



Original Research Article

Exploring the diuretic activity of Bisehri booti (*Aerva lanata*): an efficacious medication for nephrological disorders.Suhail Ahmad^{1*}, Mohammad Zaki Ahmad², Mohd. Wajeehul Qamar³, KM Yusuf Amin⁴, JA Ansari⁵¹Department of Ilmul Saidala, Sufia Unani Medical College, Hospital & Research Centre, East Champaran, Bihar, India.²Department of Ilaj Bit Tadbeer, University College of Unani, Tonk, Rajasthan, India.³Department of Ilmul Advia, Sufia Unani Medical College Hospital and Research Centre, East Champaran, Bihar, India.⁴Department of Ilmul Advia, Aligarh Muslim University, Aligarh, UP, India.⁵Department of Pharmacy Practice, Faculty of Pharmacy, Integral University, Lucknow, India.

ARTICLE INFO	ABSTRACT
<p>Article History</p> <p>Received : 15-Mar-2023 Revised : 05-Apr-2023 Accepted : 15-Apr-2023</p>	<p>Aim and Objective: The Bisehri booti (<i>Aerva lanata</i>) is an effective traditional medicine used for the treatment of nephrological disorders. This study aimed to investigate the diuretic activity of Bisehri booti by measuring urinary volume and sodium in albino rats.</p> <p>Methods: Four groups of five animals each were administered normal saline, hydrochlorothiazide, aqueous extract of Bisehri booti, and juice of Bisehri booti, respectively. Urine volume and sodium concentrations were measured to evaluate the diuretic activity of Bisehri booti.</p> <p>Results: Urine volume and sodium concentrations were significantly increased in the hydrochlorothiazide group and the groups administered with aqueous extract and juice of Bisehri booti, compared to the control group. The study showed that Bisehri booti has moderate diuretic activity, with the juice being more active than the extract. The finding was at variance with the demonstration of steroidal activity in the test drug in the present study. However, it is possible that the test plant possesses two or more active principles having steroidal and diuretic activity, respectively, making it a better drug for renal ailments as it would exert steroidal activity without the water and sodium retention, which is an undesired effect of steroidal drugs.</p> <p>Conclusion: The study provides scientific evidence supporting the traditional use of Bisehri booti as a diuretic agent. Further studies are needed to identify the active principles responsible for the observed diuretic activity and to investigate the potential of Bisehri booti in the treatment of renal ailments.</p>
<p>Key words</p> <p>Bisehri booti Aerva lanata Diuretic</p>	
<p>NonCommercial-ShareAlike 4.0 International License (CC BY-NC-SA)</p>	
	<p>*Author for Correspondence: suhailfargaleet@gmail.com</p> <p>DOI: https://doi.org/10.5281/zenodo.7893857</p>

Copyright © 2023 Biomedjournal Privacy Policy. All rights reserved.

INTRODUCTION

The practice of employing folk drugs or ethnic drugs is generally perceived as encompassing the use of drugs among tribal and rural communities as well as those utilized in traditional systems of medicine such as Tibb-e-Unani and Ayurveda, as these drugs are derived from herbs. However, this viewpoint is inadequate because Tibb-e-Unani and other traditional systems of medicine are complex systems with a systematic pharmacology based on comprehensive physiology and pathology, which cannot be equated with unspecialized, incomplete, and unorganized drug information common among forest-dwelling and rural communities, just because most traditional medicine drugs are derived from herbs. Some drugs are borderline in that they are widely used among rural communities but receive only sporadic mention in traditional medicine texts. Scientific research on such drugs could not only identify valuable agents but also improve traditional systems of medicine by shifting peripheral drugs to the mainstream. Bisehri Booti

