CHAPTER II

HISTORICAL

1. Chemical constituents of genus sapindus.

Distribution of chemical constituents in the genus sapindus.

Table 1 The chemical constituents of Sapindus mukorossi

Chemical compound	Extract	Activity	References
[1]	МеОН	molluscicidal	Huang et al., 2003
[2]	МеОН	Antifungal	Zikova and Krivenchuk, 1965
[3]	МеОН	molluscicidal	Huang et al., 2003
[4]	МеОН	molluscicidal	Huang et al., 2003
[5]	МеОН	molluscicidal	Huang et al., 2003
[6]	МеОН	molluscicidal	Huang et al., 2003
[7]	МеОН	molluscicidal	Huang et al., 2003
[8]	МеОН	molluscicidal	Huang et al., 2003
[9]	МеОН	molluscicidal	Huang et al., 2003
[10]	МеОН	Cytotoxic	Tamura, Mizutani and Yamamoto, 1990

 Table 2 The chemical constituents of Sapindus emarginatus.

Chemical compound	Extract	Activity	References
[1]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001
[2]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001
[6]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001
[7]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001
[8]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001
[9]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001
[11]	МеОН		Kanchanapoom, Kasai and Yamasaki, 2001
[12]	МеОН	-	Kanchanapoom, Kasai and Yamasaki, 2001

Table 3 The chemical constituents of Sapindus trifoliatus

Chemical compound	Extract	Activity	References
[2]	МеОН	Cytotoxic	Tamura, Mizutani and Yamamoto, 1990
[8]	МеОН	Cytotoxic	Tamura, Mizutani and Yamamoto, 1990

Table 4 The chemical constituents of Sapindus saponaria

[8]	МеОН	-	Lemos et al., 1994
[10]	МеОН	-	Lemos et al., 1994
[18]	МеОН	-	Lemos et al., 1994

Table 5 The chemical constituents of Sapindus delavayi

Chemical compound	Extract	Activity	References
[2]	МеОН	-	Nakayama et al., 1986
[8]	МеОН	-	Nakayama et al., 1986
[10]	МеОН	/-	Nakayama et al., 1986
[17]	МеОН	÷	Nakayama et al., 1986
[20]	МеОН	•	Nakayama et al., 1986
[21]	МеОН	-	Nakayama et al., 1986
[22]	МеОН		Wong et al.,1991
[23]	МеОН	-	Wong et al.,1991
[24]	МеОН	-	Wong et al.,1991
[25]	МеОН	-	Wong et al.,1991
[26]	МеОН	-	Wong et al.,1991

δ -Hederin [1]

α-Hederin [2]

Figure 8 Structures of compounds previously isolated from genus Sapindus.

Hederagenin 3-O-(2,4-O-di-acetyl- α -L-arabinopyranosyl-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [3]

Hederagenin 3-O-(3,4-O-di-acetyl- α -L-arabinopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [4]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Hederagenin 3-O-(3-O-acetyl- β -D-xylopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [5]

Hederagenin 3-*O*-(4-*O*-acetyl- β -D-xylopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside

(Mukurozi-saponin E₁) [6]

Figure 8 Structures of compounds previously isolated from genus Sapindus. (Cont.)

Hederagenin 3-O-(3,4-O-di-acetyl- β -D-xylopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [7]

Hederagenin 3-O- β -D-xylopyranosyl- $(1\rightarrow 3)$ - α -L-rhamnopyranosyl- $(1\rightarrow 2)$ - α -L-arabinopyranoside (Sapindoside B) [8]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Hederagenin 3-O-α-L-arabinopyranoside [9]

Clemontoside-C [10]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Hederagenin [11]

Hederagenin 3-*O*-(2-*O*-acetyl- β -D-xylopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [12]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

23-*O*-acetyl-Hederagenin 3-*O*-(4-*O*-acetyl- β -D-xylopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [13]

Oleanolic acid 3-O- β -D-xylopyranosyl- $(1\rightarrow 3)$ - α -L-rhamnopyranosyl- $(1\rightarrow 2)$ - α -L-arabinopyranoside (Prosapogenin CP₃) [14]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Oleanolic acid 3-O-(4-O-acetyl- β -D-xylopyranosyl)-(1 \rightarrow 3)- α -L-rhamnopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranoside [15]

Mukurozioside IIb [16]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Emarginatoside C [17]

Hederagenin 3-O- α -L-rhamnopyranosyl- $(1\rightarrow 3)$ - β -D-glucopyranoside [18]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Hederagenin 3-O- α -L-arabinofuranosyl- $(1\rightarrow 3)$ - α -L-rhamnopyranosyl- $(1\rightarrow 2)$ - α -L-arabinopyranoside [19]

$$H_3$$
C H_3 C H_4 C H_5 C

Pyishauosides IIIa [20]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Pyishauosides IVb [21]

Pyishauosides IVa [22]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Pyishauosides Ib [23]

Pyishauosides IIb [24]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Mukurozioside IIa [25]

Figure 8 Structures of compounds previously isolated from genus Sapindus (Cont.)

Mukurozioside IIb [26]



2. Literature Review

2.1 Literature review on molluscicidal activity of saponins.

Molluscicidal activity has so far only been observed for monodesmosidic saponins [Domon and Hostettmann, 1984], with bidesmosidic saponins requiring basic or enzymatic hydrolysis to the corresponding monodesmosidides before the induction of activity. Saponins of oleanolic acid or hederagenin are the most active, while the corresponding aglycones are inactive [Marston and Hosttetmann, 1985]. Other factors which are important for the activity are: The nature of sugar chains, the sequence of the monosaccharide, and the interglycosidic linkages.

Lauhachinda (1996) reported that neem seed extract at the concentration at 2-3 ppm caused 73 - 100 % death of the tested animals in 72 h. after application. Field test of the neem seed extract on various sizes of snails demonstrated that concentration at 6 ppm can caused 70-80 % death of all size snails at 72 h. [Lauhachinda et al., 1996]

Somkasettrin (1999) reported that neem extracts at concentration 3 ppm caused 100% mortality to small size snails (20-30 mm) at 48 h. while the medium (30 - 40 mm) and large size (50- 60 mm) snails reach 100% mortality at 72 h. [Somkasettrin et al., 1996]

Derris root and tea seed cake extracted by distilled water at 12 h. maceration were tested on three sizes (5-20, 21-35 and 36-50 mm.) of golden apple snail. The LC₅₀ value of derris root extract was 25.20, 45.26 and 77.00 mg/l for three sizes of snail, respectively. The hatching of eggs

showed that there was no effect on the extracts in all concentration [นันที ยา, 2543]

Somrudee (2002) studied the toxicity of indigenous plant extracts to golden apple snail. The result showed that strong activity was observed in the aqueous extracts of *Bougainvillea spectabilis*, *Calotropis gigantean* and *Croton tiglium* and many types of plant in ethanolic extract. [Somrudee, 2002]

Triterpenoid hederagenin saponins isolated from Sapindus mukorossi Garetn. (Sapindaceae) had molluscicidal effects against the golden apple snail, Pomacea canaliculata, which have become major pests of rice and other aquatic crops throughout Taiwan and other parts of Asia [Huang et al., 2003]. Seven isolated hederagenin saponins including one new hederagenin saponins, caused 70 %- 100 % mortality at 10 ppm against the golden apple snails. Hederagenin saponins with three sugar moieties had higher molluscicidal activity than triterpenoid saponins with one sugar moiety.