MICROSCOPIC ILLUSTRATION OF PELARGONIUM X HORTORUM (GERANIACEAE)

ABSTRACT: For identification of a crude drug, there are several parameters which standardize 4 it for sure. Microscopic features describe a crude drug very well. Chances of adulteration are 5 very common due to morphological similarities in different species of drugs and to avoid such 6 7 confusions, standardization via microscopy helps to create a valuable profile of a given crude drug. Involving different parts of plant drug in microscopy viz leaves, stem, roots and flower etc. 8 helps a lot in identifying the original drug. Transverse section of different parts of plant, powder 9 microscopy and determination of leaf constants like stomatal number, stomatal index, vein islet 10 number, vein termination number and palisade ratio of Pelargonium x hortorum describes the 11

KEYWORDS: Pelargonium, Identification, Microscopic evaluation.

basic features of the drug and authenticate it as the original one.

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INTRODUCTION:

Pelargonium <u>x</u> hortorum <u>L.H. Bailey</u> is a species of <u>the genus Pelargonium</u> commonly used as an ornamental plant. Pelargonium P.x hortorum species is probably a <u>hybrib hybrid species</u> between Pelargonium P. zonale and Pelargonium P. inquinans belonging to family Geraniaceae. Plant can be propagated by stem cuttings and requires peaty or loamy soil and flourishes in sunny conditions [1]. The plant varies from height of 45 to 50



Fig.1a: General view of the 1Leaves of P. x-hortorum

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cm with fragrant green colored decorative leaves up_to 5 to 7.5 cm in length with reticulate venations and crenate margin. Flowers appears in many colors like red, pink, orange or white having five petals positioned around the center as ball shaped clusters. The inflorescence is long rigid peduncle. Generally tap root system is present in the plant.



Fig.1b: Flowers of *P.x_hortorum*

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Different species of *Pelargonium* are available abundantly in nature with immense

pharmacological potential and exhibit antifungal [2], mosquito repellent [3], anxiolytic, 22

antidepressant [4] and pediculicidal activities [5]. *Pelargonium* derived essential oils (citronellol,

geraniol, p-menthone and α -pinene etc.) are extensively used in perfumery, cosmetics, soaps,

creams and aromatherapy products [6-7]. The aim of this study is to standardize the particular 25

species of Geranium i.e G.hortorum and as such no microscopic profile has been available of 26

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MATERIAL AND METHODS:

Collection of plant: 29

The plant is collected in the month of January 2019 from Govt. College of Pharmacy Rohru, 30

31 Distt. Shimla, Himachal Pradesh, India and the collected samples were subjected to microscopic

examination. 32

Microscopy: Anatomical sections of the fresh leaf, petiole, stem and roots were prepared for the 33

microscopic studies and examined under Trinocular microscope Olympus-CH-20i model and

compound microscope. 35

36 For determination of leaf constants like stomatal number, stomatal index, vein islet number, vein

termination number and palisade parenchyma ratio camera lucida-was used Camera lucida-

Stomatal number and index determination: The fragment of leaf was cleared by boiling with 38

chloral hydrate solution. Epidermal layer was then peeled out using forcep. A square of 1mm 39

was drawn on a drawing paper using Camera lucida and stomata were counted and stomatal 40

41 index was calculated using formula-

Stomatal index (S.I.) = $S/E+S \times 100$

Comment [SS2]: Fig 1a and Fig 1b don't go in

Comment [SS1]: what is the citation of this sentence ??

Comment [SS3]: Where did this species come from? This species is no the world.

Comment [SS4]: The aim of this study, is

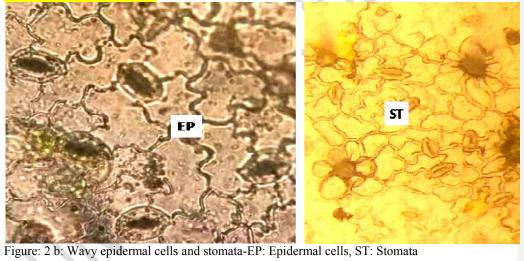
insufficient.

