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Anti-cancer Constituents of Liliaceae Plants

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INTRODUCTION

Liliaceac is one of the laegest families of flowering plants, consisting of about 220-240 genera with 3500-4000 species. According to A. Engler's classification(1964)¹, it is divided into 13 subfamilies(Table 1).

Some Liliaceae plants are well known as important sources of medicines. For example, Rhizoma Anemarrhenae(rhizome of *Anemarrhena asphodeloides*), Radix Ophiopognis(tuber of *Ophiopodon japonicus*) and Rhizoma Smilacis Glabrae(rhizome of *Smilax glabra*) are listed in Japanese Pharmacopoeia. Sap of *Aloe* spp is used for

a laxative, and colchicine, a medicine against gout, is obtained from *Colchicum autumnale*.

Liliaceae is a rich source of steroidal glycosides, and some plants, such as *Urginea scilla*, *Convallaria* spp., *Rhodea japonica*, etc. contain cardiac glycosides, but no systematic study of the chemical constituents of Liliaceae plants had been carried out until we started our research.

There are two procedures for isolating compounds with medicinal potential from plants(Chart 1). For crude drugs and medicinal plants possessing potent activity, the method indicated on the left in Chart 1 is recommended, but in the case of the plants whose

Table 1. Subfamilies and main genera of Liliaceae by Engler(1964)

Alstromerioideae: Alstromeria

Allioideae: Agapanthus, Allium, Brodiaea, Ipheion, Triteleia

Asparagoideae: Asparagus, Convallaria, Paris, Polygonatum, Rhodea, Ruscus, Trillium

Asphodelideae: Aloe, Chlorophytum, Hemerocallis, Hosta

Aaletroideae: Aletris

Lilioideae: Erithronium, Fritillaria, Lilium, Nomocaris, Tulipa

Herrerioideae: Herreria Luzuriagoideae: Luzuriaga

Melanthiodeae: Narthecium, Tricyrtis, Veratrum, Zygadenum

Ophiopogoniodeae: Liriope, Ophiopogon

Scilloideae: Camassia, Chionodoxa, Eucomis, Hyacinthus, Ornithogalum, Scilla, Urginea

Smilacoldeae: Smillax

Wurmbaeoldeae: Colchicum, Gloriosa

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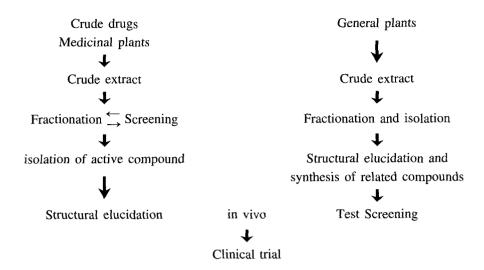


Chart 1. Procedure to identify naturally occurring compounds with medicinal potential

medical properties are undocumented, the right-hand method is suitable, Using this method, about 500 new steroidal glycosides including cholestane glycosides and steroidal alkaloids were isolated from about 50 species of Liliaceae²).

The biological activities of the compounds isolated were examined using several methods²⁾. For anticancer activities, the following two tests were carried Out:

1. in vitro Anti tumor promoter activity test

 Anti-tumor activity test:cytotoxicity towards HL-60 and MOLT-4 leukemia cells

1. In vitro Anti-tumor Promoter Activity

Some foods originating from Liliaceae plants, such as Allium sativum(garlic) and A. wakegi are effective in the prevention of mailgnant tumors. Sterol, laxogenin, was identified as the active compound³⁾. in vitro Antitumor promoter test of the steroidal glycosides isolated by our studies was carried out, and the

activity was estimated from the rate of incorporation of phospholipid into HeLa cells stimulated with TPA (12-O-teradecanoylphprbol 13-acetate).

The steroids(1-3) isolated from *Hosta longipes* have a carbonyl group as well as the laxogenin. They showed relatively strong activities.

A sterroidal glycoside(4) bearing HMG methyl ester at C-27 hydroxyl position⁴⁾ isolated from the bulls of *Lilium* spp. showed prominent activity which was ten times higher than that of the laxogenin. It also showed comparative atrength in inhibiting the

Table 2. Inhibition of human malignant tumor cell proliferation with 4

cells	inhibition(%)
pancreatic cancer(PANC-1)	95.5
Osteosarcoma (OST)	72.2
Human gastric cancer(HGC-27)	70.8
Pheochromcytoma(PC-12)	24.8

4: IC₅₀ 4µg/ml(3.8µM) Sample concentration: 5µg/ml

Table 3. Cytostatic activities of 5 and 6 towards HL-60 and MOLT-6 cells

	· ·	
	HL-60	MOLT-4
5	0.0092	0.0024
6	0.025	0.018
etoposide	0.025	0.018
adriamycin	0.0072	0.035
vincristine	0.0029	0.00059
methotrexate	0.012	0.048 $IC_{50}(\mu M)$

proliferation of several human malignant tum or cells⁵⁾ (Table 2).

2. Anti-tumor Activity

Table 4. Cytostatic activities of and some anticaner agents towards HL-60 and MOLT-4 cells

	HL-60	MOLT-4
7	0.00025	0.00049
etoposide	0.025	0.054
adriamycin	0.0072	0.035
vincristine	0.0029	0.00059
methotrexate	0.012	$0.048 IC_{50}(\mu M)$

4-O-p-methoxybenzoyl-Rha-(1→2)-Glc

The seres of cholestane glycosides isolate from *Ornithogalum saundersiae*, a bulbous plant native to South Africa, exhibited very strong cytostatic activity towards several malignant tumor cells. The activities of two rearranged cholestane glycosides(5⁶), 6⁷) against human leukemia cells, HL-60 and MOLT-4 were almost equal or more potent than those of the clinically applied anti-cancer agents, etoposide, methotrexate, adriamycin(ADM) and vincristine⁸(Table 3).

