

Anti-proliferative Activity of *Crotalaria pallida* Aiton on MCF-7 Breast Cancer Cells

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Abstract

Crotalaria pallida Aiton (*C. pallida* Aiton) is empirically used as dietary supplement to treat cancer by the people of North Sulawesi. However, its scientific pharmacology activity has not been explored yet. Therefore, this study was conducted to evaluate anti-proliferative activity of *C. pallida* Aiton on MCF-7 breast cancer cells. The extraction of leaves and seeds were performed using ethanol, ethyl acetate, n-hexane, and water. Phytochemical screening was then performed to identify secondary metabolites in this extract. Anti-proliferative activity was evaluated using 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. The results indicated that ethyl acetate fraction of *C. pallida* Aiton has the lowest IC₅₀ (29,67). In conclusion, ethyl acetate fraction of *C. pallida* Aiton is potential to be developed as anti cancer agent.

Keywords: *Crotalaria pallida* Aiton, WST assay, MCF-7 cell line

Introduction

Breast cancer is the most common life-threatening cancer in women and the leading cause of cancer death among women. Cancer cells are very similar to cells of the organism from which they originated and have similar (but not identical) DNA and RNA.¹⁻³

The toxicity of chemotherapeutic drugs sometimes creates a significant problem in the treatment of cancer using allopathy or established medicine. Various therapies have been propounded for the treatment of cancer, many of which use plant-derived products.⁴⁻⁶

Crotalaria pallida Aiton (*C. pallida* Aiton, *Leguminosae* family, was empirically used by the people of North Sulawesi to treat cancer. However, its scientific pharmacology activity has not been explored yet.

Therefore, this study was conducted to evaluate anti-proliferative activity of *Crotalaria pallida* Aiton on MCF-7 breast cancer cells. Such information is essential to provide evidence on the anti-cancer activity of this medicinal plant from Indonesia.

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Table 1. Result of phytochemical screening

Compound	Sample							
	Seeds Fraction			Leaves Fraction			Seed Extract	Leaves Extract
	n-h	ea	water	n-h	ea	water		
Alkaloid	-	+	+	+	+	+	+	+
Flavonoid	-	+	-	-	-	+	+	-
Saponin	+	+	+	+	+	+	+	+
Triterpenoide	+	+	-	+	+	-	+	-
Steroid	+	+	-	+	+	-	+	-
Fenol	-	+	+	-	+	+	-	+
Monoterpene	-	+	+	-	+	+	+	+
Sesquiterpene	-	+	+	-	+	+	+	-

n-h: n-hexane; ea: ethyl acetate.

Methods

The plant materials used in this study were 1195.45 g and 2449.93 g of *C. pallida* Aiton leaves and seeds, respectively. The solvents used were ethyl acetate, n-hexane, distilled water, fetal bovine serum (FBS) (Gibco), phosphate buffer saline (PBS), water soluble tetrazolium (WST-8) (Dojindo) kit, and sodium dodecyl sulphate (SDS). The cell line used was MCF-7 breast cancer cells (Cell Culture Laboratory of Universitas Padjadjaran).

Cold extraction of leaves and seed of *C. pallida* Aiton was performed using ethanol. for 3 x 24 hours. These extracts were then fractionated using several solvents with different polarity.

Phytochemical screening was performed with several reagents to identify the presence of secondary metabolites such as alkaloid (Mayer-Dragendorff), flavonoid, polyphenol, quinone, tannin, monoterpene, sesquiterpene, and saponin.

Alkaloid Compound

The extract was moistened by adding a

small amount of ammonia. Chloroform. was then added to the solution. The chloroform layer is then removed while being filtered.. Subsequently, hydrochloric acid was added. The mixture was shaken firmly to form several layers. The second layer was added by Mayer's reagent. The presence of white sediment indicated the presence of alkaloid group compounds. The third layer was then added with Dragendorff's reagent. The presence of yellow-orange solution indicated the presence of alkaloid compounds.

Flavonoid Compound

Magnesium powder and hydrochloric acid was added into the extract. The mixture was then heated over a water bath, then filtered. The obtained filtrate was then added with amyl alcohol. The presence of orange, red or yellow solution indicated the presence of flavonoids.

Polyphenol Compound

The extract was heated over a water bath and then filtered. The obtained filtrate was added with iron (III) chloride solution. The presence of the black-blue-green solution indicated the presence of polyphenol compounds.

Table 2. Antiproliferative activity of *C. pallida* Aiton

Sample	Concentration								IC ₅₀ (µg/ml)
	8	16	31	63	125	250	500	1000	
Leaves (ethanol extract)	0	8	33	54	86	85	84	80	56.7
N-hexane fraction	0	6	27	44	53	77	86	84	107.1
Ethyl acetate fraction	0	16	54	79	89	86	83	77	29.67
Distilled water fraction	0	0	0	0	18	65	85	82	>200
Seeds (ethanol extract)	12	18	29	46	28	35	57	84	>200
N-hexane fraction	0	0	49	57	84	89	88	89	35.71
Ethyl acetate fraction	1	22	40	65	52	68	86	88	44.25
Distilled water fraction	0	0	0	0	0	0	0	0	-

Quinone Compound

The extract was heated over a water bath and then filtered. Potassium hydroxide solution. (5%) was added into the solution. The presence of quinone compounds was shown by the formation of yellow to red solution.

