






## Composition, structure and productivity of the herbaceous vegetation of five forest stands varying in soil moisture and nitrogen in Central Himalaya, India

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### Abstract

Herb layer contributes significantly to the species diversity of forest ecosystem and reacts rapidly to changes in the soil characteristics. Composition, structural parameters and productivity of the herbaceous vegetation of five forest stands in the Central Himalaya of India, situated at Nainital district of Uttarakhand, India were investigated. At each site, 1 ha plot was established and herbaceous vegetation was analysed by placing 30 random quadrats of 50x50 cm at each site. To determine the soil moisture and soil nitrogen, soil samples were collected from each site and analysed in the laboratory. Results showed that chosen sites contrasted in terms of soil moisture, total soil nitrogen, herbaceous diversity, and biomass. In this study, 70 genera, 73 species from 31 families of herbs were recorded. Total number of herbaceous species recorded in the stands varied from 28 (Pines and Hanumangarhi) to 50 (Takula). Total individuals of all herb species were 110, 114, 141, 164 and 188 plants m<sup>-2</sup> in Rusi (RU), Hanumangarhi (HG), Pines (PI), Barapatthar (BP) and Takula (TA) forest stands, respectively. Index of similarity and species turnover ranged from 34.48 to 62.50 and 0.31 to 0.54 respectively. Herb density, diversity and biomass showed positive correlation with soil moisture and soil nitrogen. These findings suggested that the soil moisture and nitrogen enhanced the herbaceous diversity by ameliorating soil conditions. Herb species such as *Bidens pilosa*, *Commelina benghalensis*, *Erigeron kavinskianus*, *Eupatorium adenophorum*, *Micromeria biflora*, *Oplismenus compositus*, *Oxalis corniculata*, *Strobilanthes angustifrons*, *Viola canescens*, *Vitis himalayana* were most benefitted species because they were present across all the forest stands in high density where as herb species like *Ajuga bracteosa*, *Anemone vitifolia*, *Arisaema tortuosum*, *Cassia occidentalis*, *Craniotome furcata*, *Dioscorea bupleuroides*, *Lepidagathis cristata*, *Sigesbeckia orientalis*, *Veronica beccabunga* and *Vicia villosa* were the most affected species because they were recorded from among one of the forest stand and they are present in very low density. This study also demonstrated a straight relationship between herbaceous diversity and biomass indicating the significance of species diversity for ample generation of biomass in forest ecosystem.

**Keywords:** Herbaceous vegetation; productivity; soil moisture; total soil nitrogen; Central Himalaya.

### Introduction

Among all kinds of vegetation types, herb layer has a pivotal role in maintaining the ecosystem equilibrium because it is mostly present in highest numbers comprising different kinds of genera and species (Gilliam, 2007; Jhariya et al. 2013; Parihaar et al. 2014; Khan et al. 2020a). This layer is also responsible for approximately 12% of the Gross Photosynthetic Production (GPP) of a forest ecosystem (Bargali and Bargali, 2000; Muller, 2003). In the forest ecosystem, the herb layer not only determines the spatio-temporal distribution but also affect dynamics of woody seedlings through regeneration. The herbaceous layer also regulates the recruitment of woody plants (Maguire and Forman, 1983) directly through competition for nutrients, light, water and indirectly through the addition of macro and micro

nutrients (San Jose and Farinas, 1991). The fibrous root system of herbs particularly grass is beneficial in binding soil particles and maintaining soil structure, thereby substantially reducing soil erosion and water loss (Sagar et al. 2008). The herbaceous vegetation influence nutrient cycling, primary production, energy flow in the forest ecosystems (Das et al. 2008), provide forage for domestic and wild animals, exhibit attraction for many butterflies due to high richness of nectar-bearing flowers (van Swaay, 2002) and provide shelter for microbial communities (Singh et al. 2006). Therefore, the interaction between these strata promotes the structural organization and number of niches (ecosystem complexity) and finally makes the system stable.

