

Synthesis of Ermanin, 5,7-Dihydroxy-3,4'-dimethoxyflavone from Kaempferol, 3,5,7,4'-Tetrahydroxyflavone with Two *O*-Methyltransferases Expressed in *E. coli*

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Received December 9, 2005

Key Words : Ermanin, Kaempferol, *O*-Methylation

Enzymatic modifications of natural compounds have been drawn attention because they provide regioselectivity and chiral selectivity, which are hardly achievable with chemical synthesis.^{1,2} Using a mobilized enzyme or a whole cell containing a particular gene delivers useful tools for modification of natural compounds. In addition, using a whole cell, known as biotransformation, has advantage of saving expensive cofactors.

Flavonoids and alkaloids are secondary metabolites produced mainly in plants.³ Their biological activities have an impact on human health⁴ so that they serve as target molecules to develop new drugs. One of common modification reactions found in these two groups of compounds is *O*-methylation due to free hydroxyl groups. Various *O*-methylation reactions have been reported and a number of genes (*O*-methyltransferases: OMTs) that mediated *O*-methylation have been cloned and characterized.⁵ Some of OMTs that use flavonoids as substrates are highly specific and thus they could be used for the regioselective modification of flavonoids. *O*-methylation of flavonoids resulted in reduction of chemical reactivity and increase of antimicrobial activity.⁵ So far, *O*-methylated flavonoids have been mainly isolated from plants,⁶ which could be a limitation for biological assays. Kaempferol (3,5,7,4'-tetrahydroxyflavone) is one of commonly found flavonoids in nature. Even though several biological activities of kaempferol have been established, its dimethoxy form, ermanin (5,7-dihydroxy-3,4'-dimethoxyflavone) was known to have several activities including antiviral,⁷ anti-inflammatory,⁸ cytotoxic,⁹ and antibacterial activity.⁶ Lists of its biological activity might be extended if large amount of ermanin is supplied.

Previously, we cloned and characterized two OMT genes, *SOMT-2* (soybean *O*-methyltransferase-2)¹⁰ and *ROMT-9* (rice *O*-methyltransferase-9).¹¹ The transgenic *E. coli* expressing *SOMT-2* transferred a methyl group to 4'-hydroxyl group of flavonoids. *ROMT-9* expressed in *E. coli* showed different regioselectivity depending on the availability of 3'-hydroxyl groups. It transferred a methyl group to 3'-hydroxyl group if flavonoids have 3'-hydroxyl group. But, when 3'-hydroxyl group is not present, it methylated a 3-hydroxyl group. Here we report the biological synthesis of ermanin from kaempferol with two OMTs expressed in *E. coli*.

ROMT-9 and *SOMT-2* were cloned in one expression

vector¹² and both proteins were induced from *E. coli* containing both *ROMT-9* and *SOMT-2* by adding IPTG at 100 μ M. In addition, *ROMT-9* and *SOMT-2* were induced separately from *E. coli* containing either *ROMT-9* or *SOMT-2*. After 4 hr induction, the cells from each culture were harvested and resuspended in LB containing ampicillin (50 μ g/mL). 100 μ M of kaempferol (3,5,7,4'-tetrahydroxyflavone) was added. The mixture was further incubated for 15 hrs at 28 °C. Analysis of culture filtrates from three reactions using high performance liquid chromatography (HPLC)⁹ revealed that kaempferol was converted into a new product which had different HPLC retention time from kaempferol itself. *ROMT-9* produced a new peak at 15.2 min (Fig. 1B) and *SOMT-2* produced a new peak at 19.9 min

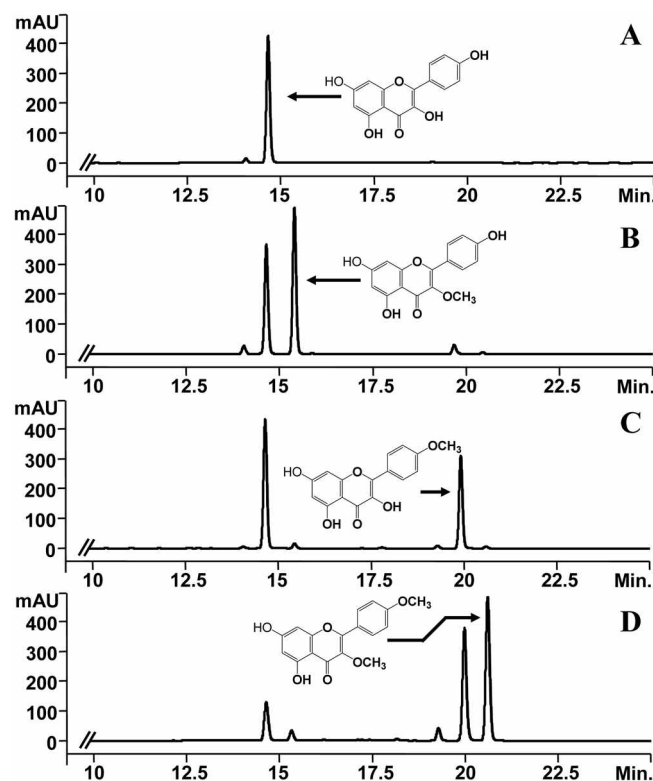
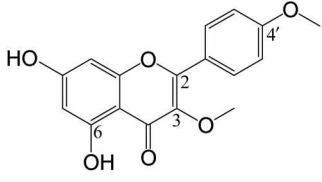


