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## BIOCHEMICAL ESTIMATION OF PRIMARY METABOLITES OF *PHYLA NODIFLORA* L. GREENE

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### ABSTRACT

*Phyla nodiflora* is an evergreen perennial herb found near the water bodies. It belongs to family Verbenaceae. It is commonly known as frog fruit or turkey tangle. The present investigation is the estimation of primary metabolite content in various plant parts. The investigation revealed that maximum amount of soluble sugar was found in stem, maximum amount of starch found in stems, leaves showed the maximum amount of protein, fruits showed maximum amount of lipids, ascorbic acid was found in highest amount in leaves, fruits showed maximum amount of phenols.

**INTRODUCTION:** Plants have the ability to synthesize wide range of chemical compounds, which are used to perform important biological function and also to defend itself from predators like insects, fungi and herbivorous mammals. Many of these phytochemicals have beneficial effects on long term health and also in the cure of human disease. At least 12000 such compounds have been isolated, which is actually about less than 10% of the total <sup>1</sup>. These medicinal plants or herbs also show the antimicrobial and chemopreventive properties <sup>2</sup>. Ethnobotany is recognized as the effective way to discover future medicine. In 2001, researchers identified 122 compounds which are used in modern medicine, 80% of them have the ethnomedical use of the active element of the plant <sup>3</sup>.

*Phyla nodiflora* is an aquatic, evergreen, perennial herb which belongs to family Verbenaceae. *Phyla* is native of South America and United States and commonly known as frog fruit, Turkey tangle. The plant is diuretic <sup>4</sup>, has gastroprotective effect <sup>5</sup>, anti inflammatory and antineoplastic activity <sup>6</sup>, antioxidant activity <sup>7</sup> useful in

pain in knee joints and in lithiasis, antispasmodic <sup>8</sup>, antibacterial activity <sup>9</sup> and antifungal activity <sup>10</sup>, chutney made from the leaves and fruits are eaten to relieve the irritation of internal piles.

The presence of these medicinal activities is due to presence of different metabolites i.e. primary metabolites and secondary bioactive compounds. Primary metabolites are of prime importance and are essentially required for example sugar, protein, lipid, starch. Many primary metabolites act as precursor of pharmacologically active metabolites.

The present study deals with the study of primary metabolites present in *phyla nodiflora*.

### EXPERIMENTAL SECTION:

**Collection of Plant Material:** Plant material was collected from the Mawtha region, near Amer, Jaipur. It was then authenticated by Herbarium, Department of Botany, Rajasthan University, Jaipur, Rajasthan, India. The identification number given by the herbarium is RUBL21061.

**Preparation of Extracts:** The stem leaves and fruits of *Phyla nodiflora* was properly washed under tap water, shade dried and powdered. All the plant parts were then evaluated quantitatively to estimate the total levels of soluble sugars, starch, proteins, lipids, phenols, and ascorbic acid following the established methods for the sugars, starch <sup>11</sup>, protein <sup>12</sup>, lipids <sup>13</sup>, phenols <sup>14</sup>, amino acid <sup>15</sup>, ascorbic acid <sup>16</sup>. All experiments were repeated five times for precision and values were expressed in mean  $\pm$  standard deviation in terms of air dried material.

**RESULTS AND DISCUSSIONS:** All the plant parts of *Phyla nodiflora* were evaluated quantitatively for the analysis of total soluble sugars, starch, protein, phenol, lipid, and ascorbic acid.

Plants are rich sources of high value metabolites like proteins, phenols, sugars, starch, lipids, amino acids and ascorbic acids which are directly involved in the normal growth, development and reproduction and are useful in flavoring, fragrances, insecticides, sweeteners and natural dyes <sup>17</sup>.

Starch is biodegradable and renewable in nature. They are increasingly being considered as an eco-friendly alternative to the use of synthetic additives in many other products, including plastics, detergents, pharmaceutical tablets, pesticides, cosmetics and even oil-drilling fluids. The highest amount of starch was observed in stems i.e. 1.98mg/gdw and minimum amount was observed in leaves i.e. .09mg/gdw (**Fig. 1**).

Quantitative estimation of sugar shows that content of sugar is highest in stems i.e. 8mg/gdw and minimum i.e.1.8mg/gdw in leaves (**Fig. 2**).

Plant sugars can be used as artificial sweeteners and they can even help diabetics by supporting the body in its rebuilding <sup>18</sup>.

