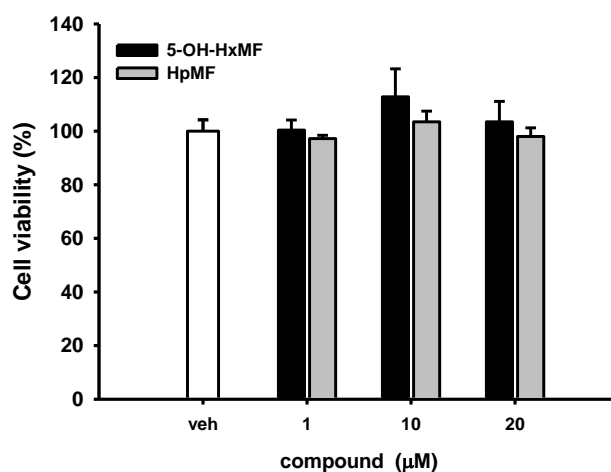
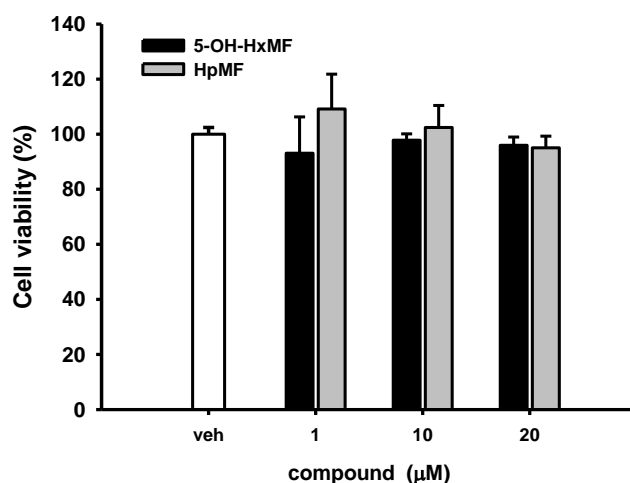


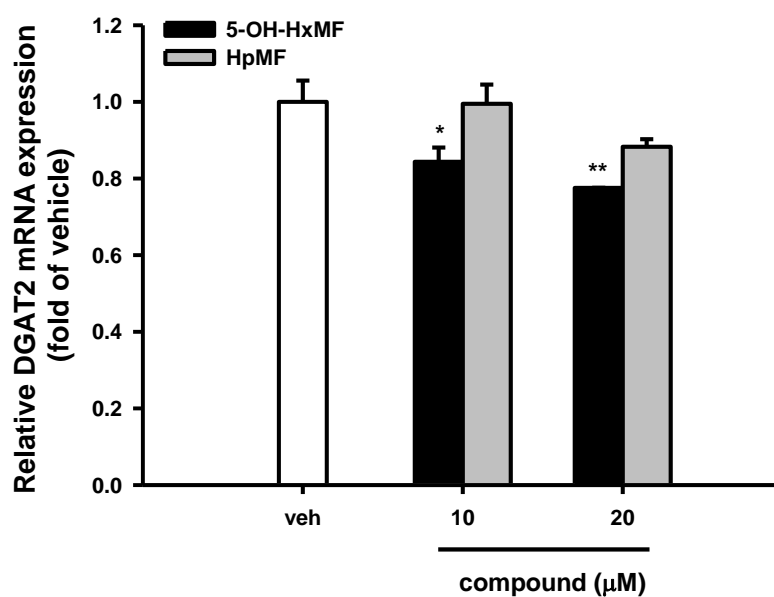
A



B



**Supplemental Figure 1.** Effects of 5-hydroxy-3,6,7,8,3',4'-hexamethoxyflavone (5-OH-HxMF) and 3,5,6,7,8,3',4'-heptamethoxyflavone (HpMF) on the cell viabilities of THP-1-derived macrophages and HepG2 cells. (A) THP-1-derived macrophages were treated with the indicated concentration of agent for 48 h and cell viability was analyzed by MTT as described in Materials and Methods. (B) HepG2 cells were cultured in 6-well plates until 80% confluent, and then changed to LPDS-containing medium for overnight. Cells were then treated with the indicated compound or control vehicle (0.1% v/v DMSO) for 48 h and cell viability was analyzed by MTT as described in Materials and Methods.



