasis

There is an urgent need to increase global awareness and support endemic countries' endeavors to develop appropriate methods to control the disease (Rollinson et al. 2013). As already mentioned, it is generally agreed that no single method will be sufficient to eliminate schistosomiasis; integrated approaches will be required due to its complexities (Mo et al. 2014). Application of molluscicides to reduce intermediate host snail populations is one of the most efficient methods for controlling the disease (Rapado et al. 2011). However, there are four major strategies for eradication: (1) treating infected individuals to reduce morbidity and mortality, and preventing the spread of *Schistosoma* parasite eggs, (2) providing communities with adequate, appropriate sanitation and accessible safe water to reduce environmental contamination and hence minimize the chances of miracidia transmission, (3) snail control to block the lifecycle of the parasite, and (4) health education (Fig. 3) (Chimbari 2012). To

accelerate progress, a wider application of existing interventions combined with implementation of new methods in a manner tailored to the socio-ecological setting is required. In China, a comprehensive control strategy to reduce rates of *S. japonicum* transmission from humans and cattle to snails was evaluated from 2005 to 2007. The strategy included removing cattle from snail-infested areas, providing farmers with mechanical equipment, improving sanitation and implementing health-education programs in the endemic areas. This integrated strategy reportedly reduced *S. japonicum* infection in humans to less than 1% in an endemic area (Wang et al. 2009).

Medical control of schistosomiasis by treating infected humans

A systematic search for chemotherapeutic drugs has been ongoing for several decades. The synthesis of PZQ in 1970 revolutionized the treatment and led to dramatic reductions in morbidity and mortality rates. PZQ is the only drug being used to treat human schistosomiasis on a large scale (Doenhoff et al. 2008; Mordvinov and Furman 2010), and is highly recommended in disease-control programs (Salvador-Reca-tà and Greenberg 2012). The scope of therapeutic programs, as treatment with PZQ has largely been

restricted to children attending school due to the lack of infrastructure and other logistical problems in many areas of the developing world (Siddiqui et al. 2011).

Although treatment of schistosomiasis in patients worldwide relies heavily on PZQ and the drug has thoroughly recognized efficacy against schistosomes (Keiser et al. 2010), evidence is now accumulating that PZQ cannot prevent re-infection, and may sometimes even exacerbate it (Chandiwana et al. 1991; Doenhoff et al. 2008). Moreover, PZQ resistance is emerging in endemic areas due to repeated use (Wang et al. 2012). Obviously, there is a need for new alternatives for treating schistosomiasis (Cioli et al. 2014; Bergquist et al. 2017). Important potential sources are plants, and already many have been tested and some showed relevant activities, especially members of the medicinal plants. Many of them have been already tested and some showed relevant activities, especially members of the plant families Euphorbiaceae, Annonaceae, Fabaceae Asteraceae, Asphodelaceae, and Asparagaceae (Tables 1, 2, 3, 4 and 5).

Strategic controls of schistosomiasis by eliminating intermediate host-snails

Schistosoma parasites, like other parasitic helminths, can survive in their hosts for a long time, but the intermediate host is the weakest link in the

transmission cycle. Thus, molluscicides are widely used as experimental models in programs to investigate the disease by killing the intermediate host snails (mollusks) (Wang et al. 2018), thereby disrupting the parasite's lifecycle and stopping transmission to people in contact with water in high-risk areas (Birley 1991).

Synthetic molluscicides

The major synthetic molluscicides that are currently used to control the snail vectors are metaldehyde, niclosamide, carbamate, organophosphate, and synthetic pyrethroids (Singh et al. 2010). Niclosamide is one of the most widely used synthetic molluscicides at present. However, it is highly toxic towards non-target organisms, including fish, and it is also ecologically destructive (He et al. 2017). Nevertheless, it is one of the most important molluscicides approved by WHO due to the lack of robust alternatives (He et al. 2017). It is most active against *O. hupensis* snails with an LC₅₀ of 0.12 µg/ml and LC₉₀ of 0.98 µg/ml after 24 h (Chen et al. 2007) but it kills amphibians and fish if used in effective concentrations. There are also several chemical molluscicides (e.g. copper sulfate, calcium cyanamide, chlorinated lime and carbamate derivatives). However, they are environmentally hazardous,



Fig. 3 Strategies for schistosomiasis eradication

Table 4 Extracts with recognized larvicidal activity

Species (plant part)	Vernacular name	Type of the extract	Results	References
<i>Amaryllidaceae</i>				
<i>Allium cepa</i> L.	Onion	ND	$LC_{50} = 50 \mu\text{g/ml}$ (miracidia) $LC_{50} = 50 \mu\text{g/ml}$ (cercariae) (<i>S. mansoni</i>)	Mantawy et al. (2012)
<i>Allium sativum</i> L.	Garlic	ND	$LC_{50} = 100 \mu\text{g/ml}$ (miracidia) $LC_{50} = 10 \mu\text{g/ml}$ (cercariae) (<i>S. mansoni</i>)	Mantawy et al. (2012)
<i>Burseraceae</i>				
<i>Commiphora molmol</i> (Engl.) Engl. ex Tschirch (Arabian or Somali gum)	Myrrh	Oil	$LC_{100} = 2.5\text{--}10 \mu\text{g/ml}$ 60–90 min (cercariae) (<i>S. haematobium</i>)	Masoud et al. (2000)
<i>Euphorbiaceae</i>				
<i>Jatropha curcas</i> L. (Seeds)	Physic nut	Methanol	$LC_{100} = 25 \mu\text{g/ml}$ (cercariae) $LC_{50} = 5 \mu\text{g/ml}$ (miracidia) (<i>S. mansoni</i>)	Rug and Ruppel (2000)
		Water	$LC_{50} = 5 \text{ mg/ml}$ (miracidia and cercariae) (<i>S. mansoni</i>)	Rug and Ruppel (2000)
		Crude oil	250 $\mu\text{g/ml}$ killed all cercariae in 80 min 100 $\mu\text{g/ml}$ kills all larvae within 2 h (<i>S. mansoni</i>)	Rug and Ruppel (2000)
<i>Lamiaceae</i>				
<i>Plectranthus tenuiflorus</i> (Vatke) Angew (Leaves)	—	Methanol	$LC_{50} = 17.39 \text{ mg/ml}$ (cercariae) $LC_{50} = 24.37 \text{ mg/ml}$ (miracidia) (<i>S. mansoni</i>)	Aziz et al. (2011)
<i>Solanaceae</i>				
<i>Solanum nigrum</i> L. (Leaves)	Nightshade	Water	$LC_{100} = 30 \mu\text{g/ml}$ (<i>S. haematobium</i>) $LC_{100} = 30 \mu\text{g/ml}$ (<i>S. mansoni</i>) $LC_{100} = 40 \mu\text{g/ml}$ (<i>F. gigantica</i>) after 30 min (cercariae)	Ahmed and Ramzy (1997)

Table 5 Compounds with antischistosomiasis of water larvacidal activity isolated from higher plant

Common name (chemical name)-/class	Species (part)/family	Separation tools	Spectral analysis	Results	References
2-hydroxychrysophanol (1,2,8-trihydroxy 3-methylanthraquinone) (Anthraquinone) (38)	<i>Hemerocallis fulva</i> (L.) L. (root)/ Hemerocallidaceae	Repeated ODS, Sephadex LH- 20 gel CC	UV, IR, (EI & HR-EI) MS, ^1H -& ^{13}C -NMR	Immobilized all cercariae within 15 s at 3.1 $\mu\text{g}/\text{ml}$	Cichewicz et al. (2002)
Kwanzoquine E/ (Anthraquinone) (41)	<i>Hemerocallis fulva</i> (L.) L. (Root)/ Hemerocallidaceae	Sephadex LH-20 CC, ODS preparative HPLC	UV, IR, (EI & HR-EI) MS, ^1H -& ^{13}C -NMR, DEPT, HMBC	Immobilized cercariae within 12–14 min at 25 $\mu\text{g}/\text{ml}$	Cichewicz et al. (2002)

and snails develop resistance to them (Bilia et al. 2000b).

Available molluscicides are also costly, which poses major problem for low-income countries where schistosomiasis is widely distributed. Despite all recent research and development current synthetic molluscicides also raise serious environmental concerns and threats to both human health and income (particularly in areas where fishing is a major source of food and income) due to their lack of specificity (Andrews et al. 1982; He et al. 2017). Thus, identifying or developing more selective molluscicidal herbal preparations that do not have adverse effects

(or at least acceptably weak effects) in non-target aquatic organisms and are biodegradable remains a high priority (El-Sherbini et al. 2009).