Table 5. Cytostatic activities of 7 and clinically applied anticancer anents on various malignant tumor cells

				<u> </u>		
IC ₅₀ (μM)						
7	MMC	ADM	CDDP	CPT	TAX	
1.5	2.0	2.0	10	2	2	
0.00013	0.01	0.003	0.05	0.005	0.01	
0.00077						
0.00010						
0.00016						
0.00068						
0.00020						
0.00020	0.01	0.002	0.001	0.001	0.002	
0.00026						
0.00016	0.02	0.01	0.005	0.005	0.001	
	1.5 0.00013 0.00077 0.00010 0.00016 0.00068 0.00020 0.00020	7 MMC 1.5 2.0 0.00013 0.01 0.00077 0.00010 0.00016 0.00068 0.00020 0.00020 0.00026	7 MMC ADM 1.5 2.0 2.0 0.00013 0.01 0.003 0.00077 0.00010 0.00016 0.00068 0.00020 0.00020 0.00026	IC ₅₀ (μM) 7 MMC ADM CDDP 1.5 2.0 2.0 10 0.00013 0.01 0.003 0.05 0.00077 0.00010 0.00016 0.00068 0.00020 0.00020 0.00026	IC ₅₀ (μM) 7 MMC ADM CDDP CPT 1.5 2.0 2.0 10 2 0.00013 0.01 0.003 0.05 0.005 0.00077 0.00010 0.00016 0.00068 0.00020 0.00020 0.00026	IC ₅₀ (μM) 7 MMC ADM CDDP CPT TAX 1.5 2.0 2.0 10 2 2 0.00013 0.01 0.003 0.05 0.005 0.01 0.00077 0.00010 0.00016 0.00068 0.00020 0.00020 0.00026

CCD-CCD-19Lu(Human normal pulmonary cell)

P388/ADM(adriamycin-resistant P388)

FM3A(mouse mastrocarcinoma)

Lu-65(human pulmonary large cell carcinnoma)

RERF-LC-AI(human pulmonary squamous cell carcinoma)

CCRF-CEM(human leukemia)

The cytostalic activities of the cholestane glycosides partly resulted from the induction of apoptosis. The apoptosis is caused by the activation of calcium dependent endonuclease. An analysis of flow cytometry showed that glycosides stopped the cell cycle of HL-60 at S2/M phase and induced apoptosis at G0/G1 phase⁹⁾.

The cholestane glycoside(7), the main constituent of the bulls of *O. saundersiae*, showed a remarkbly strong cytostatic activity, about 10-100 times stronger

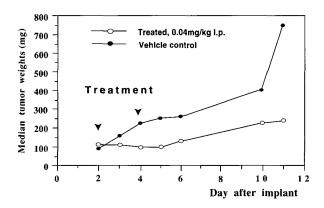


Fig. 1. Response of HEP 134 xenografts to 7

than those of mitomycin C(MMC), ADM, cisplatin(CDPP), camptpthecin(CPT) and taxol(TAX)¹⁰ (Table 4).

P388/CPT(camptothecin-resistant P388)

A-549(human pulmonary adenocarcinoma)

Lu-99(human pulmonary large cell carcinnoma)

P388(mouse leukemia)

It is worth noting that the activity of 7 towards tumor cells including carcinostatic-resistant cells is about 10,000 times more potent than that towards normal cells(CCD-19Lu)¹⁰⁾ (Table V).

The data of the 60 Cell Line Assay by National Cancer Institute(NCI) in America indicated that 7 has a wide anti-tumor spectrum and is especially effective in proliferation of melanoma cells.

In *in vivo* test, 7 increased the life span of mice with P388 leukemia by 59% with one-time administration of 0.01mg/kg¹⁰. 7 also inhibited the proliferation of human hepatoma 134 transplanted in nude mice at 0.06mg/kg¹¹ (iv) (Fig. 1).

Thus, the compounds isolated from *Ornithogalum* saundersiae have extremely potent cytostatic activity, and some other steroidal glycosides isolated from Liliaceae plants also exhibited strong cytostatic activities against several malignant tumor cells. Recently a steroidal glycosidc(8) was isolated from the bulb of *Allium*

jesdianum. 8 exhibited considerable cytostatic activity toward HL-60 cells with an GI₅₀ value of $1.5 \mu g/ml$ compared with etoposide used as positive control(GI₅₀ $0.3 \mu g/ml$). The evaluation of 8 by the National Cancer Institute 60 cell line assay showed that the mean concentrations requried to achieve GI₅₀, TGI and LC₅₀ levels against the panel of cells tested were $4.5 \mu M$ $18 \mu M$ and $54 \mu M$, respectively, and some cell lines, such as leukemia CCRF-CEM, non-small cell lung cancer HOP-62 and breast cancer MCF7 were affected by 8^{12} .

The compounds isolated from plants have attracted special attention as sources for medicines. Our studies clarify that the stcroidalgycosides isolated from Liliaceae plants are important compounds for of anti-cancer agents.

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