## Anti-inflammatory Effect of Hederagenin Glycoside Isolated from Lonicera japonica

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Lonicera japonica Thunb. is a twining shrub that has been used as an antidote and to treat urinary disorders, fever and headache. It has been known as an anti-inflammatory agent in Korea from ancient times and is used widely for treating upper-respiratory tract infections, diabetes mellitus and rheumatoid arthritis. In the previous research, we isolated several flavonoid derivatives from the EtOAc soluble fraction. Among the flavonoid derivatives, ochnaflavone, a biflavone, exhibit strong inhibitory activity against group II PLA2 enzyme purified from rat platelet ( $IC_{50} = 3\mu M$ ). In the continuing investigations into anti-inflammatory agents based on this plant extract, we identified that the n-butanol soluble fraction showed antiinflammatory activity against acute, granulomatic and chronic inflammation models in mice and rats. From the n-butanol soluble fraction, we isolated three new hederagenin glycosides and elucidated their structures as 3-0-a-L-arabinopyranosyl hederagenin 28-0-a-L-rhamnopyranosyl  $(1\rightarrow 2)$ - $[\beta$ -D-xylopyranosyl $(1\rightarrow 6)$ ]- $\beta$ -D-glucopyranosyl · ester (loniceroside A), 3-0-a-Lrhamnopyranosyl  $(1\rightarrow 2)$ -a-L-arabinopyranosyl hederagenin 28-0-a-L-rhamnopyranosyl  $(1\rightarrow 2)$ - $[\beta-D-xy]$ lopyranosyl $(1\rightarrow 6)$ ]- $\beta-D$ -glucopyranosyl ester (loniceroside B) and 3-0- $\beta$ -Dglucopyranosyl hederagenin 28-0- $\alpha$ -L-rhamnopyranosyl (1 $\rightarrow$ 2)-[ $\beta$ -D-xylopyranosyl (1 $\rightarrow$ 6)]- $\beta$ -D-glucopyranosyl ester (loniceroside C). Lonicerosides A and C showed anti-inflammatory activities against croton-oil and arachidonic acid induced mouse ear edema models at the dose of 100 mg/kg. Furthermore, loniceroside A, a major component of the n-butanol fraction, reduced adjuvant-induced arthritis in rats at the dose of 100mg/kg/day.

## References

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