34 (*Aerva lanata*) is frequently used by rural communities 88
 35 in India and Sri Lanka, but only a few Tibb-e-Unani 89
 36 physicians mention it [1]. It is used as a diuretic and 90
 37 anthelmintic agent and is employed to treat ailments 91
 38 like headaches, diarrhea, sore throat, and coughs [2-4]. 92
 39 A few Unani physicians have found it to be useful in 93
 40 treating conditions like albuminuria, haematuria, renal 94
 41 and vesicular calculi, and prostatitis, but its most 95
 42 important medicinal use appears to be in conditions 96
 43 similar to nephrotic syndrome [5-6]. Bisehri Booti was 97
 44 tested for its anti-inflammatory, analgesic, steroidal, 98
 45 diuretic, and anthelmintic activity, as it is also used by 99
 46 common people to treat headaches and worm 100
 47 infestations [7]. Udupihille et al. (1986) conducted a 101
 48 systematic study on various parts of the plant to 102
 49 determine their diuretic activity, and they found that 103
 50 the fresh flowers of the plant have the strongest 104
 51 diuretic activity. According to reports from 105
 52 experimental studies, the plant produces a hypotensive 106
 53 effect, but there are conflicting reports about its cardiac 107
 54 effect, and it is claimed to have negative chronotropic 108
 55 and extra systole at a dose level of 50 mg/kg. In the 109
 56 hypodynamic frog heart preparation, a level of 1-3 110
 57 mg/ml of perfusion fluid, the plant produced a positive 111
 58 inotropic effect, as reported by Tripathi et al. (1990) [8- 112
 59 9].

60 In the light of above considerations Bisehri booti (*A.* 113
 61 *lanata*) was tested for its diuretic and activity which is 114
 62 done by the Measurement of Urinary Volume and 115
 63 Sodium in albino rats. Since the Unani physician use the 116
 64 Juice of whole plant, therefore, it was used in the 117
 65 present study also. There are obvious difficulties in the 118
 66 use of the Juice of the plant, therefore, the aqueous 119
 67 extract of the whole plant was also studied with the 120
 68 view of comparing its effect with the Juice and thereby 121
 69 identifying the extract as a more convenient 122
 70 formulation if it was found to be equipotent with the 123
 71 Juice. The Unani clinical dose of Bisehri booti (*A. lanata*) 124
 72 In 20 mg / kg of the Juice it was converted into the 125
 73 corresponding rat dose by multiplication with 7. [10] 126
 74 The value thus obtained was rounded of to 15 mg / 100 127
 75 g of Juice and this was used as one of the doses in study. 128
 76 It is widely felt that the doses reported in Unani texts 129
 77 are generally on the lower side and this is one of the 130
 78 reasons for the failure of therapy with these drugs, 131
 79 therefore, a slightly higher dose that is 20 mg / 100 g of 132
 80 the juice was also studied. The aqueous extract yield 133
 81 was 1/10th of the crude drug W/W. Since the juice yield 134
 82 is also nearly of the same order W/W, therefore, the 135
 83 aqueous extract of Bisehri Booti (*A. lanata*) was also 136
 84 studied at the same doses as that of the juice 15 mg / 137
 85 100g and 20 mg /100 g in albino rats. 138

86
87

Aerva lanata is a perennial, herbaceous weed that 139
 140 groweth in a suberect diffuse or prostrate to erect 141
 142 manner. It is frequently found in waste lands as well as 142
 143 crop fields, spanning throughout the tropical regions of 143
 144 Asia and Africa [1, 12]. In India, it is distributed widely 144
 145 in hotter regions reaching an altitude of up to 900 145
 146 meters on hills, as well as the plains of Bengal, 146
 147 extending westward to Konkan, Madhya Pradesh, and 147
 148 also present in the Punjab and trair region of UP [13, 14]. 148
 149 The plant is also commonly located in the Philippine 149
 150 region [15, 16], and distributed in Arabia, Tropical 150
 151 Africa, and Java [7]. The season of its flowering 151
 152 spanneth from the months of October to November, 152
 153 and continued until June [1, 17].

The plant is usually pubescent or villous and can grow 154
 155 to a height of 1-4 feet, with a base that is often woody. 155
 156 In old plants, branches grow longer and arise from 156
 157 either the stem or the root stock [1, 18].