Plant molluscicides

Several thousand synthetic compounds have been tested for their ability to control host snails but none of them was proven to be as entirely satisfactory as yet. Therefore, natural sources of novel agents came into play and are considered now as attractive alterantives to seek for novel drug leads (Ribeiro et al. 2021; Xing et al. 2021). Primary target sources are medicinal

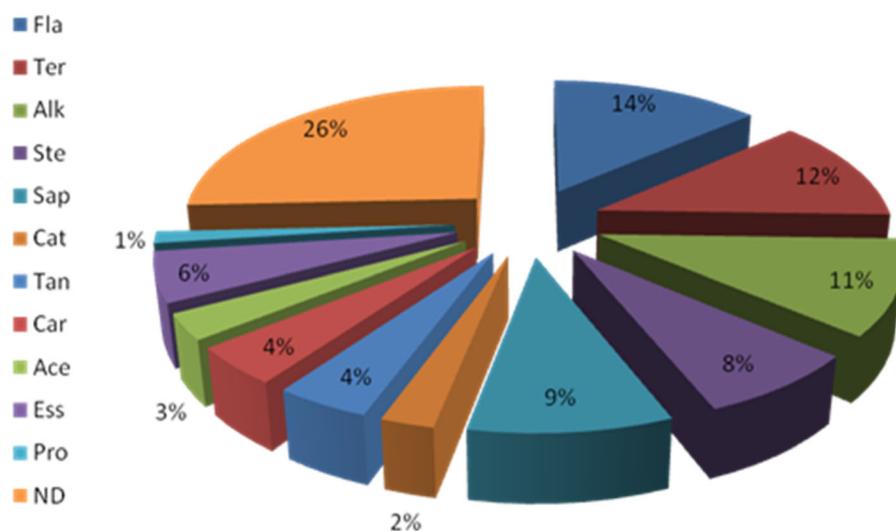


Fig. 4 The most bioactive of natural compounds against snails based on a review of literature: Fla, flavonoids; Ter, Terpenes; Alk, alkaloids; Ste, steroids; Sap, saponins; Cat, catechines; Tan, tannins; Car, carbohydrates; Ace, acetogenine; Ess, essential oils; Pro, proteins; ND, not defined

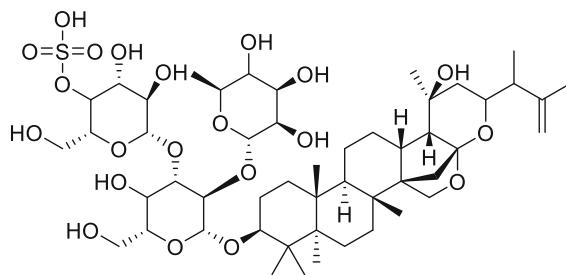


Fig. 5 A novel anti-schistosomiasis triterpene glycoside (Asparagalin A) (1)

plants because they are generally a rich source of renewable bioactive organic chemicals, and they may grow well in endemic areas, providing income for local farmers. Furthermore, plant derived molluscicides may have several advantages like low costs, high target specificity, water solubility, high biodegradability, and low toxicity towards normal organs in the human hosts (Singh et al. 2010).

Since 1982, the potential utility of more than 150 plant species for controlling freshwater snails has been tested. Numerous groups of compounds plant-derived compounds were toxic to target organisms at acceptable doses, ranging from < 1 to 100 µg/ml (Singh et al. 2010). The molluscicidal activity of plant extracts and often their active compounds have been reported (Tables 1 and 2). Commonly identified active compounds belong to saponins, alkaloids, terpenoids and tannins (Fig. 4). The plant extracts evaluated against the intermediate host (snails and larvae of *Schistosoma*) are listed in Table 4. These include, for instance, methanol and aqueous extracts of *Jatropha curcas* L. (Euphorbiaceae) on snails transmitting *S. mansoni* and *S. haematobium* (Rug and Ruppel 2000). Different parts were traditionally used as antimicrobial agents. The plant had traditional uses as antimicrobial agents. Plant species known to be rich in bioactive molluscicidal compounds are presented in Tables 2 and 5. Tannins are particularly suitable for snail control because tannin-containing plants are not only widely distributed but tannins can be also relatively easily isolated (Al-Sayed et al. 2014). The highest contents of molluscicidal agents have been found in members of Euphorbiaceae followed by Annonaceae, Mimosaceae, Polygonaceae, Verbenaceae and Caesalpinae families, the active compounds of a number of plants are presented in (Fig. 4).

More extensive efforts are warranted to isolate and identify the remaining unknown compounds from other plants, that could meet all the desirable criteria of molluscicides.

Classes of compounds used as plant molluscicides

Saponins Saponins comprise a structurally diverse class of triterpenes, based on triterpenoids often in the form of glycosides. They occur in plant species and have been also isolated from marine organisms (Challinor and De Voss 2013). Saponins derive their name from the soapwort plant, genus *Saponaria* (Caryophyllaceae). The triterpene skeletons of saponins are formed from the C₃₀ precursor oxisqualene. They are diverse, and decorated with different functional groups in different plant species (Challinor and De Voss 2013). They have hemolytic properties, toxic effects on most cold-blooded animals and proven molluscicidal activity (Francis et al. 2002; Sparg et al. 2004).

Triterpenoids and saponins are widespread in nature. The most active molluscicidal saponin compounds have an oleanolic acid-based aglycone and trisaccharide sugar moiety (Mølgaard et al. 2000). This is consistent with our recent finding that asparagalin A (Fig. 5) (1), a triterpene saponin from the Egyptian species *Asparagus stipularis* Forssk., suppresses *S. mansoni* egg-laying. Shoots and roots of this plant used in folk medicine as diuretic for curing jaundice, liver ailments, against bilharzias (El-Seedi et al. 2012). Sometimes molluscicidal compounds are formed by an enzymatic reaction during extraction of plants, for instance if *Phytolacca dodecandra* L'Hér. berries are crushed with water (Parkhurst et al. 1989).

The desert tree *Balanites aegyptiaca* (L.) Delile (common name Higleeg) (Suleiman 2015) has a well-known molluscicidal activity. It was the first plant reported for the control of schistosomiasis owing to its saponins content (Marston and Hostettmann 1985). The fruits of this plant were used as remedy to eradicate intestinal parasites. This plant species is traditionally (Table 3) used in treatment of malaria and is apparently suitable for vector control (Chothani and Vaghasiya 2011).

Pulsatilla chinensis (Bunge) Regel displayed potent molluscicidal activity against *O. hupensis*, attributable to the presence of the active triterpenoid saponins. The plant roots are widely used in traditional

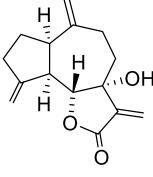
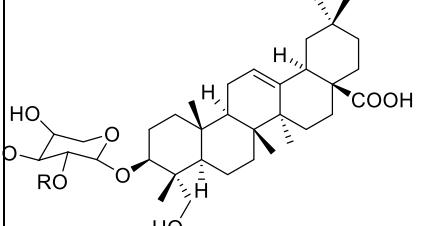
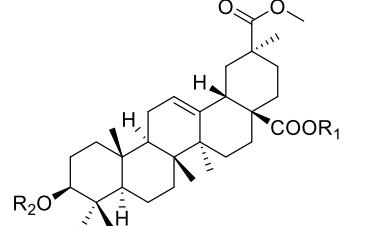
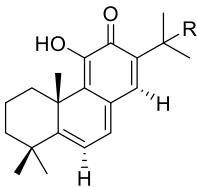
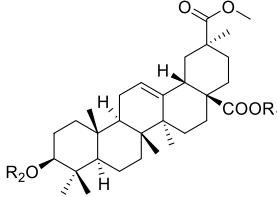
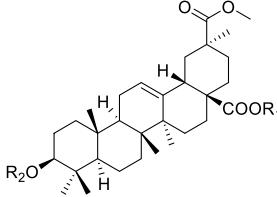
 <p>7α-hydroxy-3-desoxyzaluzanin C (5)</p>	 <p>3-O-[α-L-arabinopyranosyl]-hederagenin (6) R= H</p> <p>3-O-[α-L-rhamnopyranosyl -(1\rightarrow2)-α-L-arabinopyranosyl]-hederagenin (8) R= Rha</p> <p>3-O-[β-D-glucopyranosyl-(1\rightarrow2)-α-L-arabinopyranosyl] hederagenin (9) R= Glc</p>	 <p>3-O-(β-D-galactopyranosyl-(1\rightarrow3)-β-D-glucopyranosyl) serjanic acid 28-O-β-D-glucopyranoside (7) R₁=Glc, R₂=Glc³-Gal</p>
 <p>15-deoxyfuerstione (10) R=H</p> <p>Fuerstione (13) R=OH</p>	 <p>3-O-(α-L-rhamnopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl) serjanic acid 28-O-β-D-glucopyranoside – saponin (11) R₁=Glc, R₂=Glc²-Glc²-Rha</p> <p>3-O-(β-D-glucopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl) serjanic acid 28-O-β-D-glucopyranoside (12) R₁=Glc, R₂=Glc²-Glc</p> <p>3-O-(β-D-galactopyranosyl-(1\rightarrow3)-β-D-glucopyranosyl) serjanic acid (16) R₁=H R₂=Glc³-Gal</p> <p>3-O-(β-D-glucopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl) serjanic acid (17) R₁=H R₂=Glc²-Glc</p> <p>3-O-(β-D-glucopyranosyl-(1\rightarrow3)-β-D-glucopyranosyl-(1\rightarrow3)-β-D-glucopyranosyl) serjanic acid (19) R₁=H R₂=Glc³-Glc³-Glc</p> <p>3-O-(α-L-rhamnopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl) serjanic acid (20) R₁=H R₂=Glc²-Glc²-Rha</p>	 <p>3-O-(β-D-galactopyranosyl-(1\rightarrow3)-β-D-glucopyranosyl) serjanic acid 28-O-β-D-glucopyranoside (7) R₁=Glc, R₂=Glc³-Gal</p>