The role of herbaceous vegetation in terms of biomass production and nutrient cycling is not significantly appreciable, however, their role in ecosystem dynamics cannot be ignored (Bargali et al. 2015a Khan et al. 2020b). Their floristic role also transcends their varied ecological importance. Species diversity that is defined by the spatio-temporal alteration in species composition and their distribution (Gillet et al. 1999), govern stability and vulnerability of forest ecosystems, understanding the composition, distribution, and diversity of herbaceous vegetation is basic to the understanding of dynamics of the forest ecosystem. Though plant species diversity is affected by a variety of abiotic and biotic factors, herbaceous vegetation of any forest ecosystem is generally most affected by edaphic factors, climatic variables and livestock grazing which in return affect regeneration of trees and shrubs, resource availability and overall scenario of forest ecosystem in that region (Bushing and Brokaw, 2002; Schnitzer and Carson, 2001; Bargali et al. 2014 and 2015 a and b).

The rates of species gain or loss in the community are affected by disturbances such as physical resources, anthropogenic agencies, species interactions and propagules availability (Karki et al. 2016). Abiotic factors that have been considered many times for research purposes were soil moisture, nitrogen availability or related variables such as the water table. Plants utilize water for photosynthesis or respiration, and inadequate supply of water loses cell turgidity which leads to wilting and eventual death of plant. Water is a vital part of plant health that is why it has been chosen so many times in studies. Understanding the effects of change in soil moisture and total soil nitrogen on herbaceous vegetation will lead us to an improved knowledge of plant ecology. This study is an attempt to understand composition, structure and biomass production of herb layer in five forest stand of Central Himalaya in relation to soil moisture and nitrogen. The purpose of the present study was to describe the impacts of varying levels of soil moisture and nitrogen on species composition, phytosociological characters and biomass of the herbaceous vegetation in forest stands of Central Himalaya, India.

## Materials and methods

### Study area

This study was conducted in the Nainital Forest Division of Nainital district in Uttarakhand state, Central Himalaya, India. After conducting frequent survey in the region, five sites between 29°21'51"-29°23'21"N latitude and 79°26'31"-79°28'29"E longitude covering an altitudinal range of 1700 to 2200 m asl were selected (Table 1). All sites are within 5 km distance to each other, have similar topography, soil type, and experience similar climatic conditions and their respective vegetational scenario is given in table 2.

Table 1. Characteristics of the selected forest stands of Central Himalaya, India.

S. No.	Locality	Latitude	Longitude	Altitude (m asl)	Aspect studied
1	Rusi (RU)	29°21'57"N	79°27'23"E	1756	West
2	Hanumangarhi (HG)	29°21'55"N	79°27'32"E	1899	East
3	Pines (PI)	29°23'1"N	79°28'29"E	1902	West
4	Barapatthar (BP)	29°23'21"N	79°26'31"E	2134	South East
5	Takula (TA)	29°21'51"N	79°27'29"E	1798	North east

Table 2. Vegetation scenario of selected forest stands

S. No.	Locality	Dominating species (On the basis of IVI for tree and shrub and PV for herb) (Total species number)			Total woody vegetation cover (m <sup>2</sup> ha <sup>-1</sup> )
		Tree	Shrub	Herb	
1	Rusi (RU)	<i>P. roxburghii</i> (3)	<i>C. nepalensis</i> (3)	<i>E. karvinskianus</i> (30)	24.01
2	Hanumangarhi (HG)	<i>C. torulosa</i> (5)	<i>C. nepalensis</i> (5)	<i>O. compositus</i> (28)	20.13
3	Pines (PI)	<i>Q. leucotrichophora</i> (4)	<i>C. nepalensis</i> (7)	<i>E. karvinskianus</i> (28)	17.41
4	Barapatthar (BP)	<i>Q. leucotrichophora</i> (4)	<i>C. nepalensis</i> (10)	<i>C. dactylon</i> (35)	29.91
5	Takula (TA)	<i>C. torulosa</i> (7)	<i>C. nepalensis</i> (9)	<i>E. karvinskianus</i> (50)	25.27

## Climate

The climate of the study area is a typical temperate type. The climate is determined by the monsoon rhythms and the year can be divided into three main seasons: winter, usually cold and relatively dry (mid-December to February or sometimes mid-March); summer, warm and dry (April to mid-June); and a rainy season, which is warm and wet (mid-June to mid-September). The period of transition occur between summer and winter and between winter and summer are autumn (October to November) and spring (February to March), respectively. The rainy season accounts for about three-fourths (3/4) of the annual rainfall. During winters, some parts of the study area receive snowfall. The Mean minimum monthly temperature ranged from 7°C (January) to 21°C (August) and mean maximum monthly temperature varied from 13°C (January) to 30°C (September). Annual rainfall was 2200 mm. Average humidity fluctuated near the saturation point during the monsoon and was lower during summer, and ranged between 36.5% (May) to 88.7% (July) (Fig.1).