In microscopic view, the hairs of the plant are arranged 157
 158 in a single row. The epidermis is made up of a single 158
 159 layer with a thin cuticle, which is more pronounced at 159
 160 the ridges. The epidermal cells are square-shaped, thin- 160
 161 walled, and lacking in lignin. The cells in the ridges are 161
 162 thicker-walled. The trichomes are elongated and 162
 163 composed of interlocking cells, which are thick and 163
 164 have pointed outgrowths. The basal cells of trichomes 164
 165 have yellow walls. The cortex is composed of 165
 166 isodiametric parenchyma with intracellular spaces and 166
 167 is 5-7 cells deep in the furrows. The primary vascular 167
 168 bundles in the young stem are numerous, small, 168
 169 collateral and open within inner phloem arranged in a 169
 170 ring typical of dicot stem. The stem shows secondary 170
 171 growth, which becomes abnormal later due to the 171
 172 production of conjunctive tissue, wherein phloem and 172
 173 xylem elements are embedded. The anomalous 173
 174 secondary growth of typical *Amaranthus* type is very 174
 175 clear. In the old stem, four rings of secondary tissues are 175
 176 found. The pith is larger in the young stem composed of 176
 177 thin-walled parenchyma and reduced in old stem due 177
 178 to increase in the amount of second xylem. Prismatic 178
 179 calcium oxalate crystals are also found in this region. 179
 180 The epidermal cells of the leaf are irregular in shape 180
 181 and have anomocytic type of stomata. The leaf has 181
 182 more trichomes than the stem, which have the same 182
 183 structure as those in the stem. The transparent 183
 184 prismatic nature of calcium oxalate crystals is also 184
 185 found in the leaf, but in lesser numbers [6, 19, 20].

The plant *A. lanata* has been studied for its chemical 186
 187 composition, revealing the presence of several 187
 188 compounds including Beta-sitosteryle palmitate, Alpha- 188
 189 amyirin, and Beta-sitosterol [21]. In addition, inorganic 189
 190 salts, free sugars, tannins, flavonoids, and hexaeosinol 190
 191 have also been identified [22]. Further research by 191
 192 Chandra in 1990 resulted in the isolation of new

142 constituents including hentriacontane, Beta-sitosterol, 195
 143 Alpha-amyrin, and botulin [23]. In 1991, four 196
 144 flavonoids, two feruloylamides, and other phenolic 197
 145 compounds were also isolated and their structures 198
 146 identified [24]. Six alkaloids were also discovered, 199
 147 including two previously known ones, Canthin-6-one 200
 148 and Beta-corbolin-1-propionic acid, and four new ones: 201
 149 10-methoxy-canthin-6-one, 10-hydroxy canthin-6-one, 202
 150 10-O- β -D-glucopyrano syloxycanthine-6-one, and 6- 203
 151 methoxy- β -corbolin-1-propionic acid [5]. 204

152 The aforementioned alkaloids, excluding the second 205
 153 and sixth ones, were obtained from the roots of the 206
 154 same plant using similar methods. The first four 207
 155 alkaloids were isolated from the chloroform extract, 208
 156 whereas the fifth and sixth ones were obtained from 209
 157 the ethyl acetate extract. The plant tested positive for 210
 158 cellulose, lignin, sterol, cutin, subrin, protein, tannin, 211
 159 and saponin, while starch and fat were not present. The 212
 160 petroleum extract of the dried plant contained 213
 161 terpenes, alkaloids, and phytosterol, which were likely 214
 162 beta-sitosterol, stigmasterol, and two compounds 215
 163 identified as possibly kaempferol-3-galactoside, and 216
 164 kaempferol-3-rhamnogalactoside and kaempferol. The 217
 165 free sugars detected in this plant were fructose, 218
 166 galactose, sucrose, and rhamnose [6, 19]. 219