Fig. 6 Structures of molluscidicidal bioactive compounds

Chinese medicine as remedies for amebiasis, malaria, vaginal trichomoniasis and bacterial infections. Treatment with a sub-lethal concentration reportedly

caused significant inhibition of acetyl cholinesterase (AHE), alanine transaminase (ALT), and alkaline

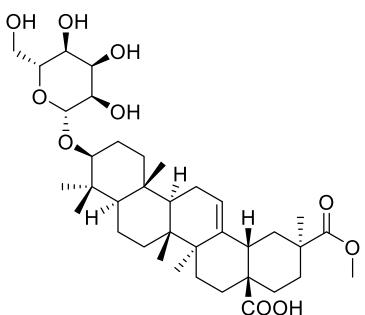
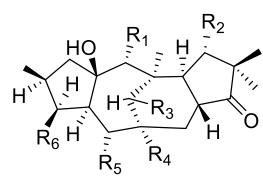
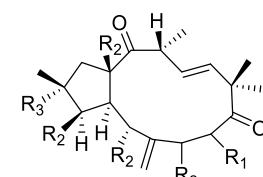
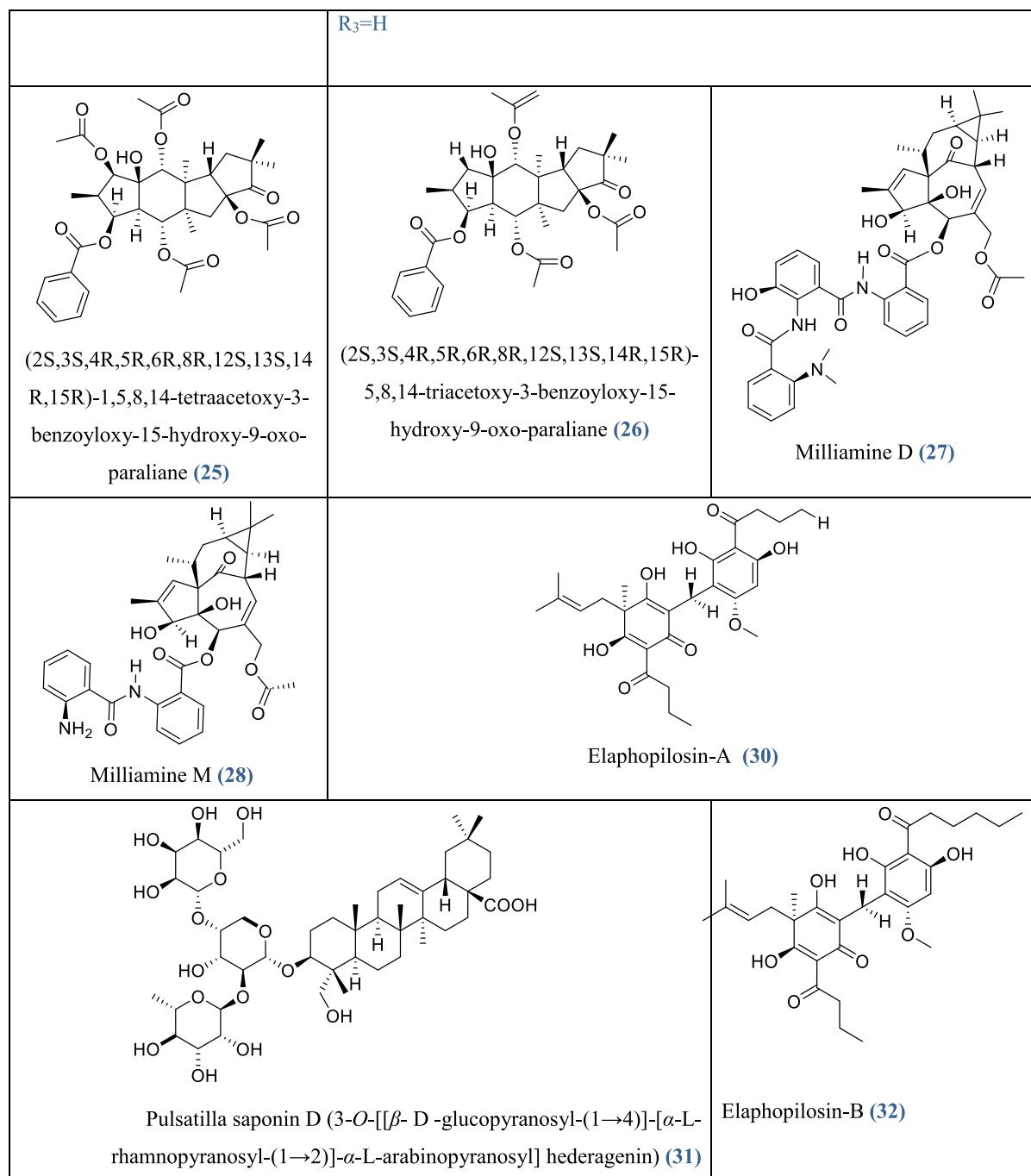
 <p>3-O-β-D-glucopyranosylserjanic acid (14)</p>	 <p>(2S,3S,4R,5R,6R,8R,11S,12S,13R,14R,15R)-6,14,17-triacetoxy-5-(2-hydroxyacetoxy)-3-benzoyloxy-15-hydroxy-9-oxo-segetane-diterpene (15) $R_1=R_3=R_4=OCOCH_3$, $R_2=H$, $R_5=OCOCH_2OH$, $R_6=OCOC_6H_5$</p> <p>(2S,3S,4R,5R,6R,8R,11S,12S,13R,14R,15R)-6,11,14,17-tetraacetoxy-5-(2-acetoxyacetoxy)-3-benzoyloxy-15-hydroxy-9-oxo-segetane (21) $R_1=R_2=R_3=R_4=OCOCH_3$, $R_5=OCOCH_2COOCH_3$, $R_6=OCOC_6H_5$</p> <p>(2S,3S,4R,5R,6R,11S,13R,14S,15R,17R)-5,6,11,14,17-pentaacetoxy-3-benzoyloxy-15-hydroxy-9-oxo-seget-8(12)-ene diterpene (22) $R_1=R_2=R_3=R_4=R_5=OCOCH_3$, $R_6=OCOC_6H_5$</p> <p>(2S,3S,4R,6R,8R,11S,12S,13R,14R,15R)-5,6,11,14,17-pentaacetoxy-3-benzoyloxy-15-hydroxy-9-oxo-segetane-diterpene (23) $R_1=R_2=R_3=R_4=R_5=OCOCH_3$, $R_6=OCOC_6H_5$</p>
	 <p>(2R,3R,4S,5R,7S,8R,13R,15R)-2,3,5,7,15-pentaacetoxy-8-iso-butyroyloxy-9,14-dioxojatropa-6(17),11E-diene (18) $R_1=OCOCH(CH_3)_2$, $R_2=R_3=OCOCH_3$</p> <p>(2R,3R,4S,5R,7S,8R,13R,15R)-2,3,5,7,15-pentaacetoxy-8-angeloyloxy-9,14-dioxo-jatropa-6(17),11E-diene (24) $R_1=R_2=R_3=OCOCH_3$</p> <p>(2R,3R,4S,5R,7S,8R,13R,15R)-3,5,7,15-tetraacetoxy-8-isobutyroyloxy-9,14-dioxo-jatropa-6(17),11E-diene (29) $R_1=OCOCH(CH_3)_2$, $R_2=OCOCH_3$</p>

Fig. 6 continued

phosphatase (ALP) activities in the liver and the cephalopodium of *O. hupensis* (Chen et al. 2012).

