

Article

Tiliacora triandra (Colebr.) Diels Leaf Aqueous Extract Inhibits Hepatic Glucose Production in HepG2 cells and Type 2 Diabetic Rats

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Supplementary Table S1. Retention time and parameters of calibration curve of standard phenolic and flavonoid compounds for HPLC-PDA method validation.

Compound	Detected at Wavelength (nm)	RT (min)	Linear Range (µg/mL)	Regression Equation	Correlation Coefficient (r ²)
Gallic acid	280	6.45	0.5–100	y = 31428x – 15432	0.9999
Catechin	280	14.52	0.5–100	y = 7333.4x + 1054.7#	0.9998
ECGC	280	17.63	0.5–100	y = 15109x – 14908	0.9993
EC	280	18.35	0.5–100	y = 7723.5x + 1562.9	0.9999
Quercetin	360	28.15	0.5–100	y = 22828x + 3052.2	0.9998

RT = retention time (min); y = peak area; x = concentration of standard (mg/l); r² = correlation coefficient of 3 data points in the calibration curve.

Supplementary Table S2. Gradient program for the HPLC analysis.

Times (min)	Solvents (%)		
	A (0.1 Phosphoric Acid)	B (Methanol)	C (Acetonitrile)
0	95	0	5
10	80	10	10
20	70	15	15
30	0	0	100
35	0	0	100

Supplementary Table S3. Primer sequences and expected amplicon sizes for gene amplification.

cDNA	Forward Primer (5'–3')	Reverse Primer (5'–3')	Amplicon Size (bp)
rCu-Zn SOD	GCAGAAGGCAAGCGGTGAAC	TAGCAGGACAGCAGATGAGT'	387
rGPx	CTCTCCGCGGTGGCACAGT	CCACCACCGGGTCGGACATAC	297
rCAT	CCTCCTCGTTCAAGATGTGGTTTTTC	CGTGGGTGACCTCAAAGTATCCAAA	122
rPEPCK	CTCACCTCTGGCCAAGATTGGTA	GTTGCAGGCCCCAGTTGTTGA	190
rG6Pase	AACGTCTGTCTGTCCCGGATCTAC	ACCTCTGGAGGCTGGCATTG	133
rActin	CCTAAGGCCAACCCTGAAAA	GGAGCGCGTAACCCTCATAG	181
h Cu-Zn SOD	AAAACACGGTGGGCCAAAG	GTGCGGCAATGATGCA	141
hGPx	TTCCCGTGCAACCAGTTTG	TTCACCTCGACTTCTCGAA	128
hCAT	TCATATACCTGTGAACTGTC	ATAGAATGACCGCACCTGAG	229
hPEPCK	AAGAGACACAGTGCCCATCC	ACGTAGGGTGAATCCGTCAG	201
hG6Pase	GAGACTGGCTCAACCTCGTC	CCTGGTCCAGTCTCACAGGT	139
hGAPDH	AGCCTTCTCCATGGTGGTGAAGAC	CGGAGTCAACGGATTGGTTCG	308

Cu-Zn SOD-copper zinc superoxide dismutase; GPx-glutathione peroxidase; CAT-catalase; PEPCK-Phosphoenolpyruvate carboxykinase; G6Pase- Glucose 6-phosphatase; GAPDH- Glyceraldehyde 3-phosphate dehydrogenase.