167 FTIR analysis of the different parts of the plant, namely 220
 168 the roots, stems, leaves, and flowers, showed the 221
 169 presence of several functional groups like amides, 222
 170 alcohols, carboxylic acids, nitro compounds, amines, 223
 171 and alkyl halides. The leaves of the plant are rich in 224
 172 minerals such as K, Na, Ca, Mg, Zn, Fe, and Mn, while 225
 173 the entire plant contains essential trace elements such 226
 174 as calcium, silicon, magnesium, potassium, chloride, 227
 175 carbon, and oxygen. Gallic acid is present in significant 228
 176 quantities in the roots, and the roots extract also 229
 177 contains quinones, phenols, triterpenoids, phytosterols, 230
 178 and phlobatannins. The stem's aqueous extract revealed 231
 179 the presence of gallic acid, apigenin-7-O-glucoside, 232
 180 quercetin-3-O-rutinoside, and myricetin upon 233
 181 phytochemical screening. The GC-MS analysis of 234
 182 different parts of the plant such as leaves, stems, roots, 235
 183 flowers, and seeds showed the existence of a variety of 236
 184 compounds like pyridine, hydroquinone monobenzyl 237
 185 ether, docosane, dotriacontane, (R,Z)-12-hydroxy-9- 238
 186 octadecenoic acid, 2-isopropyl-2,5-dihydrofuran and 239
 187 more [25-31]. 240

188 Medicinal Uses

189 Unani Medicine: In his tome titled "Mujarrabat-e-
 190 Qadri", Hakeem Abdul Qadir (1931) hath commended 242
 191 the usage of Bisehri-Booti to alleviate renal and 243
 192 vesicular calculi.[5] Certain reputable practitioners of 244
 193 Tibb-e-Unani in Aligarh and Amroha hath employed 245
 194 this plant with efficacious outcomes in treating other

maladies, such as albuminuria, haematuria, and
 prostatitis. The root is purported to possess demulcent
 diuretic properties and is deemed beneficial in treating
 strangury according to Ayurveda [7, 32] The entire herb
 is considered Sitayam and is utilized in Rikta-pita
 (Haematamasis) and Diabetes [13].

The usage of *A. lanata* is widespread in Sri Lanka to
 alleviate symptoms of indigestion and coughing [33]. In
 the state of Bihar, this plant is employed to treat white
 urine, diarrhoea, cholera, dysentery, and even snake
 bites [2]. The extract of the plant is believed to have
 anthelmintic and diuretic properties [3, 34].
 Additionally, it is claimed that the plant possesses
 refrigerant, diuretic, sodiferic, stimulant, and vermifuge
 properties.[35] The flowers or flowering tops of the
 plant have been reported to be effective in treating
 headaches [17, 36]. Furthermore, the juice extracted
 from the root is applied on the forehead to provide
 relief from headaches.

Tripathi and colleagues conducted a study to
 investigate the impact of an alcoholic extract of *A.*
lanata on the blood pressure and ECG of anaesthetized
 rats, as well as on isolated frog heart preparations. They
 found that at a dose of 10mg/kg, the extract reduced
 blood pressure without affecting the ECG pattern,
 whereas higher doses (50mg/kg) produced negative
 chronotropic effects and extrasystoles. Additionally, at
 doses of 1-3mg/ml of perfusion fluid, the extract had an
 inotropic effect on frog heart preparations.
 Furthermore, the researchers tested various parts of the
 plant, including the flower, leaf, stem, and root, in raw
 and dried forms, for their diuretic properties. The study
 found that fresh plant extracts significantly increased
 urine output, and the extract obtained from fresh
 flowers induced the most significant diuresis. Raw
 flower extract was found to be the most effective in
 producing diuresis compared to other parts of the raw
 plant and the dried form. The study suggests that *A.*
lanata has a potent diuretic activity, with dried *A. lanata*
 having a moderate diuretic activity, possibly due to the
 destruction or partial transformation of the diuretic
 factor during the drying process. Furthermore, the
 flowers of the plant contained a higher concentration of
 the diuretic factor compared to the leaves, roots, and
 stem, which had a diminishing diuretic effect in
 descending order [8, 9].