**Supplemental Figure 2.** Effects of 5-hydroxy-3,6,7,8,3',4'-hexamethoxyflavone (5-OH-HxMF) and 3,5,6,7,8,3',4'-heptamethoxyflavone (HpMF) on the expression of DGAT2 in HepG2 cells. HepG2 cells were cultured in 6-well plates until 80% confluent, and then changed to LPDS-containing medium for overnight. Cells were then treated with the indicated compound or control vehicle (0.1% v/v DMSO) for 24 h. Total cellular RNA was prepared and the expression of DGAT2 was analyzed, as described in the Materials and methods section. Data represent the mean  $\pm$  SD of three independent experiments relative to the value of vehicle control. \* $p$ <0.05 and \*\* $p$ <0.01 represent significant differences compared with the vehicle control.

**Supplemental Table 1.** Effects of nobiletin (NOB), 5-demethylnobiletin (5-demethyl NOB), 3',4'-didemethylnobiletin (DTF), 5-hydroxy-3,6,7,8,3',4'- hexamethoxyflavone (5-OH-HxMF) and 3,5,6,7,8,3',4'-heptamethoxyflavone (HpMF) on DiI-oxLDL uptake activity and mRNA expression of CD36 and SR-A in THP-1-derived macrophages <sup>a</sup>.

<b>Citrus PMF</b>	DiI-oxLDL uptake (%)	oxLDL-induced CD36 (%) <sup>a</sup>	oxLDL-induced SR-A (%)
<b>NOB</b>			
10 μM	93.3 ± 10.2 <sup>1</sup>	N/A	N/A
20 μM	61.8 ± 2.3 <sup>**1</sup>	N/A	N/A
<b>5-demethyl NOB</b>			
10 μM	74.4 ± 1.2 <sup>**2</sup>	42.6 ± 0.4 <sup>**2</sup>	60.5 ± 2.4 <sup>**2</sup>
20 μM	62.7 ± 0.9 <sup>**2</sup>	53.4 ± 9.2 <sup>**2</sup>	70.2 ± 9.3 <sup>**2</sup>
<b>DTF</b>			
10 μM	61.8 ± 2.2 <sup>**1</sup>	N/A	N/A
20 μM	36.9 ± 1.5 <sup>**1</sup>	61.2 ± 5.7 <sup>*</sup>	38.4 ± 7.9 <sup>*</sup>
<b>5-OH-HxMF</b>			
10 μM	67.4 ± 7.5 <sup>*</sup>	82.2 ± 3.5 <sup>*</sup>	87.3 ± 3.4 <sup>*</sup>
20 μM	60.1 ± 6.1 <sup>*</sup>	82.1 ± 3.0 <sup>*</sup>	89.4 ± 4.1 <sup>*</sup>
<b>HpMF</b>			
10 μM	74.6 ± 9.4 <sup>*</sup>	85.6 ± 5.2 <sup>*</sup>	81.3 ± 20.2
20 μM	70.5 ± 3.2 <sup>*</sup>	88.8 ± 5.8 <sup>*</sup>	98.2 ± 9.0

<sup>a</sup>Data (mean ± SD) are expressed as percentage of the vehicle control (n=3). \*, *p*<0.05 and \*\*, *p*<0.01 represent significant differences compared with the vehicle control in the presence of oxLDL.

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2. J. H. Yen, C. Y. Weng, S. Li, Y. H. Lo, M. H. Pan, S. H. Fu, C. T. Ho and M. J. Wu, *Mol. Nutr. Food Res.*, 2011, **55**, 733-748.