Abstract

The Tamarix dioica (T. dioica) is widely used medicinal plant to cure many chronic ailments. The dioica is being used to treat diabetes mellitus in traditional medicinal system, however little scientific evidence is available on this plant in this context. The plant was collected from Khyber Pakhtunkhwa province of Pakistan. The current study involves the fractionation of crude methanolic extract of T. dioica using, n-hexane, ethyl acetate, chloroform and n-tranol, the dried plant material obtained from each solvent was dissolved in methanol and further investigated for its antimicrobial activity and other studies. The screening for antioxidant activity using 2,2-diphenyl-1-picrylhydrazyl (DPPH) assay was carried out. The total antioxidant activity was determined by phosphomolybdenum method The *in vitro* antidiabetic potential was assessed by measuring α -glucosidase inhibition. Total phenolic and flavonoid contents along with metal chelating activity were also determined for each fraction. The metabolites were identified using highly sensitive and emerging α -NMR technique.

The antibacterial activity was tested by agar well diffusion method against three Grampositive strains (Bacillus sp., Actinomyces viscosus, Corynebacterium minutissium) and three Gram-negative strains (Escherichia coli, Pseudomonas fluorescens, Klebsilla pneumoniae). The antifungal activity was tested by agar diffusion method against three fungal strains (Aspegillus flavus, Aspergillus niger, Penicillium digitatum). The chloroform fraction exhibited the highest inhibitory effect in all the tested bacterial strains especially against Escherichia coli and Klebsilla pneumoniae with zone of inhibition of 7.50mm and 11.10mm respectively. The ethyl acetate fraction was also active against Escherichia coli with zone of inhibition of 11.50mm. In antifungal activity ethyl acetate fraction showed highest %age of zone of inhibition of 71.58% against Aspegillus flavus and 68.57% against Aspergillus niger, It was noted that all results were concentration dependant that is higher the concentration of plant extract more will be the activity.

The results revealed, the ethyl acetate fraction as the most potent with DPPH scavenging activity of 84.44±0.21% and α-glucosidase inhibition with IC50 value of 122.81±2.05 μg/mL. The total antioxidant activity of ethyl acetate fraction was 163.12±1.66mg ASE/g PE followed by chloroform fraction 152.12±1.18 mg ASE/g PE. The total phenolic and flavonoid content values of 205.45±1.36 mg gallic acid equivalent per gram dried extract and 156.85±1.33 mg quercetin equivalent per gram dried extract were obtained for ethyl acetate

The results of metal chelating ability of T.dioica showed that its ethyl acetate and form fractions possess almost the same iron chelating ability. Ethyl acetate fraction percentage iron chelating of $58.28\pm0.54\%$ followed by chloroform with $57.92\pm0.28\%$ chelating ability, these values were much lower than the value given by the standard $1.38\pm0.05\%$.

bucketing of H-NMR spectra indentified 22 metabolites including some marmacologically important like tamarixetin, tamaridone, quercetin, rutin, apigenin, carechin, kaempferol, myricetin and isorhamnetin. Leucine, lysine, glutamic acid, aspartic acid, serine and tyrosine were the major amino acids identified in ethyl acetate fraction. The molecular docking analysis provided significant information on the binding affinity among secondary metabolites and α -glucosidase. These metabolites were most probably responsible for the antioxidant activity and α -glucosidase inhibitory potential of ethyl acetate fraction. The study ascertained the ethnomedicinal use of T. dioica to manage diabetes mellitus and may be a helpful lead towards naturopathic mode for anti-hyperglecemia.

Keywords: Tamarix dioica, antibacterial, antifungal, antioxidant, α-glucosidase, ¹H-NMR, metabolites.