The total levels of lipid were found to be higher in fruits i.e. pods 60mg/gdw and lowest in stems i.e. 30mg/gdw (**Fig. 3**). The higher amount of plant lipid can be used as essential oils, spice oleoresins and natural food colors. With a strong foundation in research and development, plant lipids have developed products that work with diverse requirements, be it culinary, medicinal or cosmetic <sup>19</sup>.

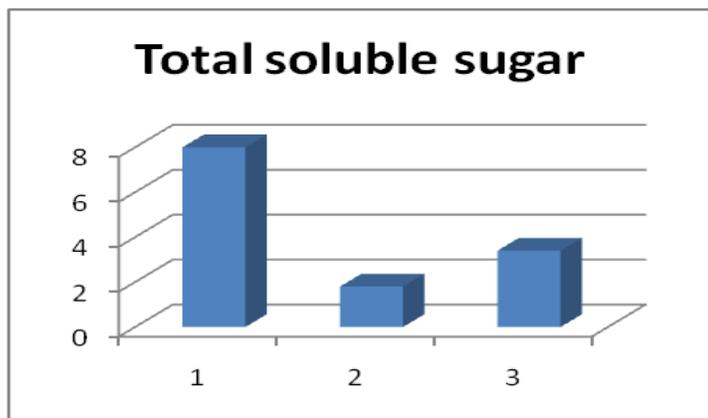
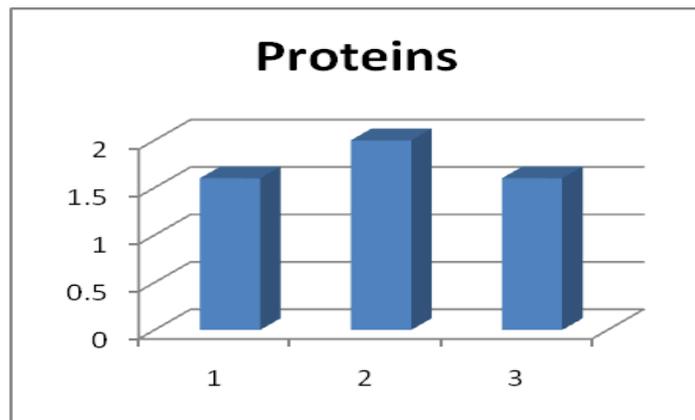
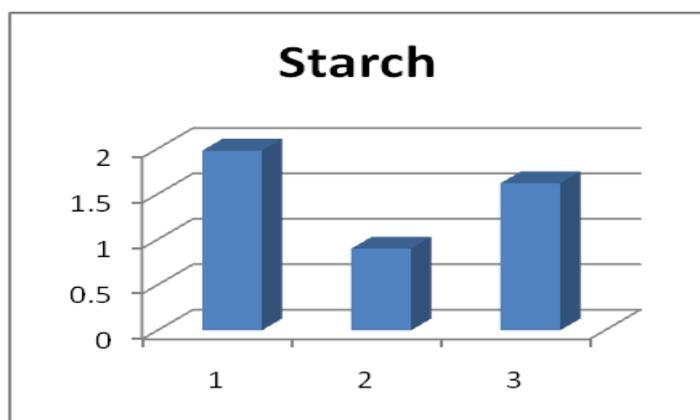
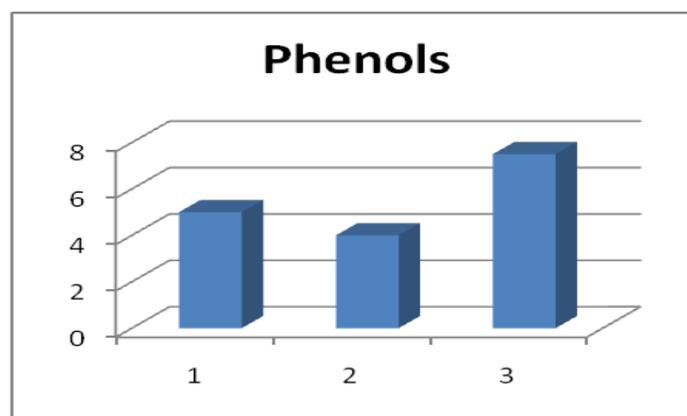
Ascorbic acid (vitamin C) is a familiar molecule because of its dietary significance, most aspects of its metabolism and some aspects of its function in plants are very poorly understood <sup>20</sup>. Total levels of ascorbic acid were found to be highest in leaves i.e. .030 mg/dw and minimum in stems i.e 0.0125mg/gdw (Fig. 4).

