

- 品檢驗局中程綱要研究計畫報告。
2. Koide, T., Iwata, M., Maekawa, K., Saito, H., Tanimoto, T., Okada, S., Nakane, T., Kawahara, N., Sekita, S., Satake, M., Yokota, Y., Tsuno, T., Suzuki, H., Matano, Y. and Yamamoto, K. 2002. Qualitative evaluation for the establishment of NIHS puerarin reference standard. *Iyakuhin Kenkyu*, 33(2): 118-123.
 3. 行政院衛生署中華藥典編修委員會。2006。中華藥典。第六版。行政院衛生署，台北。
 4. 日本藥局方編輯委員會。2006。第十五改正日本藥局方。pp. 1190。廣州書局，東京。
 5. 原田正敏。1989。繁用生藥之成分定量。廣川書店，東京。

Qualitative Evaluation for the Preparation of Chinese Medicine Reference Standard-Puerarin

YA-HUI HSU, SZU-HUI CHEN, FANG-SU LIU, YI-CHU LIU,
JER-HUEI LIN AND CHI-FANG LO

Division of Research and Analysis

ABSTRACT

The raw material of Puerarin was examined prior to the preparation of the "Puerarin Reference Standard". The physico-chemical properties of the candidate material were evaluated by a collaborative study among seven laboratories. Analytical data obtained were summarized as follows: the UV maximum absorption wavelengths and the corresponding specific absorbencies ($E_{1\text{cm}}^{1\%}$) were 249.9, 305.7 nm, and 736.8, 248.8, respectively. IR Spectra showed that IR absorption at 3366, 3234, 1633, 1515 and 1059 cm^{-1} . HPLC analysis revealed 1-6 impurities which individual amount was $\leq 0.50\%$ and the total amount from any single laboratory was $\leq 0.84\%$.

Based on the above results, the candidate material met the requirement of authorization as the Puerarin Reference Standard.

Key words: puerarin, Chinese medicine reference standard, collaborative study