241 MATERIAL AND METHODS

The test drug namely Bisehri booti (*A. lanata*) was
 obtained from the herbarium of Department of Ilmul
 Advia, A.M.U., Aligarh, after proper identification in the
 light of botanical information. The test drug was used

246 for experimental study in two forms, i.e. aqueous
247 extract and fresh plant juice.

248 Extract: The drug was collected in the month of
249 October, dried in air and powdered. The powder then
250 was extracted for 6 hours, in distilled water. The extract
251 was filtered and the solvent was evaporated on water
252 bath. The extract obtained was 1/10th of the air dried
253 plant.

254 Juice: The Juice was made from fresh plant at the time
255 of experiment; the fresh plants were collected, crushed
256 and squeezed with the help of a special squeezer.

257 Study for Diuretic Activity

258 The test for diuretic activity was carried out by the
259 method of Taylor et al. (1962) modified by us [37].
260 Albino rats of approximately the same weight were
261 divided into 4 groups of 5 animals each. The animals in
262 Group I serving as control fed with 3.0 ml of normal
263 were saline. The animals in Group II serving as the
264 standard were administered with standard drug viz.
265 0.5mg/100gm of Hydrochlorothiazide, by oral route. The
266 animals in Group III and IV were administered with
267 20mg/100gm of juice and the aqueous extract of the
268 plant dissolved in distilled water, respectively, by oral
269 route. The animals in Group II, III, and IV were also
270 administered NaCl solution of a strength and in a
271 volume so that they received water and NaCl equal to
272 3.00 ml of normal saline given to standard and control
273 animals. All the animals were immediately placed,
274 individually, in metabolic cages and the urine passed
275 over 6 hours was collected. The volume of the urine
276 was measured and the concentration of sodium
277 estimated by Flame Photometry. The volume and
278 sodium content of the animals of various groups was
279 the urine statistically compared and analysed by
280 Student's "t" Test.

281 RESULTS

282 The mean urine volume measured over 6 hours in the
283 group treated with the Juice of *A. lanata* was 1.675 ml
284 which is 3 times more than in the control group i.e.
285 0.575 ml and sodium content of this group was
286 2058.312 ± 95.62 ppm, which is also higher than the
287 urinary sodium of control group i.e. 1816.648 ± 190.63
288 ppm ($p > 0.025$). The mean urine volume in group
289 treated with extract of *A. lanata* amounted to 1.4 ml
290 and the sodium content of urine was 1974.98 ± 170.84
291 ppm which is also higher than the urinary sodium of
292 the control group ($p < 0.025$). The urine volume in the
293 standard group treated with hydrochlorothiazide (0.5
294 mg / 100 g) was 4.3 ml ($p < 0.0010$) and sodium content
295 of urine was 2291.643 ± 164.638 ($p < 0.010$). The results
296 are presented in table no. 1 & 2 and figure no 1 & 2.

Table No. 1: Effect of bisehri booti (*A. lanata*) on urine output.

Group	Urine passed
Normal Saline	0.575 ± 0.147
Hydrochlorothiazide (0.5mg / 100g)	4.3 ± 0.61**
Extract (20mg /100g)	1.4 ± 0.24*
Juice (20mg /100g)	1.675 ± 0.23*

(mean ± S.E.M.), n=5, * = $p < 0.025$, ** = $p < 0.0005$.

Table No. 2: Effect of Bisehri booti (*A. lanata*) on sodium excretion.

Groups	Sodium excreted
Normal Saline	1816.648 ± 190.63
Hydrochlorothiazide (0.5mg/ 100g)	2291.643 ± 164.638**
Extract (20mg /100g)	1974.98 ± 170.84*
Juice (20mg /100g)	2058.312 ± 95.62*

(mean ± S.E.M.), n = 5, * = $p < 0.025$, ** = $p < 0.0005$

DISCUSSION

The aqueous extract and the juice of Bisehri Booti (*A. lanata*) were studied, each at the dose 15 mg / 100 g and 20 mg / 100 g, in albino rats. The test drug was studied for diuretic activity by the measurement of urinary volume and sodium in albino rats.