Tanin Compound

The extract was heated over a water bath and then filtered. Gelatin solution (1%) was added into the extract. The presence of white sediment indicated the presence of tannin compound.

Monoterpenoid and Sesquiterpenoid

The extract was grinded and evaporated. Dried extract was then added with 10% vanillin solution in concentrated sulfuric acid solution. The color changes indicated the presence of monoterpenoid and sesquiterpenoid.

Triterpenoid and Steroid

The extract was added with ether solution. Approximately 0.5 ml of anhydrous acetic acid, 0.5 ml of chloroform, and 1 ml of concentrated sulphuric acid were added into the solution. The formation of purple-brown ring on the second layer of the

solution indicated the presence of steroid or triterpenoid.

Saponin Compound

The extract was heated over a water bath and then filtered. After the extract was cold, it was shaken rigorously for approximately 30 seconds. Hydrochloric acid. was then added into the foam to examine the presence of saponin compound.

Anti-proliferative test was performed using MTT assay. Various concentrations of extracts were put in a 96 well plate containing MCF-7 cells, and were then incubated for 24 hours. Ten µL of WST-8 solution was added to each well and incubated for 3 hours at 37 °C. The reaction was discontinued with the addition of SDS reagent (100 µl). Cell proliferation rate was determined by reading the microtiter plate and measuring the absorbance at 450 nm.

Results and Discussion

Breast cancer is one of the most prevalent types of malignancy among woman in both developed and developing countries.⁸ Cancer

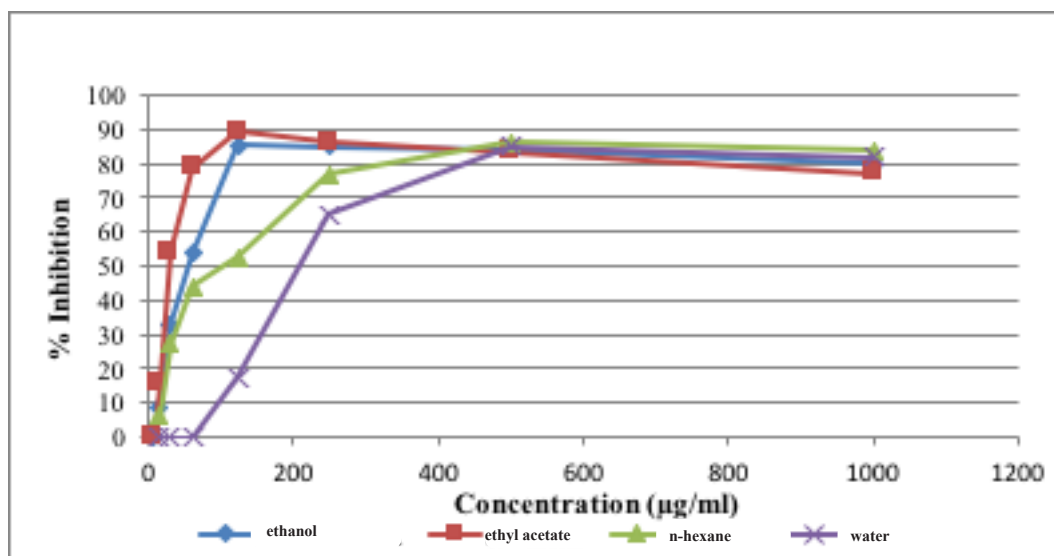


Figure 1. Anti-Proliferative Activity of *C. Pallida* Aiton Leaves

chemotherapy plays a vital role in this disease management. The main objective of such treatment is eliminate the cancer, without causing harm to the host cells. Researchers are predominantly interested in the treatment that are selective and can possibly induce cellular apoptosis, activity that is satisfied by several secondary metabolites of the plant.⁹

The yields of concentrated extracts of *C. pallida* Aiton leaves and seeds were 57 g and 42 g, respectively. In this study, ethanol was used since it is a universal solvent which attract polar, semi polar and non polar secondary metabolites.⁷ The results of phytochemical screening can be found in Table 1.

The WST assay results showed that the extract and fraction of *C. pallida* Aiton had anti-proliferative activity. The lowest IC₅₀ was observed in ethyl acetate leaves fraction (29.67 µg/ml) and n-hexane seeds fraction (35.41 µg/ml) (Table 2).

The presence of several phytochemicals may contribute to its anti-cancer activities, such as alkaloid, flavonoid, and terpenoid. Previous studies showed that alkaloid might modulate microtubule activity. Microtubules contribute in the formation of mitotic spindle, making it becomes suitable target for anti-cancer drugs. Vinblastine is one type of the alkaloid compound which has been isolated for anti-cancer drug development.⁸⁻¹¹

Flavonoids might also induce anti-proliferative activity of *C. pallida* extract. Previous study showed that it might interferes with multiple mechanism, including signal transduction. Several flavonoid compounds could also activate apoptotic proteins and induce DNA damage in cancer cells.¹²

Conclusion

Ethyl acetate fraction of *C. pallida* Aiton is potential to be developed as anti cancer agent.

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Conflict of Interest

None declared.

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