Extracts of *Calendula officinalis* L. and *Ammi majus* L. (common name, Khilla sheitani) have molluscicidal activity against *Bullinus truncates* and

**Fig. 6** continued

B. Alexandria (less strongly). The recorded LC₅₀ and LC₉₀ values indicated that *C. officinalis* L. is more toxic to both snails where *A. majus*, and *Bullinus truncatus* snails are more sensitive to the extracts of both plants than *B. alexandrina*. Prolonged exposure

to sub-lethal concentrations of *A. majus* L. affected the egg-laying and survival of both snails. In addition, treatment with sub-lethal doses of extracts of both plants clearly inhibited transaminase activity, diminished the total protein content, and markedly increased

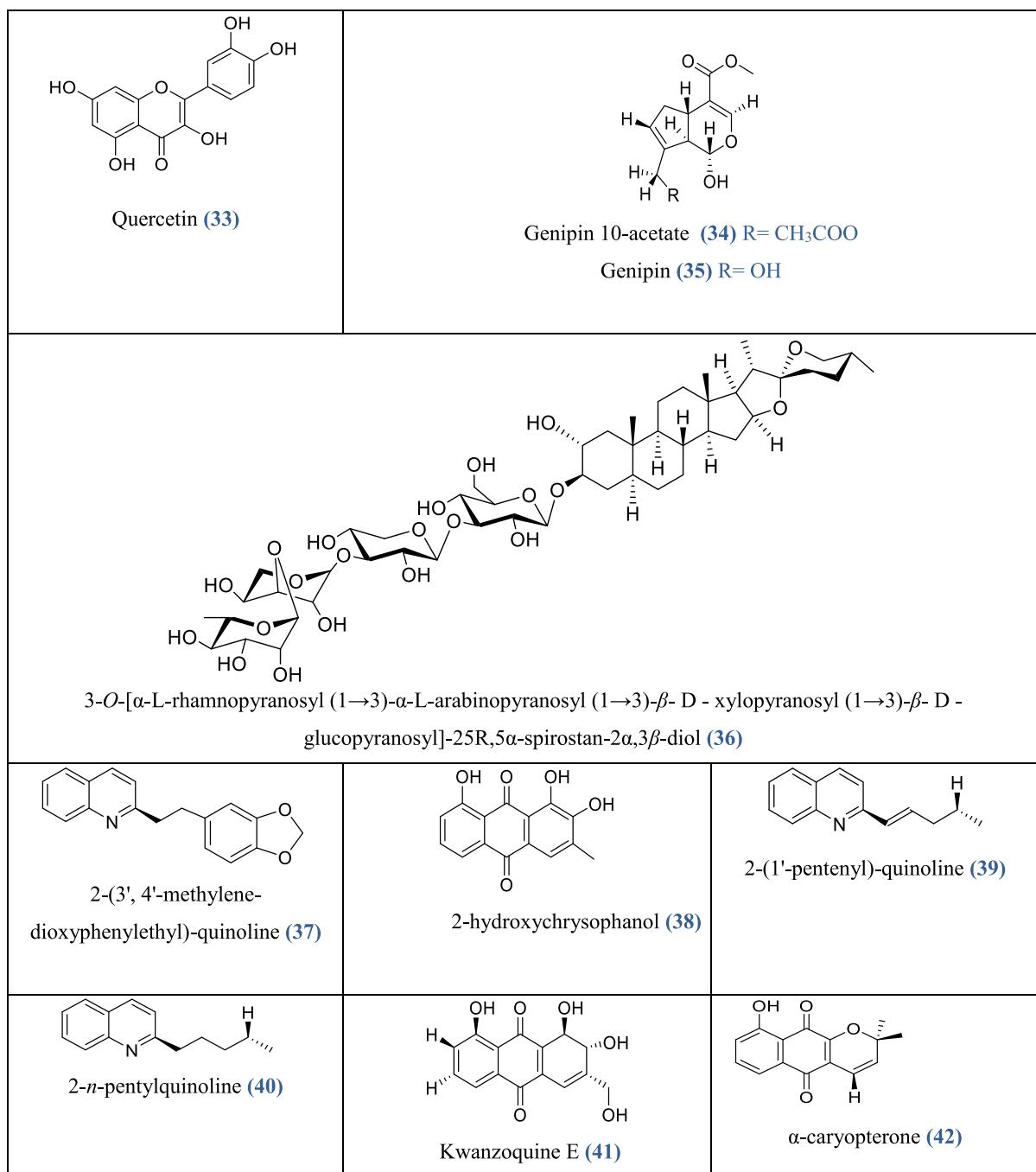
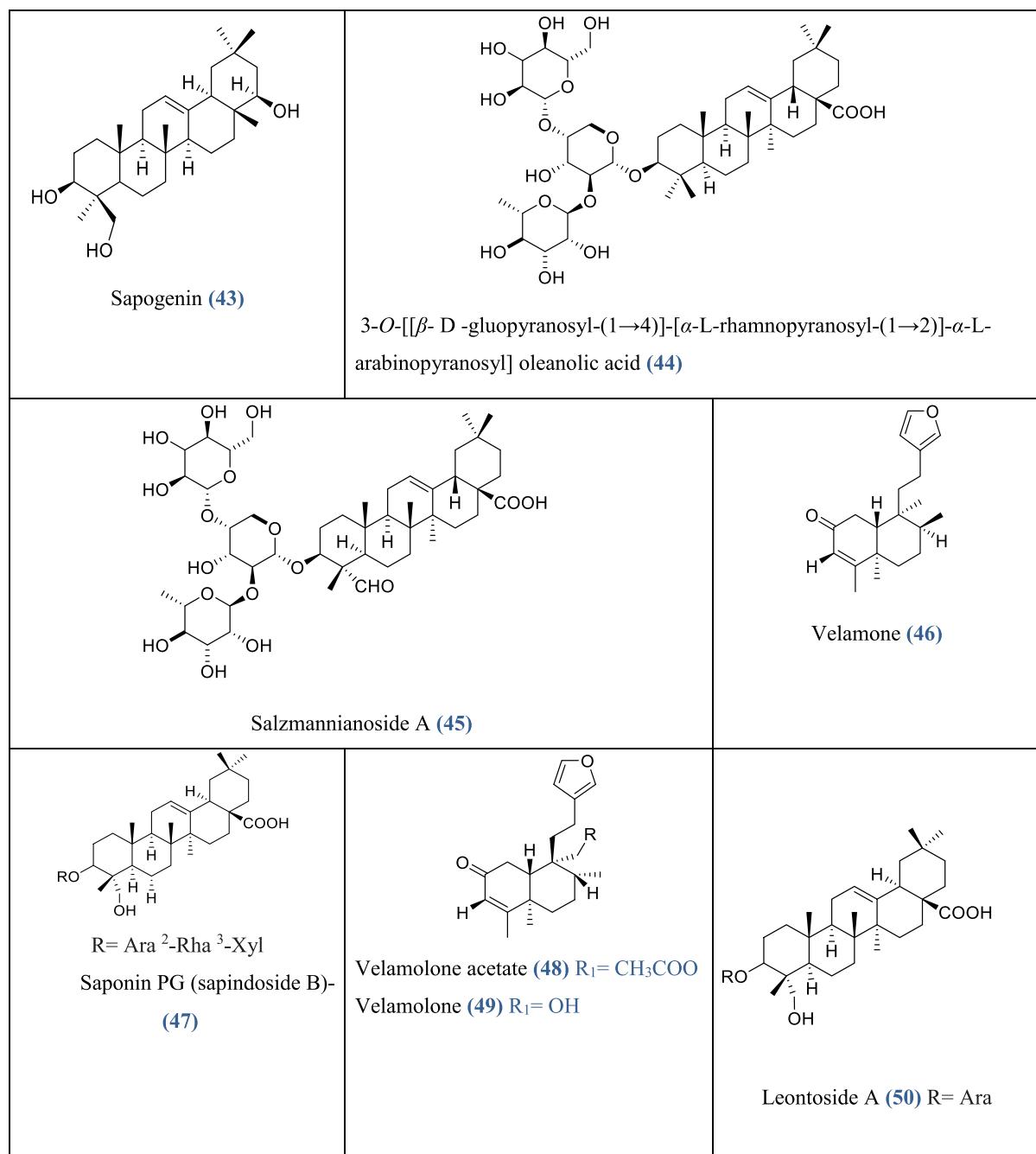


Fig. 6 continued

total lipid contents in both snails hemolymph (Rawi et al. 1996). The activity of *A. majus* L. is thought to be due to saponins, and furocoumarins, although the

effect seems to be highly snail species-dependent (Marston and Hostettmann 1985; Rawi et al. 1996).