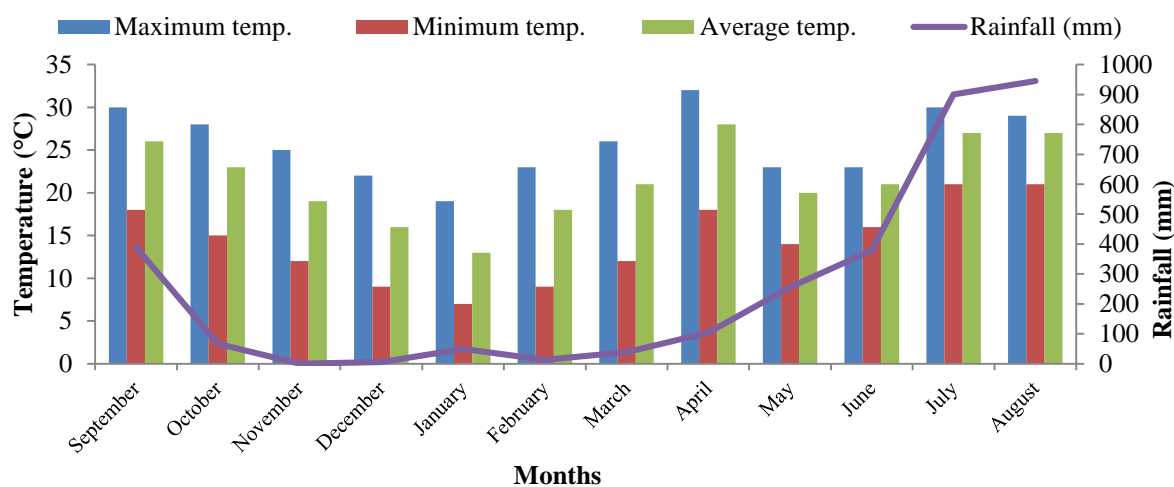


Figure 1. Meteorological data of study sites during September 2017-August 2018 (Source: ARIES, Nainital)

## Sampling procedure

Five sites at five different locations were selected in five different forest stands dominated with different kinds of genera and species of trees such as *Pinus roxburghii*, *Quercus leucotrichophora*, *Cedrus deodara*, *Cupressus torulosa* etc. For the herbaceous vegetation analysis, 1 ha plot was established in that forest stand and 10 quadrats of 10 m x 10 m were placed in it and in that each quadrat, three 50 cm x 50 cm size quadrats (5 site x 30 quadrats=150 quadrats) were placed randomly (Misra, 1968). The sampling was done in the month of September which is considered as a peak growth period as well as maximum availability of herb diversity and biomass production (Bargali and Bargali, 2000). The number and size of the quadrats were determined by the running mean method (Kershaw, 1973) and species area curve (Misra, 1968) method. Herb species of every quadrat were uprooted and species

number, density, provenance value and biomass were recorded for every species separately. Total biomass for herbs was calculated by adding the aboveground and belowground biomass. Unidentified species were put between paper sheets and marked according to their quadrat number. These unidentified herb species were then brought to laboratory and with the help of experts and relevant flora, they were identified. Uprooted (Both above ground and belowground parts) herb species were packed in paper bags and oven dried at 80°C for 48 hrs or till they achieve a constant weight. From each site soil samples were collected from two depths *i.e.* surface layer (0-15cm) and sub- surface layer (15-30cm) with the help of soil corer. For soil moisture, immediate soil weight was taken at respective sites and these samples were then put into polythene zipper bags, oven dried at 80°C for 48 hrs or till they completely devoid of any kind of moisture. Total soil nitrogen was determined by the micro-Kjeldahl method (Peach and Tracey, 1956).