Proteins are the primary components of living things. The presence of protein level in the plant points towards their possible increase food value or that a protein base bioactive compound could also be isolated in future <sup>21</sup>. Total levels of protein were found to be highest in leaves i.e. 2mg/gdw and both fruits and stem showed 1.6mg (Fig. 3). Total levels of phenols were found to be highest in fruits i.e. 7.5mg/gdw and lowest in leaves i.e. 4.0mg/gdw (Fig. 4).

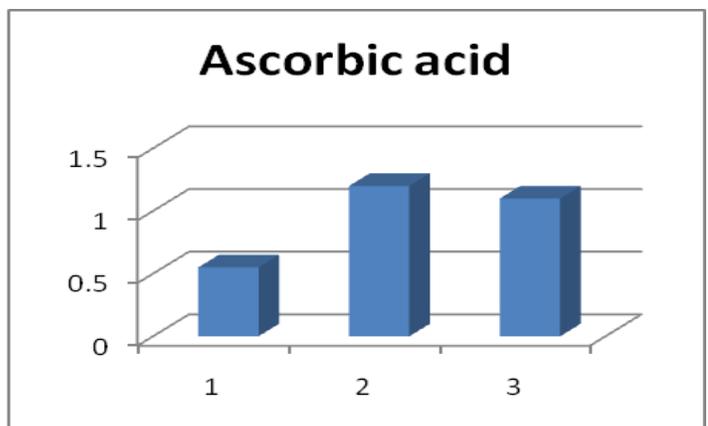
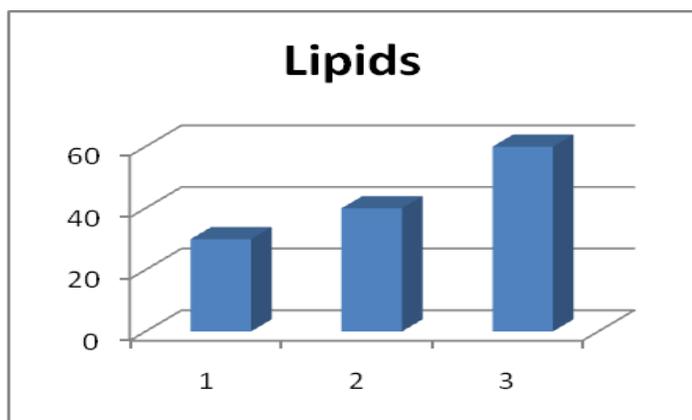
The higher amount of phenols is important in the regulation of plant growth, development and diseases resistance. It can be used as fungicide, pesticides, an antiseptic, disinfectant and in the manufacture of resins, explosives, plastics, detergents and pharmaceutical substances.

**TABLE 1: CONCENTRATION OF PRIMARY METABOLITES IN PHYLA NODIFLORA L. GREENE (MG/GDW)\***

	Stem	Leaves	fruits
Starch	1.98 $\pm$ 0.126	0.9 $\pm$ 0.042	1.62 $\pm$ 0.064
Total soluble sugar	8.0 $\pm$ 0.19	1.8 $\pm$ 0.132	3.4 $\pm$ 0.051
Ascorbic acid	0.0125 $\pm$ 0.63	0.030 $\pm$ 0.04	0.025 $\pm$ 0.025
Lipids	30.0	40.0	60.0
Proteins	1.6 $\pm$ 0.029	2.0 $\pm$ 0.052	1.6 $\pm$ 0.036
Phenols	5.0 $\pm$ 0.080	4.0 $\pm$ .10	7.5 $\pm$ 0.210

FIGURE 1: SOLUBLE SUGARS CONCENTRATION OF *P. NODIFLORA*FIGURE 5: PROTEINS CONCENTRATION OF *P. NODIFLORA*FIGURE 2: STARCH CONCENTRATION OF *P. NODIFLORA*FIGURE 6: PHENOLS CONCENTRATION OF *P. NODIFLORA*

1= Stem; 2= leaves; 3= Fruits

FIGURE 3 : ASCORBIC ACID CONCENTRATION OF *P. NODIFLORA*FIGURE 4: LIPID CONCENTRATION OF *P. NODIFLORA*

**CONCLUSION:** *Phyla nodiflora* contain many primary metabolites like carbohydrates, proteins, phenols, lipids, amino acids and ascorbic acids. Highest amount of soluble sugar were found to be in stems (8.0mg/gdw), starch in stems (1.98mg/gdw), lipids in fruits (60mg/gdw), proteins in leaves (2.0mg/gdw), phenol in fruits (7.5/gdw) and ascorbic acid in leaves (1.20mg/gdw). These results are suggestive of primary bioactive compound of commercial importance and may result in great interest in plants pharmaceuticals. These primary metabolites further used for biosynthesis of secondary metabolites or bioactive compounds<sup>22</sup>.