Since the test drug is reported to be effective in renal disorders it was thought that it may possess diuretic activity also. Therefore, it tested for diuretic was activity by the method of Taylor et. al. (1962). Albino rats of either sex was divided into 4 groups of 5 animals each. The animals in Group I were fed with normal saline and the animals in Group II were administered with standard drug viz. 0.5 mg/100 g hydrochlorothiazide by oral route. The animals in group III and IV were administered with 20 mg/100 gm of aqueous extract and juice of the Bisehri booti (*A. lanata*), respectively. The animal was placed in metabolic cages immediately and volume of the urine passed over six hours collected and measured. The sodium concentration was estimated by flame photometry. In the control animals the urine volume and Sodium were found to be 0.575 ± 0.147 ml and 1816.648 ± 190.63 ppm, while in the standard animals treated with hydrochlorothiazide they were significantly increased to 4.3 ± 0.61 ml and 2291.643 ± 164.638 ppm ($p < 0.0005$), respectively. In the animals treated with aqueous extract and the juice, these parameters were found to be significantly increased to 1.4 ± 0.24 ml and 1974.98 ± 170.84 ppm and 1.675 ± 0.23 ml and 2058.312 ± 95.62 pmm ($p < 0.025$), respectively. Therefore, the study shows that Bisehri booti possesses moderate diuretic activity with the

336 juice being more active than the extract. This finding is 384
 337 at variance with the demonstration of steroidal activity 385
 338 in the test drug in present study. However, it is possible 386
 339 that the test plant possesses two or more active 387
 340 principles having steroidal and diuretic activity, 388
 341 respectively. If this is the case Bisehri Booti will be a 389
 342 better drug for renal ailments as it would exert
 343 steroidal activity without the water and sodium 390
 344 retention which is an undesired effect of steroidal 391
 345 drugs. Since steroidal agents have metabolic effects also 392
 346 therefore due to the demonstration of steroidal activity. 393

347 CONCLUSION 394

348 In conclusion, this study examined the diuretic activity 395
 349 of the aqueous extract and juice of Bisehri Booti (*A. 396*
 350 *lanata*) in albino rats at doses of 15 mg / 100 g and 20 397
 351 mg / 100 g. The results showed that both the extract 398
 352 and juice increased urine volume and sodium 399
 353 concentration, indicating moderate diuretic activity. 400
 354 Interestingly, the juice was found to be more effective 401
 355 than the extract. While this finding is inconsistent with 402
 356 the previously demonstrated steroidal activity of the 403
 357 plant, it is possible that Bisehri Booti contains multiple 404
 358 active principles with both steroidal and diuretic 405
 359 effects. If this is the case, Bisehri Booti could be a 406
 360 superior treatment option for renal disorders as it 407
 361 would provide the benefits of steroidal activity without 408
 362 the undesirable effects of water and sodium retention. 409
 363 This study provides further support for the potential 410
 364 medicinal uses of Bisehri Booti in traditional medicine. 411

365 CONFLICT OF INTEREST 412

366 None declared. 413

367 REFERENCES 414

368 1. Trimien H: A Hand Book to the Flora of Cylon. 417
 369 Part. III. M / S Bishen Singh Mahendra Pal Singh, 418
 370 New Cannought Place, Dehli, India, 402-403, 419
 371 1974.
 372 2. Jain SK, Trafdar CR.: Medicinal Plant - lore of 420
 373 Santals (A review of P.O. Bodding 's work). Eco 421
 374 Bot. 1970;24:244. 422
 375 3. Asolker LV, Kakkar KK, Chakre J: Second 423
 376 Supplement to Glossary of Indian Medicinal 424
 377 Plants with Active Principles . Part-1. 425
 378 Publication & Information Directorate (C.S.I.R.), 426
 379 New Delhi, India p.27, 1992. 427
 380 4. Nadkarni KM: Indian Mataria Medica. Popular 428
 381 Book Depot, Bombay, India, p.49, 1976. 429
 382 5. Qadir MA: Mujarrabat-E-Qadri, Mohan Printing 430
 383 Press, Aligarh, India, p. 207, 1930. 431
 432