### **Data Analysis**

For the determination of soil moisture following formulae was used

$$\text{Soil moisture (\%)} = \frac{\text{Fresh weight of soil} - \text{dry weight of soil}}{\text{dry weight of soil}} \times 100$$

Provenance value (PV) of each species was calculated by summing up their respective relative frequency (RF) and relative density (RD):

$$\text{Provenance Value (PV)} = \text{RF} + \text{RD}$$

The species with highest value of PV was identified as dominant and that having the second-highest value was defined as co-dominant species.  $\alpha$ -diversity was calculated using the Shannon–Wiener index ( $H'$ ).

$$H' = -\sum p_i \log_2 p_i \quad (\text{Shannon and Weaver, 1963})$$

Where,

$p_i$  is the proportion of total stand basal area represented by the  $i^{\text{th}}$  species.

The working formula given by Smith (1974) was used here as;

$$H' = 3.322 \left[ \sum \frac{N_i}{N} \log_{10} \frac{N}{N_i} \right]$$

Where,

$N_i$  is the total density of species and  $N$  is the total density of all the species. The factor 3.322 was used to convert the index value to  $\log_2$ .

Species evenness ( $e$ ) and  $\beta$ -diversity for each site was calculated using the following formulae:

$$e = \frac{H'}{\ln S} \quad (\text{Whittaker, 1972})$$

$$\beta = \frac{S_c}{\bar{S}} \quad (\text{Whittaker, 1972})$$

Where,  $H'$ =Diversity of species;  $S$ =Number of species,  $p_i$  = proportion of provenance value belonging to species ' $i$ ',  $S_c$  =total number of species,  $\bar{S}$  =Average number of species per sample.

Index of similarity (%) between two sites was calculated using Sorensen's index for presence/absence data as

$$SI = \frac{2C}{(a+b)}$$

Where, ' $C$ ' is the number of species of species common to both sites, ' $a$ ' is a number of species at site first and ' $b$ ' is the number of species at site two.

Species turnover (ST), a measure of floristic change between two selected sites, was calculated as:

$$ST = \frac{(1+g)}{(a+b)}$$

Where  $l$  is the number of lost and  $g$  is the number of species gained between two selected sites; 'a' and 'b' are the same as in equation for SI (Schoemaker and McKee, 1988).

The Statistical analysis was done by using SPSS 25.

## Results

Recorded herb species in the five forest stands are presented in Table 3. Total 73 herb species from 31 families were reported from five sites covering 37.5 m<sup>2</sup> area. Highest family Asteraceae (16) recorded maximum number of herb species followed by Lamiaceae (7) and Poaceae and Polygonaceae with 6 species each (Table 3). Total number of herb species per site ranged from 28 (PI and HG) to 50 (TA) (Table 5) and number of unique species per site varied from 2 (PI) to 11 (BP and TA) (Table 3), while 9 species were present in all the five sites. On the basis of provenance value (PV), out of five sites, three sites (PI, RU and TA) showed same dominant and co-dominant species as *Erigeron karvinskianus-Oxalis corniculata* respectively. At site- BP *Cynodon dactylon-Erigeron karvinskianus* whereas at site-HG *Oplismenus compositus-Cynodon dactylon* were dominant and co- dominant species, respectively. Density of herbs (plant m<sup>-2</sup>), soil type, soil moisture and total soil nitrogen of selected studied sites are given in Table 4. Highest density of herbs was 188 plant m<sup>-2</sup> at site-TA followed by site-BP (164 plant m<sup>-2</sup>) and site-PI (141 plant m<sup>-2</sup>) whereas the soil moisture varied between 5.56%(RU) to 11.43% (BP) in surface layer (0-15 cm) and 7.09% (RU) to 15.40% (BP) in the sub-surface layer (15-30 cm) . Total soil nitrogen ranged from 0.20% (RU) to 0.41% (TA) (Table 4). Herb density increased with increasing soil moisture (%) and total soil nitrogen (%) indicating that soil fertility increased growth of herbs (Table 4).  $\beta$ -diversity, Shannon index, evenness and biomass per site varied from 1.46 (TA) to 2.61 (PI and HG), 4.01 (PI) to 5.09 (TA), 1.204 (PI) to 1.316 (RU) and 2.36 kg m<sup>-2</sup> (PI) to 6.46 kg m<sup>-2</sup> (TA), respectively (Table 5). Figure 2 showed dominance diversity curve in which bottom curve (site-TA, *Erigeron karvinskianus-Oxalis corniculata* community) represented the highest diversity, while the uppermost curve (site-HG, *Oplismenus compositus-Cynodon dactylon* community) represented the lowest diversity.

