

## INFORMATION ABOUT NEW CONTRIBUTIONS OF THE THESIS

**Full name of the PhD student:** Nguyen Viet Dung

**Thesis name:** “Research on plant characteristics, chemical compositions and some biological effects of *Hypericum sampsonii* Hance., (Hypericaceae) family”.

**Major:** Medicinal herbs - Traditional medicine

**Code:** 9720206

**Full name of the instructor** (Academic title, degree):

1. Assoc. Prof. PhD. Nguyen Duy Thuan

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**Training institution:** National Institute of Medicinal Materials

**Summary of new results of the thesis:**

### ***1. In term of botany***

This thesis is the first document in Vietnam describing in detail morphological characteristics of plants, characteristics of micro-dissection of the stems, leaves, roots and powder of the stems, leaves and roots of *Hypericum sampsonii* Hance., (Hypericaceae) family.

### ***2. In term of chemistry***

From the aboveground part, *Hypericum sampsonii* Hance., (Hypericaceae) family is isolated with 15 compounds. Their chemical structures are determined by HR-ESI-MS, 1D-, 2D NMR spectroscopy.

Including:

- 5 compounds are firstly isolated from *Hypericum sampsonii* Hance: 3,5,6-trihydroxy-1-methoxyxanthone; Petiolin F; Quercetin-3'-O- $\beta$ -D-galactopyranosid; Quercetin-3-O- $\beta$ -D-galactopyranosid; Cratoxyarborenone F.

- The remaining 10 compounds include (Mangiferin, Quercetin, 3,5-dihydroxy-2',4',6'-trimethoxybenzophenone-3-O- $\alpha$ -L-rhamnopyranosid, Neolancerin, Euxanthone, 2-hydroxyxanthone, acid betulinic, 3,5-dihydroxy-2',4',6'-trimethoxybenzophenone, I3-II8-biapigenin và Daucosterol).

### ***3. In term of acute toxicity and biological activity***

**3.1. Acute toxicity:** The aboveground part of *Hypericum sampsonii* Hance leaves has been evaluated to be at very low toxicity in 24 hours at a dose of 225.0 grams of dried medicinal herbs/Kg of body weight.

### ***3.2. Biological activity***

- The antioxidant activity of isolated compounds and extracts of *Hypericum sampsonii* Hance leaves on DPPH-free radical scavenging model shows that Mangiferin (HSA1), 3,5,6-trihydroxy-1-methoxyxanthone (HSA2) and the extract of *Hypericum sampsonii* Hance has a low antioxidant activity, in which compound 3,5,6-trihydroxy-1-methoxyxanthone (HSA2) is a xanthone which is firstly isolated from *Hypericum sampsonii* Hance; furthermore, and this is also the first time on which the antioxidant activity of this compound has been mentioned.

- It evaluates the ability to inhibit NO production for the isolated substances, resulting compounds Petiolin F (HSA4), 3,5-dihydroxy-2',4',6'-trimethoxybenzophenone-3-O- $\alpha$ -L-rhamnopyranosid (HSA9) and 3,5-dihydroxy-2',4',6'-trimethoxybenzophenone (HSA20); in which, Petiolin F (HSA4) is the first compound isolated from of *Hypericum sampsonii* Hance and this is also the first time that the inhibitory activity of NO production of these compounds has been mentioned.

- The acetylcholinesterase enzyme inhibitory activity is evaluated for 9 samples of isolates (Mangiferin; 3,5,6-trihydroxy-1-methoxyxanthone; Petiolin F; 3,5-dihydroxy-2',4',6'-trimethoxybenzophenone-3-O- $\alpha$ -L-rhamnopyranosid; Neolancerin; Cratoxyarborenone F; Euxanthone; 2-hydroxyxanthone and 3,5-dihydroxy-2',4',6'-trimethoxybenzophenone) and extract of *Hypericum sampsonii* Hance. The result shows that the extract of *Hypericum sampsonii* Hance has a very good inhibitory activity of acetyl cholinesterase enzyme with value of  $IC_{50} = 19,95 \pm 1,09 \mu\text{g/ml}$  at 2 test concentrations of 100  $\mu\text{g/ml}$  and 500  $\mu\text{g/ml}$  compared with HSA15; in addition, in 9 samples of compounds, Cratoxyarborenone F (HSA15) is firstly mentioned with the ability to inhibit acetyl cholinesterase enzyme.

- The aboveground extract of *Hypericum sampsonii* Hance has been highly evaluated to have a hepatoprotective antioxidant effect equivalent to and slightly better than Sylymarin at a dose of 140 mg/kg with two dosage levels (3.6 grams of DL/kg and 10.8 grams of DL/kg).

- The aboveground extract of *Hypericum sampsonii* Hance has been highly evaluated to improve memory. The results at a dose of 10.8 grams of DL/kg of body weight of rats have an effect on improving memory equivalent to Scopolamin control.

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