#### REFERENCES:

1. Tapsell, L.C, Hemphill I, Cobiac L, *et al.* Health benefits of herbs and spices: the past, the present, the future. *Med J Aust* 2006; 185 (4 suppl) S4-24
2. Lai, PK, Roy J. Antimicrobial and chemopreventive properties of herbs and spices. *Curr. Med. Chem.* 2004. :1451-60
3. Fabricant DS, Farnsworth NR The value of plants used in the traditional Medicine for drug discovery. *Environmental health perspective* .March 2001 :109 suppl1(69-75)
4. Sangita Shukala, Rashmika Patel and Rajiv Kukkar. Study of phytochemical and diuretic potential of methanol and aqueous extracts of aerial parts of *phyla nodiflora* linn. *International*

- Journal of Pharmacy and Pharmaceutical Sciences.2009;1(1)85-91
5. Khalil.H, Ismail H, Taye A and Kamel M. Gastro protective effect of *Lippia nodiflora* Linn. extracts in ethanol induced gastric lesions. Pharmacognosy magazine .2007;3(12):258-261
  6. Ahmed F, Selim MST, Das AK and Chouduri MSK. Anti-inflammatory and antinoceptive activities in methanolic extracts of *Lippie nodiflora* Linn. Die pharmazie 2004; 59(4):329-30.
  7. Durairaj Ashok Kumar, Vaiyapuri Tamilselvan, Upal kanti Mazumdar and Malaya Gupta. Antioxidant and free radical scavenging effects of *Lippia nodiflora*. Pharmaceutical biology. 2009;1(1):85-91
  8. Bhakuni DS, Dhar ML, Dhar MM, Dhawan BMN and Mehrotra BN. Screening of Indian plants for biological activity part II Indian journal of Experimental Biology,1969;7:250
  9. Gopal RH, Balkrishna K, Vasanth S, Bhima Rao B: Activity of *lippia nodiflora* essential oil on bacteria. Semin research in ayurveda and Siddha. New delhi;CCRAS:1995
  10. Pirzada AJ, Iqbal P, Shaikh W, Kazi TG and Ghani KV: Studies on the elemental composition and antifungal activity of medicinal plant *L. nodiflora* against skin fungi . J of Pak Assoc.Derma, 2005; 15(2):113-118.
  11. Dubois M, Gilles K, Hamilton JK, Rebers PA and Smith F .A colorimetric method for the determination of sugar. *Nature*. 1951: 167-168
  12. Lowery OH, Rosebrough NJ, Farr AL and Randall RJ .Protein measurement with the Folin phenol reagent. *J.Biol.Chem*. 1951: 193 265-275
  13. Jayaraman J Laboratory Manual in Biochemistry. New Delhi: Wiley Eastern Limited, New Delhi 1981.
  14. Bray HG and Thrope WV Analysis of phenolic compounds of interest in metabolism. *Meth. Biochem. Anal*.1954 (1) 27-52.
  15. Roe JH and Kuether CA .The determination of ascorbic acid in hole blood and urine through 2, 4- dinitrophenyl hydrazine derivative of dehydro ascorbic acid. *J. Biol. Chem*. 1943:147:399.
  16. Lee YP and Takahashi T. An improved colorimetric determination of amino acid with the use of ninhydrin. *Anal.Biochem* 1966. 14: 71-77
  17. Kaufman PB, Duke JA, Briclman H, Cseke S and Warber S Natural Products From plants. CRC Press Boca Raton, F.L 1999
  18. Freeze H Disorders in Protein Glycosylation and Protein Therapy. *The Journal of Pediatrics*. 1998: 133(5) 543-600
  19. Yadav PR and Tyagi R Lipid Biotechnology. 1 Discovery Publishing House-New Delhi.89. 2006.
  20. Nicholas The function and metabolism of ascorbic acid in plants. *Annals of Botany*. 1996 78: 661-669
  21. Thomsen S, Handen HS and Nyman V Ribosome inhibiting proteins from in vitro cultures of *Phytolacea dodecandra*. *Planta. Med*. 1991: (57) 232-236.
  22. Vijayvergia R and Kumar J .Quantificaton of primary metabolites of *Nerium indicum* Mill. *Asian J. Exp. Sci*. 2007: 21(1): 123-128.

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