Phytolacca dodecandra L'Hér. is one of the most active molluscicidal plants, and thus has been widely

**Fig. 6** continued

studied. Its activity was initially reported by Lemma's group (Lemma 1965, 1970) and the presence of an active saponin named lemmatxin was demonstrated (Parkhurst et al. 1974). Aqueous extracts of the dried berries contain up to 25% saponins (Treyvaud et al.

2000) which were stable in water for two days at room temperature (Mølgaard et al. 2000). Some isolated saponins are lethal to snails at very low concentrations. However, a major disadvantage of using saponins as molluscicides is that they are also

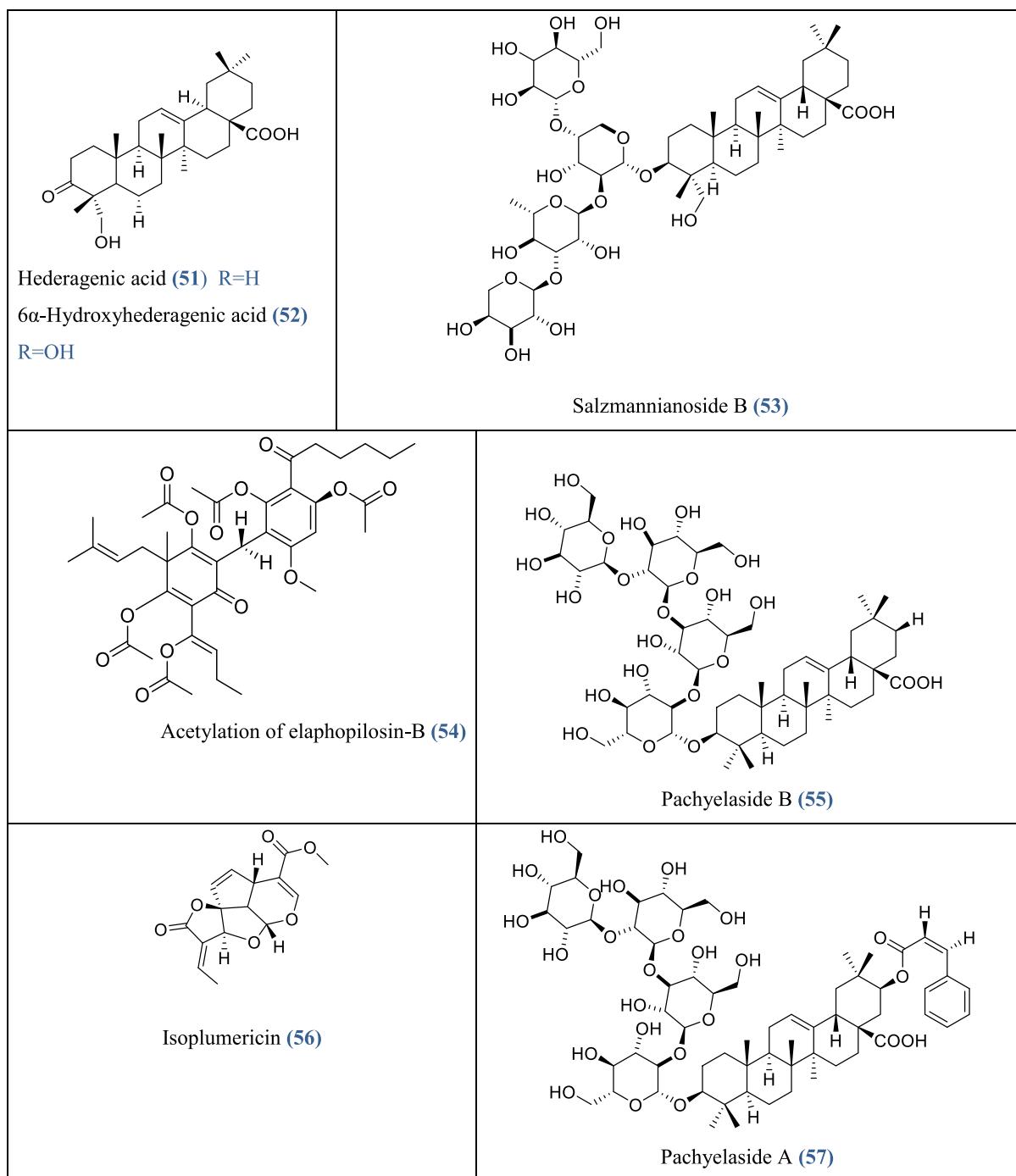
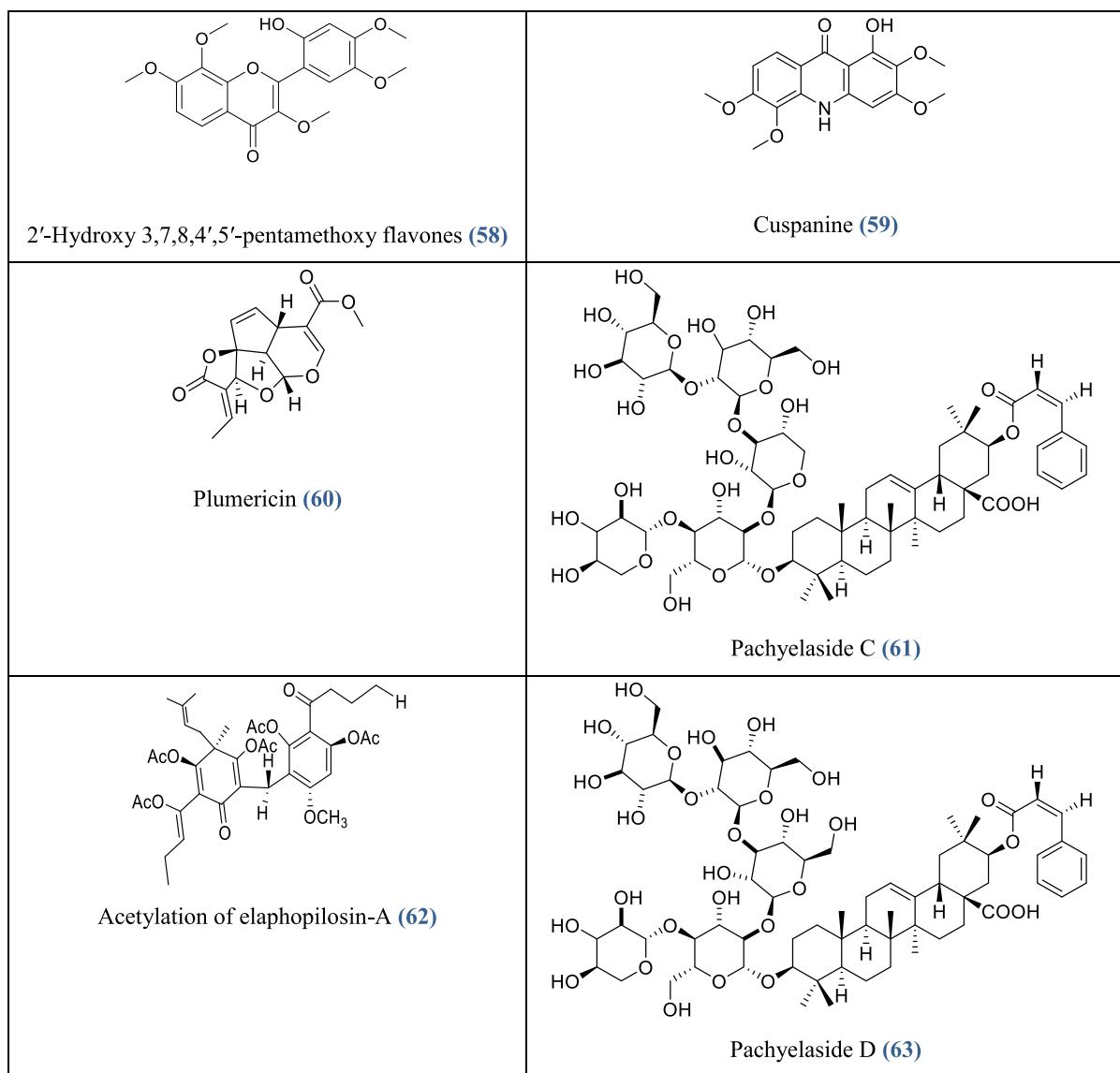


Fig. 6 continued

generally lethal to fish compared to flavonoids and other phenolics that are generally less toxic to non-target organisms (Wei et al. 2002).

