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**Phytochemical study of some medicinal compounds
present in *Cordia myxa* L. plant cultivated in Iraq**

A Thesis

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Abstract

Cordia myxa L. (Arabic name: Bamber) of the family Boraginaceae. The origin of this plant is tropical Asia. This plant is distributed in a wide range in the Arab Gulf region while in Iraq it is cultivated in a wide range in the middle and south regions such as Baghdad and Al-Basrah.

This plant contains many active constituents including: flavonoids (robinin, rutin, datiscoside and hesperidin, one flavone aglycone, dihydrorobinetin), two phenolic acids (chlorogenic acid and caffeic acid), saponins, terpenes and sterols (betasitosterol) and coumarins.

C. myxa L. has high medicinal uses such as anti-inflammatory, analgesic, antimicrobial, antiasthmatic, gastroprotective, anticancer and antileishmania.

This study concerned with extraction, identification, isolation, and purification of some active constituents which are flavonoid (rutin) and phenolic acid (chlorogenic acid) from *C. myxa* L. leaves and detection of presence of these compounds in crude extracts of *C. myxa* L. fruits. In addition, detection of the presence of other active constituents which are (betasitosterol and coumarin) in the crude extracts of *C. myxa* L. leaves and fruits.

One extraction method was tried for flavonoid (rutin) by Soxhlet apparatus with 70% ethanol then fractionated with hexane, then with ethylacetate, one extraction method for phenolic acid (chlorogenic acid), by Soxhlet extraction with 80% methanol, one extraction method for sterols (betasitosterol) by petroleum ether (60-80)°C and one extraction method for coumarin with 50% ethanol.

The preliminary identification of the flavonoid, phenolic acid, sterol and coumarin was done by thin layer chromatography (TLC) where different solvent systems were tried and ultra-violet at (254 and 365) nm were used

for detection of rutin, chlorogenic acid , coumarin. while 5% H₂SO₄ spray reagent with heating at 100°C was used for betasitosterol detection. This identification was further augmented by high performance liquid chromatography (HPLC) and then isolation and purification of rutin and chlorogenic acid.

The identification of isolated compounds (rutin and chlorogenic acid) were done by a direct comparison with authentic standards using thin layer chromatography (TLC), melting point (M.P), fourier transform infrared (FTIR), and High performance liquid chromatography (HPLC) analysis.