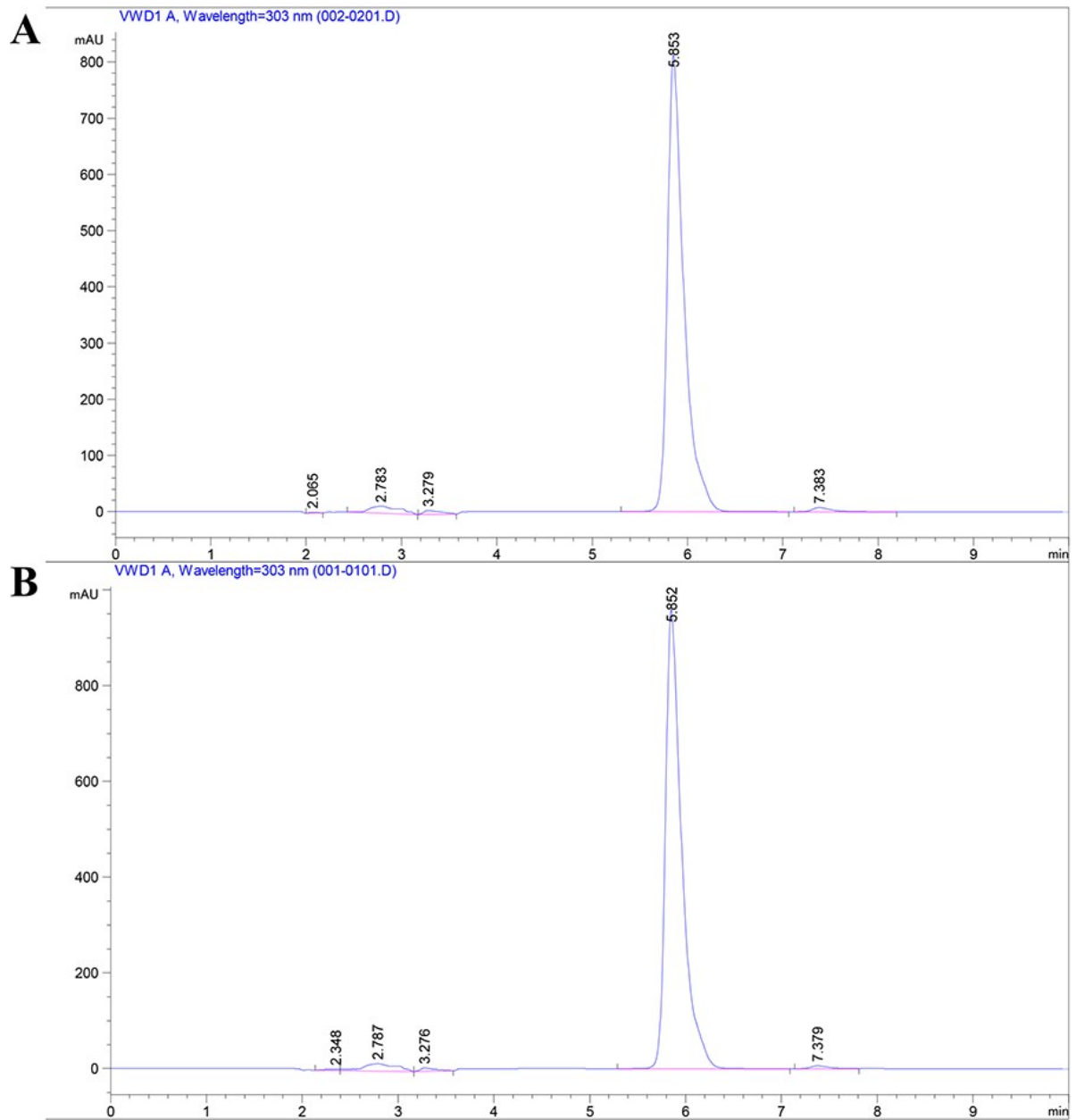


**Table S1** Primers used in this study

Primers name	Primers sequence	Recognition sites
<i>bgl</i> -1-F	CGGGATCCTTGTCAAACAAAGAACGGTTTTT	<i>Bam</i> H I
<i>bgl</i> -1-R	ATAAGAATGCGGCCGCTTAGTTGATCAGACGATGGTATTC	<i>Not</i> I
A		
<i>bgl</i> -2-F	CGGGATCCATGGTCAAAGAAAGAAGCTTTCTT	<i>Bam</i> H I
<i>bgl</i> -2-R	ATAAGAATGCGGCCGCCTATCCTCCTTGAGGCAGGAT	<i>Not</i> I
<i>bgl</i> -3-F	GGAATCCATATGATGAAAAAGTTGGCTCAAAGAGT	<i>Nde</i> I
<i>bgl</i> -3-R	CGGGATCCTCATCGGTTCTTTTCATTCATT	<i>Bam</i> H I
<i>bgl</i> -4-F	CGGGATCCTTATTCGTCTGTTTCCTCATAGC	<i>Bam</i> H I
<i>bgl</i> -4-R	ATAAGAATGCGGCCGCTTAATGAAGATTACCAATCCAGTG	<i>Not</i> I
C		
<i>bgl</i> -5-F	CGGAATCCTTAAAAACCGTTATTCTTTGAAAC	<i>Eco</i> R I
<i>bgl</i> -5-R	CCCAAGCTTTTAATGTCGATGGAAACTAACAAGC	<i>Hind</i> III