Glycoalkaloids Several glycoalkaloids (Fig. 6) identified from *Solanum* species are molluscicidal. Extracts of *S. elaeagnifolium* Cav. berries, and *S.*

**Fig. 6** continued

sodomaeum Dunal leaves and seeds reveal molluscicidal activity against *Bu. truncates*, the intermediate host of *S. haematobium* (Table 2) (Bekkouche et al. 2000). In addition, *S. sisymbriifolium* Lam. fruits are active against the snails, the active fractions contain the steroid alkaloids solamargine (Fig. 7) (2) and β -solamarine (Bagalwa et al. 2010), but solamargine is the most effective of these compounds, due to its specific sugar substitution (Miranda et al. 2012). Solamargine

(containing a chacotriose sugar chain moiety) is also more active than solasonine, (containing a solatriose sugar chain moiety) against adult *S. mansoni* worms (Miranda et al. 2012).

Phorbol esters Phorbol (Fig. 8) (3) was initially isolated in 1934 as a hydrolysis product of *Croton tiglium* L. vegetable oil (Flaschenträger, v. Wolffersdorff 1934; Abegaz and Kinfe 2020), and its structure was subsequently elucidated by Hecker and

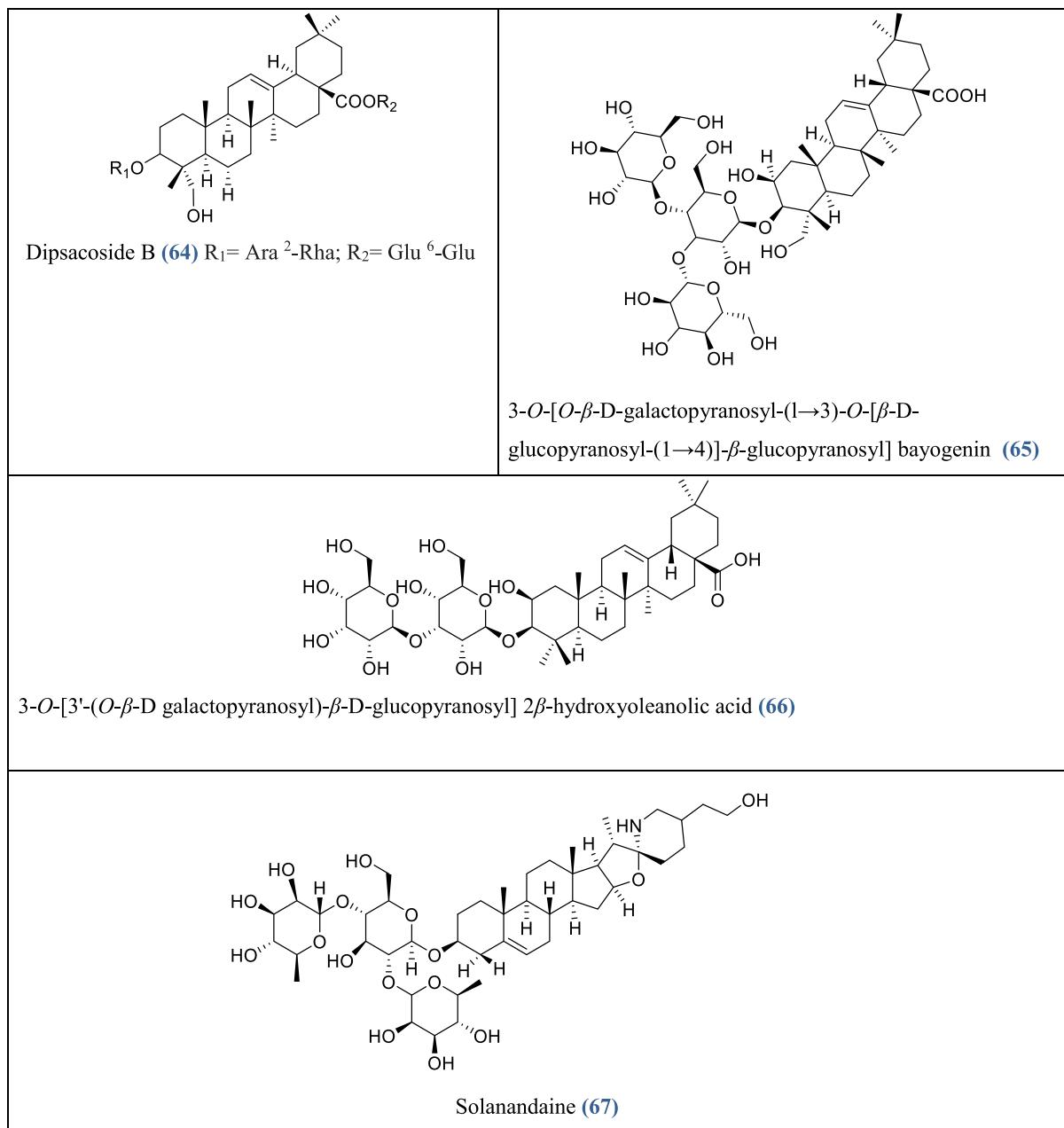
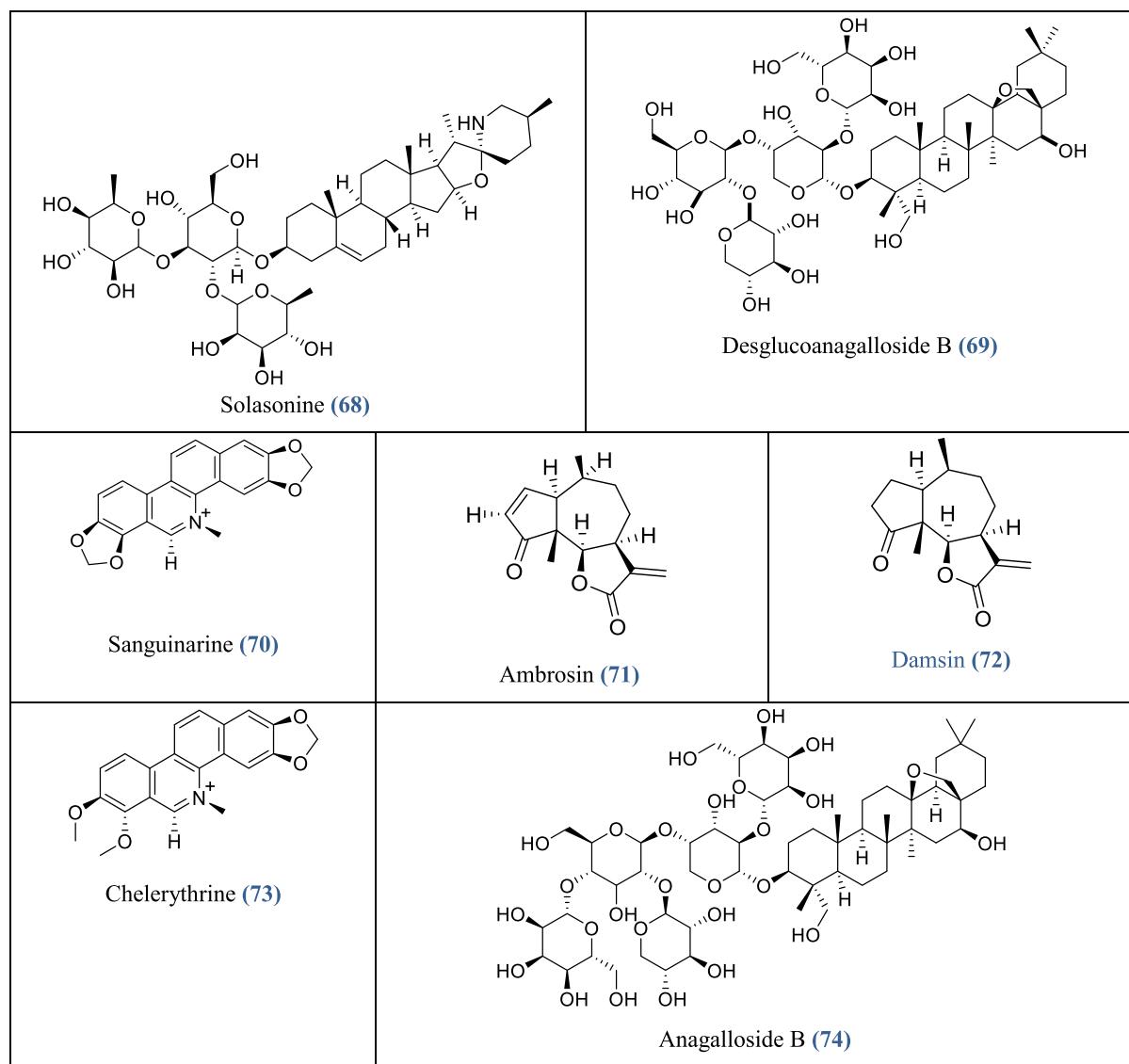


