Mineral Analysis Oftamilnadia Uliginosa Retz.Tirveng& Sastre (Rubiaceae).

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Abstract

Fresh leaves and fruits of Tamilnadiauliginosa were used for the present study. In the present study, elements like Mg,K,Na,Fe,Mn,Ca,Pb,Cu present in T.uliginosawas carried out to assess its medicinal properties. The resultalso is helpful in standardization and identification of plant as well as crude drug for the future analysis and preparation of drug.

Keywords: Mineral Analysis, Elements, Tamilnadiauliginosa,

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Introduction

The earth is endowed with a rich wealth of medicinal plants. In India, plant derived drugs have been used in traditional systems of medicines such as Unani, Ayurveda, Siddha and Homoeopathysince ancient timesCharakasamhita(1000BC) detailing the use of medicinal plants used inAyurvedic system of medicine, recorded the use of over 340 drugs from plants. Plants providedirectly usable drugs, and a variety of chemical compounds that can be used as starting points for the synthesis of new drugs with improved pharmacological properties (Ballabh et al., 2008). Medicinal herbs and their preparations (hot and cold infusions, decoction, and tinctures) are widely used by human beings all over the world (Arpadjan et al., 2008). Medicinal plants can be defined as plants that are commonly used for treating and preventing specific ailments and diseases (Anselem, 2004). These plants are either "wild plant species" those which grow spontaneously in self maintaining populations in natural or semi-natural ecosystems existing independently of direct human actions in contrastto the "domesticated plants species" those that have arisen through human actions such as selection or breeding and depend on management for their existence. Even though herbs had been priced for their medicinal, flavoring and aromatic qualities for centuries; the synthetic products of the modern age surpassed their

importance.Today however, the herbal products symbolize safety in comparison to the synthetics that are considered unsafe to both human and environment. According to World Health Organization (WHO), more than 80% of the world's populations, mostly in developing countries depend on traditional plant based medicines for their primary health care needs (Pier Angeli et al., 2009). Medicinal plants are the richest bioresource of drugs in traditional systems of medicines(Adnan et al., 2014). The ancient people observed that some ingredients used in food had the specific properties for maintaining or eliminating certain diseases and maintaining good health (Vijayakumar et al., 2009). Scientific validation, including the nutritional, pharma-cognostic and phytochemical characterization is the essential step in the characterization of a medicinal plant (Habibul Hassan et al., 2014).

Rubiaceae, the coffee family is the fourth largest flowering plant family which includes about 500 genera and 6,000 species of herbs and shrubs (Evans, 2002). Members of the family tend to be concentrated in warmer and tropical climates around the world. A wide variety of growth forms are present in the family. While shrubs are the most common, members can also be trees, lianas or herbs. The family name isafter the madder genus *Rubia*whichisderived from the Latin word "ruber" meaning "red". The DeepthymolM.J. & PraveenDharT.

family was described by Antoine Laurent de Jussieu. The group contains many commonly known plants, including the economically important coffee (*Coffea*), quinine (*Cinchona*), gambier (*Uncaria*), horticulturally valuable madder (*Rubia*), West Indian jasmine (*Ixora*), partridge berry (*Mitchella*), *Morinda*, *Gardenia*, and *Pentas*.

Materials and Methods Study area

The study selected area isMuthangaWild Life Sanctuary located in WayanadDistrict of Kerala, comingunder Nilgiri Biosphere Reserve on the Western Ghats (Figs.1and 2). The MuthangaWild Life Sanctuary is contiguous with the protected area network of Nagarhole and Bandipur National Park of Karnataka on the northeast and Mudumalai National Park of Tamil Nadu on the southeast. Rich in biodiversity, the sanctuary is an integral part of the Nilgiri Biosphere Reserve, which has been established with the specific objective of conserving the biological heritage of the region. The total extent of the area is 344.44km² and is divided into two discontinuous portions with revenue lands in between. TheNorth West portion of the sanctuary has only one range namely, Tholpetty covering an area of 77.67km². This range is contiguous with Nagarhole National Park, also known as Rajiv Gandhi National park, Nagarhole in the

northeast, Kakkankotte reserve forest in the north and Brahmagiri hills of North Wayanad forest division in the east. The southern portion of the sanctuary comprises an area of about 266.77km²(Ratheesh *et al.*, 2011).

Plant material

Fresh leaves and fruits of Tamilnadiauliginosa were collected from the Muthanga Wild Life Sanctuary and Kuruva Island.Fresh leaves and fruitswere washed and useddirectly for the analysis of macroscopic and microscopic characters. For other analysesdried powder of the plant parts were used. For this the plant parts were subjected to shade drying for about ten weeks. The dried plant materials were powderedand the powder was passed through a mesh sieve and stored in air tight containers. The species for the proposed study was identified and authenticated by using the Flora of Presidency of Madras by Gamble.A voucher specimen of the plant has beendeposited in the Herbarium of Department of Botany, University of Kerala, Kariavattom. (KUBH5810). Relevant photographs of the plants were taken from the field itselfand important points were noted in the field book on the spot.

Method of Mineral analysis

The estimation of mineral content wascarried outas per the procedure mentioned by health protection branch laboratories, bureau of nutritional science (Okoro*et* al., 2012). The sample was

prepared by dry ashing and various elements were calculated by flame atomic absorption spectroscopy. The samples were burned to ashes in muffle furnace at 450°C for 16 h. Then the ash was wetted with water and sufficient amount of nitric acid was added to cover the ash. Refluxed on a hot plate for 1h.Then heat was reduced and gentlyevaporated the acid. Again samples were returned to muffle furnace at 375°Cfor 1hour.Repeated heatinguntil a white ash is obtained. Added 2.5ml dilute HCl and dissolved the ash by boiling the solution, cooled and made up to appropriate volume with water. Determination of elements was done using flame atomic absorption spectroscopy.

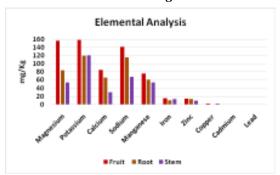
Results

The amounts of various elements in decreasing order in fruits were potassium, magnesium, sodium, calcium, manganese, iron, zinc and copper respectively. In stem the occurrence of these elementswas in the decreasing order ofpotassium, sodium, manganese, magnesium, iron, zinc and copper while in roots it was in the orderpotassium, sodium, magnesium, calcium, manganese, zinc, iron and copper.

Result of mineral analysis of fruit, root, and stemis represented in Fig. 1

Fig. 1 Elemental analysis of fruit,root and stem of *T.uliginosa*

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Summary and Conclusions

Plants have been the source of many important drugs because they are able to produce various chemical entities and bioactive molecules through the process known as metabolism. These bioactive molecules are used as the active ingredients of modern medicine or as the lead compounds for new drug discovery. Though the Ayurvedicsystem of medicine has a long history of use of medicinal plants, yet they lack adequate scientific documentation of plants and standardization of drugs particularly based on modern scientific knowledge. The medicinal properties of aplant depend upon the bioactive phytochemical constituents present in it and which shows various physiological effects on human body. So through phytochemical DeepthymolM.J. & PraveenDharT.

screening of plants can detect the various important compounds which can be used as the base of modern drugs for curing various diseases. In the present study,elements like Mg,K,Na,Fe,Mn,Ca,Pb,Cu present in

T.uliginosa was carried out to assess its medicinal properties. The resultalso is helpful in standardization and identification of plant as well as crude drug for the future analysis and preparation of drug.

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