

## Supplementary Information for

# The preparation of ginsenoside Rg5, its antitumor activity against breast cancer cells and its targeting of PI3K

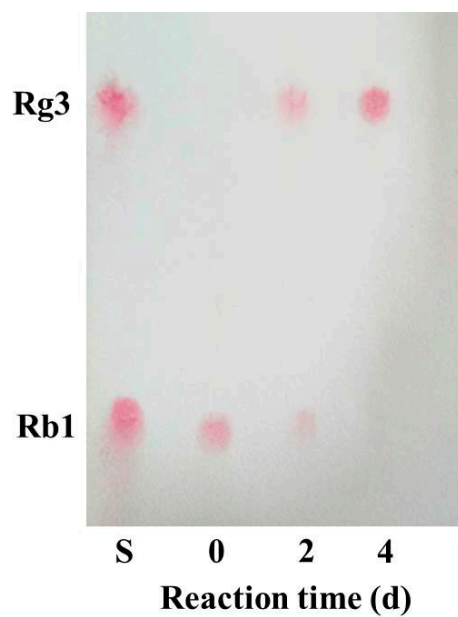
Yannan Liu<sup>a,b,c</sup>, Daidi Fan<sup>a,b,c\*</sup>

<sup>a</sup>Shaanxi Key Laboratory of Degradable Biomedical Materials, School of Chemical Engineering, Northwest University, 229 North Taibai Road, Xi'an, Shaanxi 710069, China

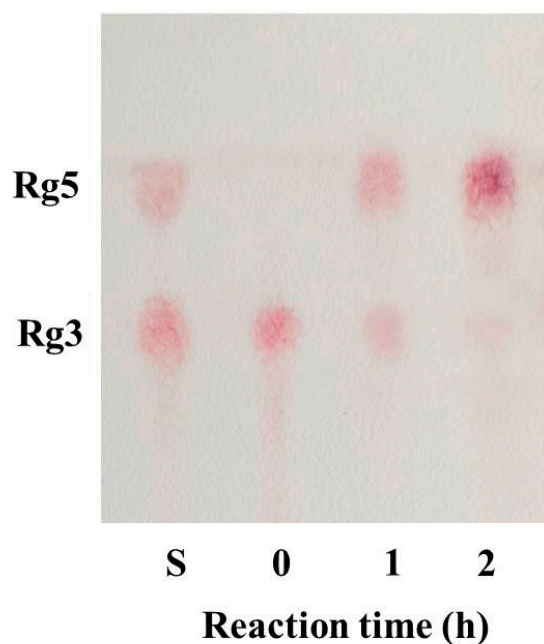
<sup>b</sup>Shaanxi R&D Center of Biomaterials and Fermentation Engineering, School of Chemical Engineering, Northwest University, 229 North Taibai Road, Xi'an, Shaanxi 710069, China

<sup>c</sup>Biotech. & Biomed. Reserch Institute, Northwest University, Taibai North Road 229, Xi'an 710069, Shaanxi, China.

Correspondence: Daidi Fan \*E-mail addresses: fandaidi@nwu.edu.cn



**Supplementary Figure S1.** Thin layer chromatography (TLC) analysis of time-course transformation of ginsenoside Rb1 by  $\beta$ -glucosidase. S is the mixture of ginsenoside standards.



**Supplementary Figure S2.** Thin layer chromatography (TLC) analysis of time-course transformation of ginsenoside Rg3 at 121°C with high pressure processing. S is the mixture of ginsenoside standards.

**Supplementary Table S1.** ADMET property evaluation

Drug	BBB (%)	PPB (%)	Chronic toxicity(%)	hERG inhibition	Developmental toxicity	Reproductive toxicity	Hepatotoxicity	AMES toxicity
Rg5	Non (82.47)	81.41	Non (73.67)	Non (75.01)	Non (98.28)	Non (91.86)	Non (71.45)	Non (91.65)