Fig. 6 continued

colleagues (Hecker 1967). Phorbol esters are diterpenes, which are, among others, major constituents of *Jatropha curcas* L. oil. They exhibit molluscicidal activity against aquatic snails (Liu et al. 1997). Some of the phorbol esters are mutagenic and

thus not suited as a molluscicidal medicine (Liu et al. 1997).

Phorbol esters (and anthraquinone) are also the most bioactive compounds tested against miracidia and cercaria (Fig. 9). There have been few attempts to

**Fig. 6** continued

isolate bioactive water larvicidal compounds; however, more extensive investigations are warranted.

Neolignins Neolignans are a group of dimeric phenylpropanoids that are formed in Myristicaceae and other primitive plant families (e.g. Piperaceae, Eucommiaceae and Lauraceae) by oxidative coupling of allyl and phenyl propanoids (Alves et al. 2002). The biological activities of neolignan derivatives have been investigated against fungi such as *Microsporum canis*, *M. gypseum*, *Trichophyton mentagrophytes*, *T.*

rubrum, and *Epidermophyton floccosum* (Zacchino et al. 1997), bacteria (e.g. *Staphylococcus aureus* and *Bacillus subtilis*) (Pessini et al. 2003), and a panel of cancer cell lines (Siripong et al. 2006). A number of these compounds displayed anti-schistosomiasis activity (Mengarda et al. 2021). Structure–activity relationships of 18 synthetic neolignan derivatives have also been studied (Alves et al. 2002). Two neolignans isolated from the leaves of *Virola surinamensis* (Rol. ex Rottb.) Warb., virolin

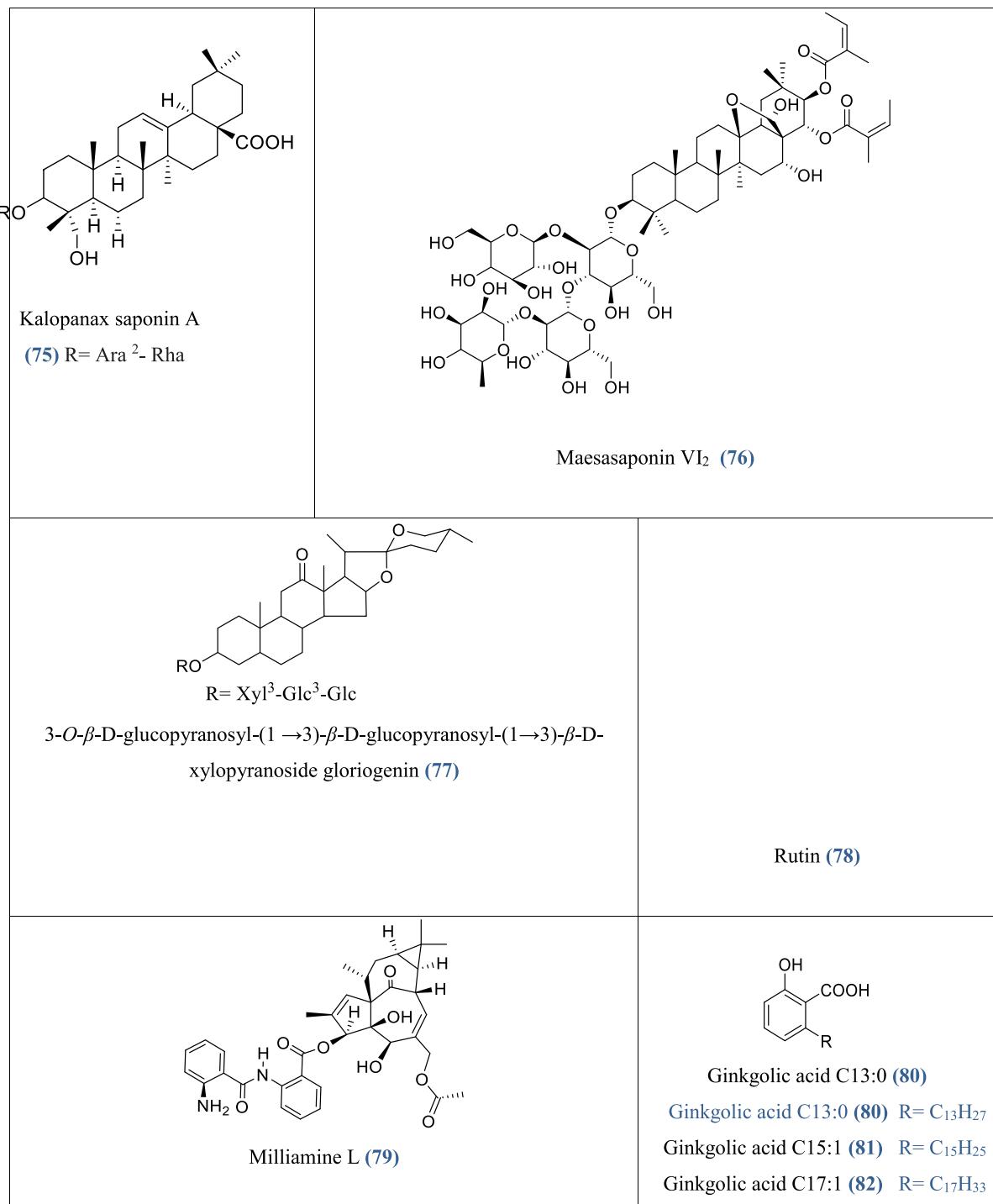


Fig. 6 continued

(Fig. 10) (4) and surinamensin have been active against *S. mansoni* (Alves et al. 1998).

Latex with molluscicidal activity Latex from Euphorbiaceae species has strong molluscicidal activity, particularly against aquatic snails (Bah et al.

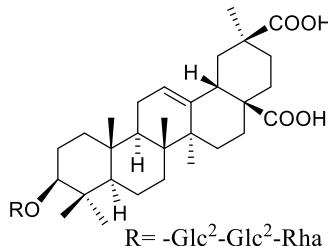
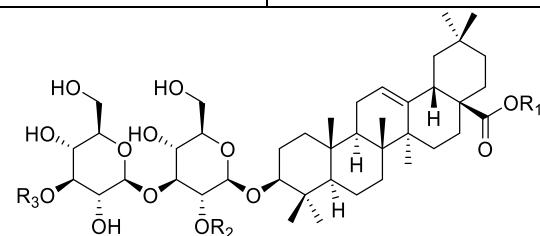
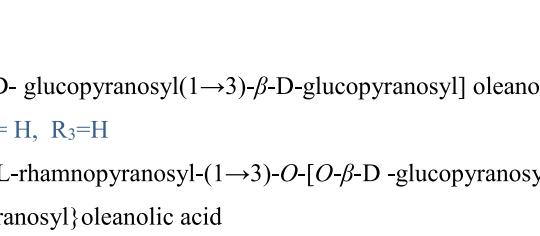
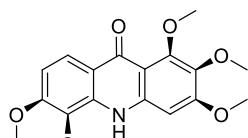
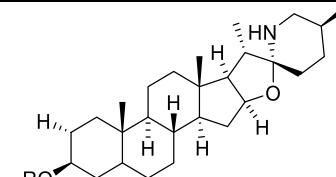
 <p>R= -Glc²-Glc²-Rha</p> <p>3-O-(α-L-rhamnopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl-(1\rightarrow2)-β-D-glucopyranosyl)spergulagenic acid-saponin (83)</p>	 <p>3-O-[O-β-D- glucopyranosyl(1\rightarrow3)-β-D- glucopyranosyl] oleanolic acid (84)</p> <p>R₁=H, R₂=H, R₃=H</p>
	 <p>3-O-{O-α-L-rhamnopyranosyl-(1\rightarrow3)-O-[O-β-D- glucopyranosyl-(1\rightarrow3)]-β-D- glucopyranosyl} oleanolic acid (85) R₁= H, R₂= H, R₃= α-L-Rhap</p> <p>3-O-[2',3'-di- O-(β-D- glucopyranosyl)-β-D- glucopyranosyl]oleanolic acid (86) R₁= H, R₂= β-D-Glup , R₃= H</p> <p>28-O-D- glucopyranosyl- 3- O-{O- α-L-rhamnopyranosyl-(1\rightarrow3)-O-[O-β-D- glucopyranosyl-(1\rightarrow3)]-β- D- glucopyranosyl} oleanolate (87) R₁= β-D-Glup, R₂=H, R₃= α-L-Rhap</p>
 <p>Cusculine (88)</p>	 <p>R= Xyl-2Glc-Gal</p> <p>Tomatine (89)</p>