6. Afaq SH, Tajuddin, Afridi RM: Bisehri Booti (*A. 433*
lanata): Some lesser known uses and 434
 435 pharmaconosy. Ethnobotany. 1991;3:91-94.
 7. Kirtikar KR, Basu BD: Indian Medicinal Plant 436
 Vol. IIIrd. International Book Distributors, 437
 Rajpur, Dehra Dun, India, p. 206, 1987.
 8. Tripathi HC, Chandra S, Lal J. Cardiovascular 438
 Effect of Aerva lanata. Indian J Pharm. 1985;17:
 439 42.
 9. Udupihille M, Jiffry M: Diuretic Effect of Aerva 440
 lanata with Water, Normal Saline and Corander 441
 as Control, Indian J Physiol Pharmacol. 1986;
 442 30: 91-97.
 10. Dhawan BN, Duley MP, Mehrotra BN, Rastogi 443
 RP, Tandon JS: Screening of Indian Plants for 444
 Biological Activity. Part IX. Indian J Exp Biol. 445
 1980;18: 594.
 11. [https://plants.usda.gov/home/classification/556](https://plants.usda.gov/home/classification/55659)
 446 [59](https://plants.usda.gov/home/classification/55659) [Accessed on Mar 2023].
 12. Collett H: Flora Simlensis, A Hand Book of the 447
 Flowering Plants of Simla and the 448
 Neighbourhood. Calcutta; Simla. Thacker, 449
 Spink; Co. India, p.414, 1921.
 13. Karnick CR: Some Aspects of Crude Indian Drug 450
 Plants in Ayurvedic System of Medicine for 451
 Madhumya (Diatbetes) Actaphytotherapeutics. 452
 1972;8:141-147.
 14. Kapoor SL, Kapoor LD: on the Botany and 453
 Distribution of "Pashanbhed" Sachitra 454
 Ayurveda. 1976; 769-791.
 15. Duthie JF : Flora of Upper Gangotic Plain and of 455
 the Adjacent Siwalik and Sub Himalyan Tract 456
 (1960) Vol.1 , Part 1. Bishen Singh Mahendra 457
 Pal Singh. Delhi, India.
 16. Mary TN, Prabha SA: Geology of *A. lanata* Juss. 458
 Geobios. 1980;7(6):285-86.
 17. Patnaik H: Some Medicinal Plants Around 459
 Cuttak. J Bomb Nat Hist Soc. 1956;54(1)140-
 460 152.
 18. Thiselton-Dyre; WT: Flora of Tropical Africa 461
 Vol.VI. Published Under the Authority of the 462
 State for the Colonies. L. Reeve & Co. Ltd. The 463
 Oast House, Brook Ashfort, Ken, England,
 1913:39-40.
 19. Afridi RM: Pharmacognostical Studies of certain 464
 Unani Medicinal Plants. M.D. Thesis. 1992.
 Dept. of Ilmul Advia, AMU, Aligarh, India.
 20. Bamber CJ: Plants of Punjab, A Descriptive Key 465
 to the Flora of the Punjab, North-west, Frontier