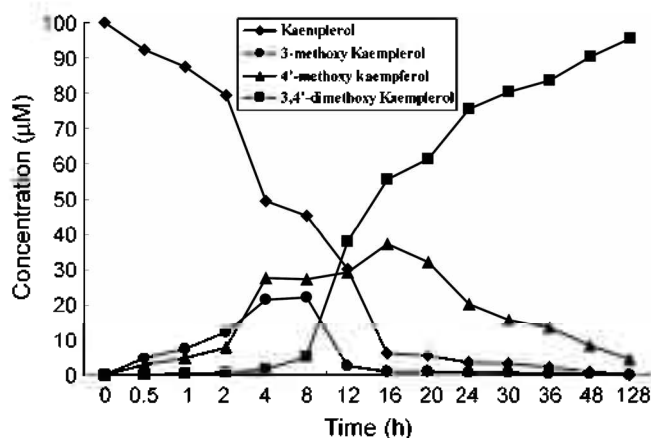
Figure 1. HPLC analysis of kaempferol reaction products with *SOMT-2* and *ROMT-9*. A, authentic kaempferol; B, reaction product of kaempferol with *ROMT-9*; C, reaction product of kaempferol with *SOMT-2*; D, reaction product of kaempferol with *SOMT-2* and *ROMT-9*

Table 1. The assignments of the NMR data of the metabolite of kaempferol produced by SOMT-2 and ROMT-9


position	δ of ^1H	δ of ^{13}C	HMBC	NOESY
2	—	155.0	—	—
3	—	139.1	—	—
4	—	177.8	—	—
5	—	161.1	—	—
6	6.22 (d; 2.0)	98.3	C-5, C-7, C-8, C-10	—
7	—	164.2	—	—
8	6.45 (d; 2.0)	93.6	C-6, C-7, C-11	—
9	—	156.3	—	—
10	—	104.3	—	—
3-OCH ₃	3.76 (s)	59.6	C-3	H-2'
1'	—	113.8	—	—
2'	8.02 (d; 9.0)	129.9	C-2, C-4'	H-3',4'-OCH ₃
3'	7.13 (d; 9.0)	122.1	C-1'	H-2',3'-OCH ₃
4'	—	161.2	—	—
4'-OCH ₃	3.84 (s)	55.3	C-4'	H-3'

(Fig. 1C) while two reaction products appeared from the reaction with *E. coli* expressing both ROMT-9 and SOMT-2 (Fig. 1D). One of them had the same retention time with the reaction product of ROMT-9. The peak at 20.6 min was likely to be a dimethylated product. The structure of this compound was clarified by NMR experiments as described in Kim *et al.*¹⁰ Each eluent containing the metabolite of kaempferol produced by SOMT-2 and ROMT-9 was collected twenty times on HPLC and evaporated under reduced pressure. The dried remnant was dissolved in ethylacetate, and the supernatant was separated by centrifuge and evaporated again under reduced pressure. The final remnant was dissolved in dimethylsulfoxide- d_6 for the NMR experiments such as ^1H , ^{13}C NMR, HMQC, HMBC, and NOESY. The complete assignment of the NMR data of the reaction product is listed in Table 1. Two methoxy peaks were observed at both the ^1H NMR spectrum and the ^{13}C NMR spectrum: 3.76 ppm/59.6 ppm and 3.84 ppm/55.3 ppm. The methoxy proton at 3.76 ppm showed a nOe cross peak with H-2' in NOESY and was long-range coupled to C-3 in HMBC. Therefore, it should be 3-OCH₃. Likewise, because the methoxy proton at 3.84 ppm showed a nOe peak with H-3' and long ranged coupling peak with C-4' in NOESY and HMBC, respectively, the methylated position is 4'-OH. As a result, the final structure of the reaction product was 3,4'-dimethoxy, 5,7-dihydroxyl kaempferol, which was regioselectively methylated product from kaempferol. Ermanin was successfully synthesized from kaempferol with two transgenic *E. coli*.

Production of ermanin was monitored periodically for 128 hr. SOMT-2 and ROMT-9 was induced as above. The culture

**Figure 2.** Production of ermanin using *E. coli* transformant expressing SOMT-2 and ROMT-9.

was collected periodically and the amounts of reactant and product quantified with HPLC. As shown in Figure 2, the amount of kaempferol continued to decrease over time and was completely metabolized after 24 hrs. In contrast, 3-methoxy kaempferol increased in quantity until 8 hrs and 4'-methoxy kaempferol increased until 16 hrs. The ermanin appeared after 8 hrs and continued to increase until 128 hrs. After 48 hrs incubation, more than 90% of kaempferol was converted into ermanin. Thus, 13.1 mg of ermanin was obtained from 14.3 mg of kaempferol with a yield of 91%. This approach does not require an enzyme purification step and *S*-adenosyl-L-methionine as a cofactor. Thus, it may be suitable for the production of large amounts of ermanin.

Acknowledgements. This work was supported by a grant of R01-2004-10688-0 (KOSEF) and KRF 2004-F00019 (KRF).

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