Fig. 6 continued

2006). The crude latex of *Euphorbia milii* var. *hislopii*, the most powerful molluscicidal agent, has activity against the schistosomiasis-transmitting snails, *B. glabrata* and *B. tenagophila*, (Yadav and Jagannadham 2008). This could be due to the presence of triterpenes, flavonoids, macrolides, ingenol and a phorbol ester (Zani et al. 1993). The latex had stronger effects than the molluscicide niclosamide and was less harmful to nontarget aquatic organisms (Oliveira-Filho et al. 1999; dos

Santos et al. 2007). It also affects the cercariae and larval stage of *Schistosoma* species (De-Carvalho et al. 1998).

Latex of *E. splendens* Bojer ex Hook. is also a potent and specific molluscicide ($LC_{90} < 1.5 \mu\text{g/ml}$) against the vector snails (Schall et al. 1998). It did not show acute toxicity or mutagenic activity towards non-target species at the concentrations of 10–12 $\mu\text{g/ml}$ (Schall et al. 1991). Attempts to identify the components responsible for the effects on snails

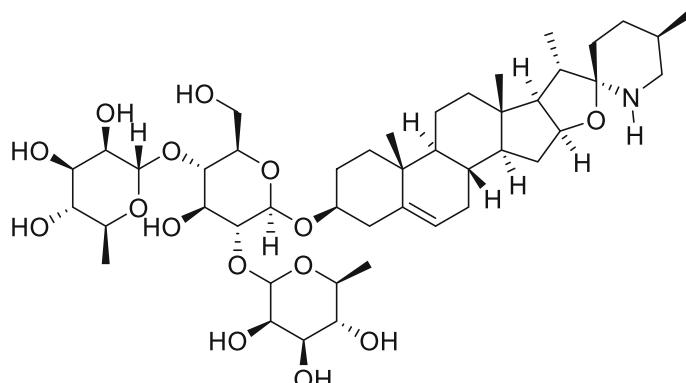


Fig. 7 Solamargine, a glucoalkaloid containing a chacotriose sugar chain moiety with molluscicidal activity (2)

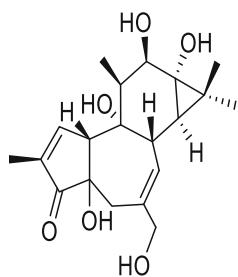


Fig. 8 Chemical structure of phorbol (3)

showed that normal, processed, non-proteinaceous and proteinaceous fractions had molluscicidal activity, but the proteinaceous fraction containing alkaloids had the strongest physiological and lethal effects on fresh water snails (*B. glabarata*) (Yadav and Singh 2011).

Clinical trials for uses medicinal plants against schistosomiasis 70 schistosomiasis haematobium patients both sexes (aged > 15–60 years old) were treated with Mirazid at 10 mg/Kg. The cure rate reached 91.9% after two months and 95.2% on the 3rd post-Mirazid treatment month (El Baz et al. 2003).

In total, 268 school children infected with *S. mansoni* were divided into three groups: PZQ (87), arachidonic acid (ARA, 91), and PZQ plus ARA (90). Over the course of three weeks, and for 15 days, PZQ 40 mg/kg/day, ARA 10 mg/kg/day, or PZQ combined with ARA (40 mg/kg on the first day of treatment, then 15 doses of ARA 10 mg/kg per day for 5 doses/week) were administrated. In children with light and heavy

infections, PZQ and ARA together evoked cure rates of 83% and 78%, respectively. ARA, like PZQ, induced moderate cure rates (50% and 60%, respectively) in school children with light infection and modest cure rates (21% and 20%, respectively) in school children with high infection. Taken together, combination of PZQ and ARA might be useful for treatment of children with schistosomiasis in high-endemicity regions (Barakat et al. 2015).

Conclusion

Schistosomiasis is a major neglected tropical disease with high public health impact. It is difficult to control the disease due to the complex life cycle of the parasite and its wide distribution in tropical and subtropical areas where health and sanitation services are poorly developed. Field control currently relies mostly on some synthetic compounds, which are costly, especially for the low-income regions where the disease is endemic. Since 1981 more than 150 plant species have been tested for molluscicidal activity, and more than 60 natural molluscicidal compounds have been isolated. Preliminary results are encouraging but further investigations are needed to bring more significant progress to a large-scale application in the field, including rigorous pharmaceutical validation, particularly to assess candidate compounds' specificity. More screening of natural sources is also needed, with particular emphasis on structure–activity relationships and action mechanisms. Saponins, for example, specifically block homeostatic circuits.

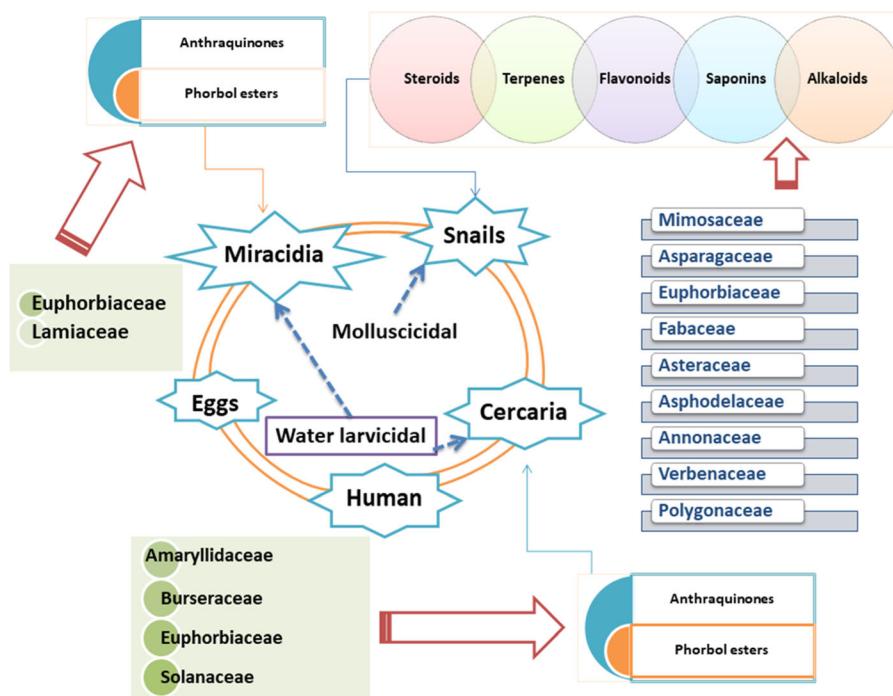


Fig. 9 Lifecycle stages of *Schistosoma* targeted by natural compounds, and the most active plant families

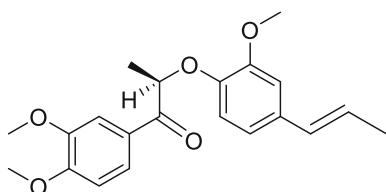


Fig. 10 Chemical structure of virolin (4)

More research is also needed to discover and identify new environmentally friendly molluscicides and larvicides of plant origin that are not toxic to non-target aquatic organisms. Establishment of a global database of the distribution of schistosomiasis vector snails is also necessary for large-scale application of plant molluscicides and optimization of control programs.

The present review here shows that nature offers a plethora of possibilities for improving schistosomiasis control. Production of locally growing plants with molluscicidal and/or anti-schistosome activities would be an attractive alternative, since they could provide abundant agents to interfere with multiple stages of the parasite's lifecycle without imposing excessive

financial burdens. Natural products are likely to be the strongest defense; myriads remain to be discovered.

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Declarations

Conflict of interest The authors declare no conflict of interest.

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