NOTE

Free Radical Scavenging Activities of β -Sitosterol Extracted from *Abies yuanbaoshanensis*

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From *Abies yuanbaoshanensis*, β -sitosterol was isolated and purified by column chromatography. The free radical scavenging activities of β -sitosterol were evaluated against DPPH, ABTS and hydroxyl radicals, respectively. The results showed that all IC₅₀ of β -sitosterol were much lower than the standard value 10 mg/mL, demonstrating good radical scavenging activities of β -sitosterol.

Key Words: *Abies yuanbaoshanensis*, β -Sitosterol, Free radical scavenging activity.

Free radical scavengers or antioxidants are currently forged as the drug candidates to counter many diseases, such as drug-associated toxicity, inflammation, atherogenesis and aging in aerobic organisms¹⁻³. Thus the significance of free radicals in the pathogenesis of various diseases has attracted considerable interest. Minor dietary compositions have been considerably deemed to combat the ill effects of free radicals. In addition, since that what may be learn from natural plants could overcome the toxicity problem of synthetic radical scavengers or antioxidants, so isolation of natural radical scavengers or antioxidants from natural plants has become a popular method.

Abies yuanbaoshanensis, which distributed in Rongshui county in Guangxi Province of China, is a peculiar endangered precious plant. The pharmacological research has showed that it possess of wide biological activities⁴, including antitumor, antibacterial, antifungal, antiinflammatory and antitussive activities. Previous studies have indicated that various diseases such as antitumor, antibacterial and antifungal are characteristically associated with free radicals and ROS⁵. It is possible that some constitutes of *Abies yuanbaoshanensis* should get good free radical scavenging activities. So herein the extraction and radical-scavenging activities of *Abies yuanbaoshanensis* was studied.

Abies yuanbaoshanensis were collected from Rongshui county of Guangxi Zhuang Autonomous Region in China, in July 2009 and verified by Prof. Zong-Ming Su. The air-dried needles and stems of *Abies yuanbaoshanensis* (10 Kg) was extracted with 95 % ethanol (50 L $\times$ 3, each 7 days) at room

temperature and the combined extracts were concentrated in vacuum to give a residue (450 g), which was suspended in the water and re-extracted with petroleum ether, ethyl acetate successively to get three fractions. The petroleum ether extract (150 g) was subjected to column chromatography on silica-gel eluting with petroleum ether/ethyl acetate (100:0-50:50, v/v) to give 9 fractions. The petroleum ether/ethyl acetate 10:1 fraction (8 g) was further purified by silica-gel column chromatography developing with petroleum ether/CHCl₃/acetone (70:18:1) and a white needle crystals of β -sitosterol (960 mg) was obtained. The flow diagram was shown in Fig. 1. m.p. 135-137 °C. ¹H NMR (500 MHz, CDCl₃): δ 5.35 (1H, br, d, J = 4.8 Hz, H-6), 3.50 (1H, m, H-3), 1.01 (3H, s, CH₃-19), 0.94 (3H, d, J = 8.0 Hz, CH₃-21), 0.85 (3H, d, J = 8.0 Hz, CH₃-29), 0.82 (3H, d, J = 7.6 Hz, CH₃-26), 0.80 (3H, d, J = 7.6 Hz, CH₃-27), 0.67 (3H, s, CH₃-18). ¹³C NMR (125 MHz, CDCl₃): δ 37.2 (C-1), 31.6 (C-2), 71.6 (C-3), 42.3 (C-4), 140.7 (C-5), 121.7 (C-6), 31.7 (C-7), 31.9 (C-8), 50.1 (C-9), 36.5 (C-10), 21.1 (C-11), 39.6 (C-12), 42.3 (C-13), 56.6 (C-14), 24.3 (C-15), 28.2 (C-16), 56.2 (C-17), 11.8 (C-18), 19.8 (C-19), 36.4 (C-20), 18.8 (C-21), 33.7 (C-22), 26.0 (C-23), 45.4 (C-24), 29.1 (C-25), 19.5 (C-26), 19.2 (C-27), 23.0 (C-28), 12.0 (C-29). The above data were consistent with β -sitosterol (Fig. 2) according with literature⁶.

In vitro antioxidant activities were measured against DPPH⁶, ABTS⁷ and hydroxyl radicals⁸, respectively, according to the literatures⁶⁻⁹ with a little modification. The values of IC₅₀, the effective concentration at which 50 % of the radicals were scavenged, were calculated to evaluate the antioxidant

Fig. 1. Flow diagram of the optimal process

Fig. 2. Structure of β -sitosterol

activities. A lower IC_{50} value indicated greater antioxidant activity. IC_{50} values of lower than 10 mg/mL usually implied effective activities in antioxidant properties⁶. The tested results were shown in Fig. 3 (Table-1).

Fig. 3. Radical scavenging activities of β -sitosterol from *Abies yuanbaoshanensis*. Values are means $\pm$ SD of three determinations. $p < 0.05$, when compared with control

TABLE-1			
IC_{50} (mg/mL) OF THE β -SITOSTEROL FROM <i>Abies yuanbaoshanensis</i>			
IC_{50} (mg/mL)	DPPH [•]	ABTS ^{•+}	OH [•]
β -Sitosterol	6.298	0.629	0.03

DPPH radical scavenging activity evaluation, a classical assay in radical scavenging activity studies, could offer rapid techniques for screening the radical scavenging activity of the antioxidants or radical scavenger. As showed in Fig. 3 (Table-1), IC_{50} of β -sitosterol in DPPH assay was found to be 6.298 mg/mL, which was evidently lower than the standard value 10 mg/mL⁶, indicating that β -sitosterol has good potent inhibition of DPPH radical.

ABTS^{•+} radical evaluation assay is another conventional and simple tool for determining the radical activity of hydrogen-donating and chain breaking radical scavengers⁷. As can be seen in Fig. 3 (Table-1), IC_{50} of β -sitosterol on ABTS^{•+} radical was found to be 0.629 mg/mL and much lower than 10 mg/mL⁶, demonstrating its good potent inhibition of ABTS^{•+} radical.

The radical scavenging activities were also tested in the present study using hydroxyl radicals generated by Fenton reagent⁹. As shown in Fig. 3 (Table-1), β -sitosterol exhibited good activity in an amount dependent manner and its IC_{50} was found to be 0.03 mg/mL, which was much lower than 10 mg/mL⁶, indicating its good radical scavenging activities on hydroxyl radicals.

Conclusion

β -Sitosterol was extracted and isolated from *Abies yuanbaoshanensis* and it was found to get good radical scavenging activity on DPPH, ABTS^{•+} and hydroxyl radicals.

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