Where S= Number of stomata, E= Number of epidermal cells

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43	Determination of Vein-islet and termination number: Fragments of leaf was cut in 2 mm x 2 mm
44	rectangular shape and boiled in chloral hydrate solution followed by dilute hydrochloric acid for
45	few minutes. A square of 1mm was drawn on a drawing paper using Camera lucida and veir
46	islets and terminations were counted.
47	Determination of palisade ratio: Fragments of leaf was cut in 2 mm x 2 mm rectangular shape
48	and boiled in chloral hydrate solution followed by dilute hydrochloric acid for few minutes. A
49	square of 1mm was drawn on a drawing paper using Camera lucida and palisade cells were
50	focused underlying four epidermal cells [8-9].
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56	RESULTS:
57	The microscopic examination of the plant consists of its transverse section of leaf, petiole, stem
58	and root. The results of the T.S, powder characteristics and leaf constants are given in the Figure
59	2(a-h).
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Figure: 2 a: Trichomes- CTR: Unicellular, uniseriate covering trichome, GTR: Unicellular, uniseriate Glandular trichome



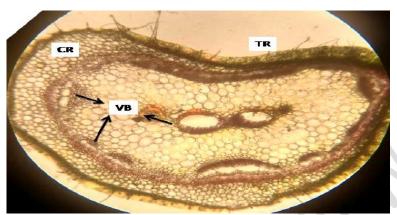


Figure: 2 c: Transverse section of Petiole-VB: Vascular bundles, CR: Cortex, TR: Trichome

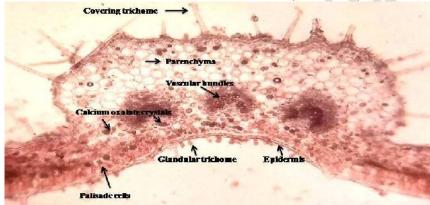


Figure: 2 d: Transverse section of leaf

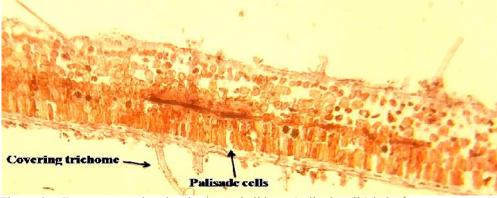


Figure: 2 e: Transverse section showing mesophyll layer (palisade cells) in leaf

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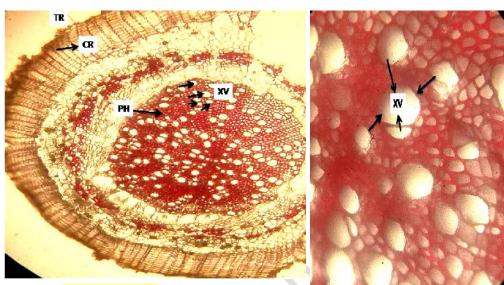


Figure: 2 f: Transverse section of root-TR: Trichome, CR: Cortex, PH: Pith, XV: Xylem vessles

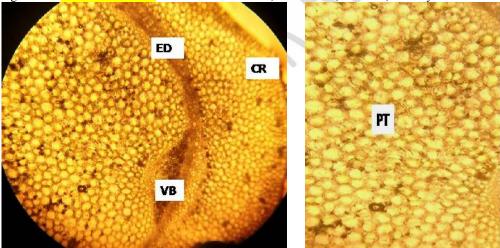
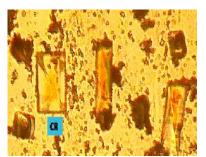
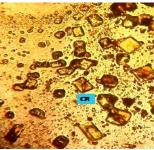


Figure: 2 g:Transverse section of stem-ED: Endodermis, VB: Vascular bundle, CR: Cortex, PT: Pith





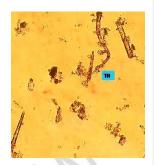


Figure: 2 h: Powder microscopy of leaf-CR: Crystals, TR: Trichomes

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Conclusion:

The macroscopic study reveals the physical characteristics of plant whereas the microscopic studies give us vital information about the histological arrangement of different plant parts. Transverse section of leaf showed the presence of unicellular covering, glandular trichomes, anomocytic stomata, mesophyll type is dorsiventral bilayered rectangular cylindrical arrangement of palisade parenchyma cells, wavy walled epidermal cells, five to six layered parenchymatous tissue, few calcium oxalate crystals and vascular bundles. In petiole five to ten celled layered cortex are present with circularly arranged vascular bundles in parenchymatous tissue. Stem portion-cross-section-showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles and pith. Roots cross-section-showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles and pith. Roots cross-section-showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles and pith. Roots cross-section-showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles and pith. Roots cross-section-showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles and pith. Roots cross-section-showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles are section showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles are section showed presence of endodermis with 15-20 layered cortex, circularly arranged vascular bundles are section showed presence of endodermis with 15

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Comment [SS7]: How many layers of endodermis?

Comment [SS8]: The roots do not contain covering trichomes. This is scientifically a big mistake. Comment the root section well.

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