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In vitro α-amylase inhibitory activity of Gmelina arborea Roxb. aqueous extract encapsulated chitosan nanoparticles

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Background: *Gmelina arborea* Roxb. is a medicinal plant used in complementary medicine with antidiabetic properties. *G. arborea* aqueous extract (GAE) has been reported to exert α -amylase inhibitory activity. However, low bioavailability of polyphenols and flavonoids, and lack of appropriate delivery systems delimit use of GAE as a commercially viable antidiabetic agent. Chitosan, a promising carbohydrate biopolymer is gaining scientific attention in encapsulation of herbal extracts. The present study was to determine *in vitro* α -amylase inhibitory activity of GAE encapsulated chitosan nanoparticles.

Objective: To investigate the α -amylase inhibitory activity of GAE encapsulated chitosantripolyphosphate (CS-TPP) nanoparticles.

Method: Dried plant materials of *G. arborea* were extracted using distilled water under ultrasonication (40 kHz, 37 °C, 30 minutes) and the aqueous extract was freeze-dried. The CS-TPP nanoparticles loaded with freeze-dried GAE (37.5 mg) were prepared using the ionotropic gelation method (750 rpm, 35 °C, 60 minutes). Resulting nanoparticles were dissolved in DMSO and a concentration series (2.53-8.23 mg/mL) was prepared. The α-amylase inhibitory assay was conducted in triplicate and the absorbance was measured (λ_{max} =540 nm) for the GAE encapsulated and unloaded CS-TPP nanoparticles using acarbose as the standard. The α-amylase inhibitory activity is expressed as IC₅₀.

Results: Percentage inhibition of α -amylase inhibitory activity of unloaded chitosan nanoparticles were -8.4±1.3%. The percentage inhibition of encapsulated GAE (0.029 mg/mL) with the CS-TPP was 21.1±1.5%. The IC₅₀ value of encapsulated CS-TPP nanoparticles with GAE was 7.8±0.2 mg/mL whereas the IC₅₀ value of GAE was 32.0±6.7 μ g/mL. The acarbose had an IC₅₀ value of 11.7±0.3 μ g/mL.

Conclusion: The results reveal that the α -amylase inhibitory activity of GAE was retained in the encapsulated GAE and the matrix material (CS-TPP) did not interfere with the α -amylase inhibition. Future studies are warranted on the stability and *in vitro* releasing profile of GAE encapsulated CS-TPP nanoparticles.

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