- 433 Province and Kashmir. M / S Periodical Experts.482
 434 Delhi, India, p. 315, 1976. 483
- 435 21. Aiyer VN, Narayan V, Sheshadri TR, 484
 436 Vaidyasaran: Chemical Components of Some 485
 437 Indian Medicinal Plants. Indian J Chem. 1973;486
 438 11:89-90. 487
- 439 22. Sastry CSP, Siraj PG: Analytical Studies Leading 488
 440 to Standardisation of *Aerva lanata* Juss 489
 441 (Ayurvedic Medicine) Indian J Pharmacy. 1977;490
 442 6:174. 491
- 443 23. Yognarasimhan SN, Nair KV, Murthy KRK. 492
 444 Medico-Botany of Karnataka, Utilisation of 493
 445 Floristic of Kankapur Wealth for the Economic 494
 446 Development Taluk, Bangalore District. J Econ 495
 447 Texo Bot. 1985;6(1):97-107. 496
- 448 24. Gertruda Z, Vladimir K, Victor O, Anatoly M: 497
 449 Canthin - 6 - one and Beta corboline Alkaloids 498
 450 from *Aerva lanata*. Planta Medica. 499
 451 1991;58(2):192-196. 500
- 452 25. Yamunadevi M, Wesely EG, Johnson MA. FTIR 501
 453 spectroscopic studies on *Aerva lanata* (L.) Juss. 502
 454 Ex Schult. Asian J Pharm Clin Res. 2012;5:82- 503
 455 86.
- 456 26. Omoyeni OA, Adeyeye EI. Chemical 504
 457 composition, calcium, zinc and phytate 505
 458 interrelationships in *Aerva lanata* (Linn) Juss. 506
 459 ex Schult leaves. Orient J Chem. 2009; 25:485- 507
 460 488.
- 461 27. Ragavendran P, Arul Raj C, Sophia D, Starlin T, 508
 462 Gopalakrishnan VK. Elemental analysis of *Aerva 509*
 463 *lanata* (L.) by EDX method. Int Res J Pharm. 510
 464 2012;3:218-220.
- 465 28. Vijayalakshmi R, Ravindhran R. HPTLC method 511
 466 for quantitative determination of gallic acid in 512
 467 ethanolic root extract of *Diospyrus ferrea 513*
 468 (Willd.) Bakh and *Aerva lanata* (L.) Juss. Ex 514
 469 Schultes-A potent Indian medicinal plants. 515
 470 Asian J Pharm Clin Res. 2012;5:82-86.
- 471 29. Vijayalakshmi R, Ravindhran R. Preliminary 516
 472 comparative phytochemical screening of root 517
 473 extracts of *Diospyrus ferrea* (Willd.) Bakh and 518
 474 *Aerva lanata* (L.) Juss. Ex Schultes. Asian J Plant 519
 475 Sci Res. 2012; 2:581-587.
- 476 30. Kumar G, Karthik L, Bhaskara Rao KV. 520
 477 Phytochemical composition and in vitro 521
 478 antioxidant activity of aqueous extract of *Aerva 522*
 479 *lanata* (L.) Juss. ex Schult. Stem 523
 480 (Amaranthaceae). Asian Pac J Trop Med. 2013; 524
 481 6:180-187.
31. Mariswamy Y, Gnanaraj WE, Antonisamy JN, 525
 Adaikalam AA, Jamesraj V. GC-MS studies on 526
 methanolic extracts of *Aerva lanata* L. Indo 527
 American J Pharmaceut Res. 2013; 3:2687- 528
 2717.
32. Bedi SJ: Ethnobotany of Ratan Mahal Hills, 529
 Gujarat, India. Eco Bot. 1978;32 (3):278-84.
33. Trimen H: A Hand Book to the Flora of Ceylon. 530
 Part. III. M/s Bishen Singh Mahendra Pal Singh, 531
 New Cannought Place, Delhi, India, 402-403, 532
 1974.
34. Srivastava JG: "Pashanbhed" Quart. J. Crud. 533
 Drug Res. 1971;11(1):1683-1669.
35. Chandra S, Sastry MS: Chemical Constituents of 534
Aerva lanata . Fitotrophia. 1990;61(1):188.
36. Maheshwari JK: The Flora of Delhi. Publication 535
 & Information Directorate (C.S.I.R.), New Delhi, 536
 pp. 294, 1963.
37. Taylor RM, Topliss JG: Structure Activity 537
 Relationship of 3-Substituted 538
 Dihydrobenzothiazine Diuretics. J Med Pharma 539
 Chem. 1962;4:312.