**Table S2.** universal citric acid–disodium hydrogen phosphate buffers

pH	0.2 mol/L Na <sub>2</sub> HPO <sub>4</sub> (mL)	0.1 mol/L C <sub>6</sub> H <sub>8</sub> O <sub>7</sub> (mL)
3.0	4.11	15.89
3.2	4.94	15.06
3.4	5.70	14.30
3.6	6.44	13.56
3.8	7.10	12.90
4.0	7.71	12.29
4.2	8.28	11.72
4.4	8.82	11.18
4.6	9.35	10.65
4.8	9.86	10.14
5.0	10.30	9.70
5.2	8.28	11.72
5.4	8.82	11.18
5.6	9.35	10.65
5.8	9.86	10.14
6.0	12.63	7.37
6.2	13.22	6.78
6.4	13.85	6.15
6.6	14.55	5.45
6.8	15.45	4.55
7.0	16.47	3.53
7.2	17.39	2.61
7.4	18.17	1.83
7.6	18.73	1.27
7.8	19.15	0.85
8.0	19.45	0.55



**Fig. S1** The original copies of HPLC  
(A) HPLC of product resveratrol (B) HPLC of resveratrol standard

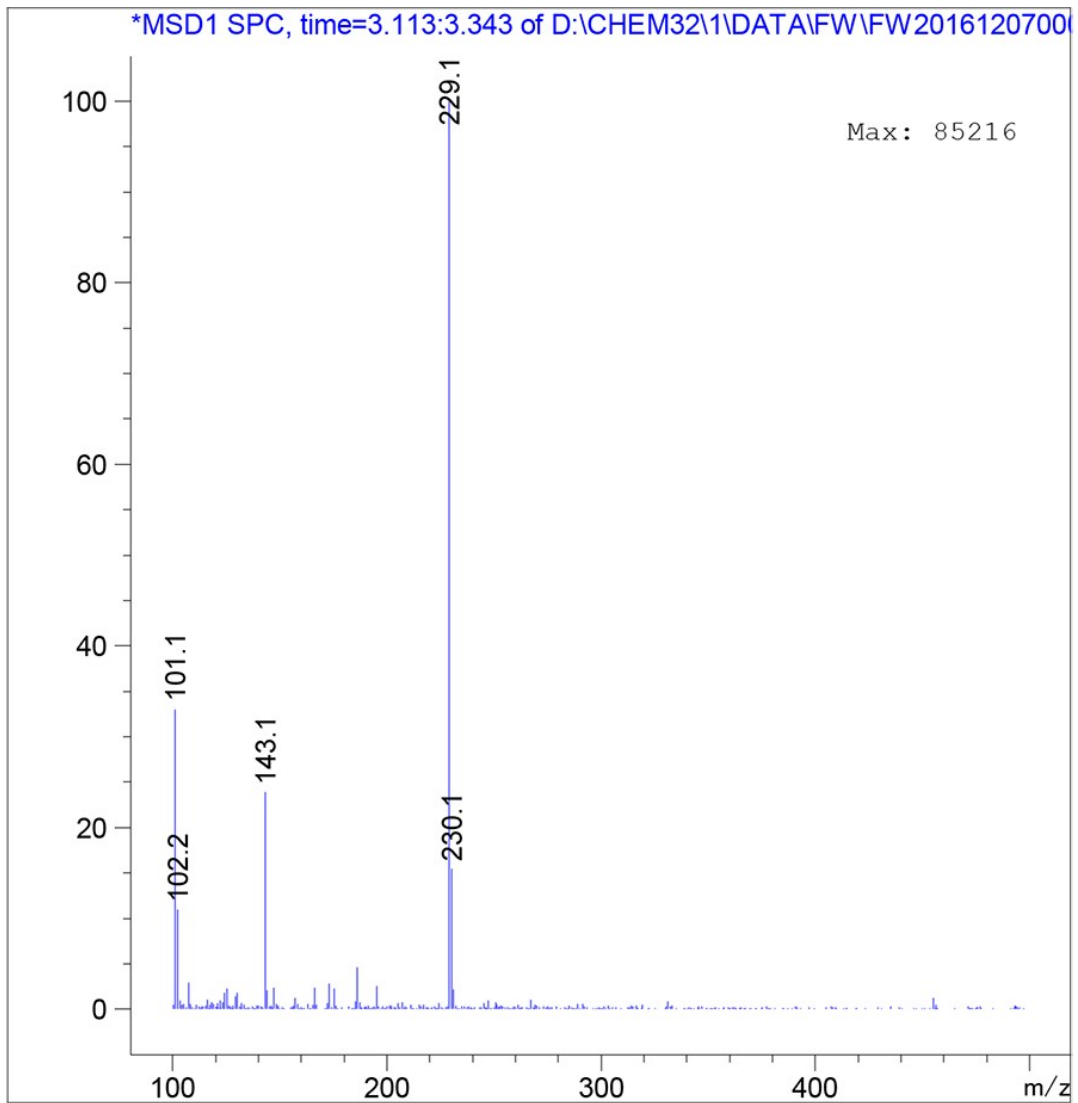
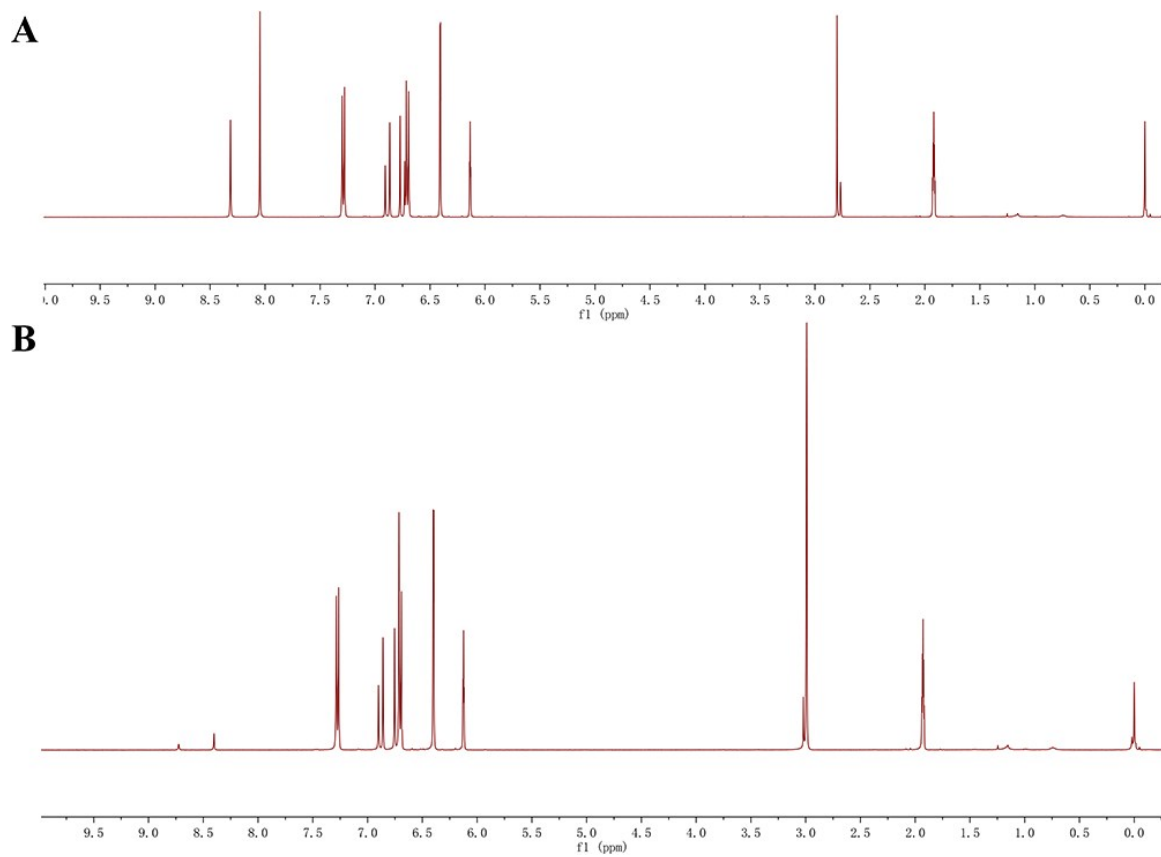


Fig. S2 The original copies of MS



**Fig. S3** The original copies of  $^1\text{H-NMR}$   
(A)  $^1\text{H-NMR}$  of resveratrol in Acetone-D6 (B)  $^1\text{H-NMR}$  of resveratrol in Acetone-D6 adding  $\text{D}_2\text{O}$