Highest index of similarity was recorded between sites RU-TA (62.50%) followed by between sites PI-TA (61.33%). Among all the selected sites, only two combinations of sites has similarity index below 50% i.e. RU-BP (46.15%) and RU-PI (34.48%) (Table 6). Species turnover ranged from 0.31 to 0.54 with highest turnover between sites RU-BP (0.31) and lowest was between sites RU-PI (0.54) (Table 7). Across the sites, herbs showed all kinds of distribution pattern i.e. regular, random and contagious. Maximum number of herb species showed contagious distribution pattern followed by random and regular distribution (Fig. 3).

One-way ANOVA showed significant ( $p \leq 0.001$ ) changes in soil moisture (%) and total soil nitrogen with respect to selected forest stands (Table 8). Soil moisture and total soil nitrogen significantly affected species number and diversity. Shannon's index and biomass, species number and herb density are significantly related to each other (Fig. 4). Shannon's index and biomass of herbs showed non-significant relationship with both linear and non-linear models used in the analysis but linear model exhibited a better fit to the data compared to non-linear models (Table 9).

Table 3. Provenance value (PV) of herbaceous species at five sites in forests of Central Himalaya, India.

Herb species	Family	RU	HG	PI	BP	TA
<i>Achyranthes bidentata</i> Blume	Amaranthaceae	0.00	5.65	9.33	0.00	3.99
<i>Ageratum conyzoides</i> L.	Asteraceae	7.19	10.59	0.00	6.07	5.99
<i>Ajuga bracteosa</i> Wall. Ex Benth	Lamiaceae	0.00	0.00	0.00	0.00	5.14
<i>Anemone vitifolia</i> Buch.-Ham.ex DC	Asteraceae	0.00	0.00	0.00	3.54	0.00
<i>Anaphalis contorta</i> Hook.f.	Asteraceae	0.00	6.36	0.00	5.58	2.78
<i>Apluda mutica</i> L.	Poaceae	7.89	0.00	8.65	6.55	6.57
<i>Argemone mexicana</i> L.	Papaveraceae	0.00	2.83	0.00	3.06	3.36
<i>Arisaema tortuosum</i> (Wall.) Schott	Araceae	0.00	0.00	0.00	2.04	0.00

<i>Artemisia annua</i> L.	Asteraceae	0.00	0.00	0.00	0.00	5.57
<i>Artemisia nilagirica</i> C.B. Clarke	Asteraceae	0.00	0.00	0.00	6.07	0.00
<i>Arthraxon prionodes</i> (Steud.) Dandy	Poaceae	0.00	0.00	0.00	0.00	6.11
<i>Begonia picta</i> Smith	Begoniaceae	0.00	0.00	0.00	2.04	0.00
<i>Bergenia ciliata</i> (Haw.) Sternb	Saxifragaceae	0.00	0.00	0.00	4.56	0.00
<i>Boenninghausenia albiflora</i> (Hook.) Rchb. ex Meisn.	Rutaceae	4.28	0.00	5.82	3.79	3.78
<i>Bidens pilosa</i> L.	Asteraceae	10.05	5.65	8.48	4.56	1.85
<i>Bidens biternata</i> (Lour) Sheriff.	Asteraceae	1.79	9.53	0.00	0.00	3.36
<i>Bupleurum tenue</i> Buch.-ham ex D. Don	Apiaceae	2.85	0.00	0.00	0.00	2.00
<i>Cassia occidentalis</i> L.	Fabaceae	0.00	0.00	1.24	0.00	0.00
<i>Carex hirta</i> L.	Cyperaceae	0.00	4.59	0.00	0.00	0.00
<i>Cerastium vulgare</i> Hartm.	Caryophyllaceae	0.00	0.00	0.00	0.00	2.57
<i>Clematis burchananiana</i> DC.	Ranunculaceae	0.00	0.00	0.00	0.00	3.42
<i>Commelina benghalensis</i> L.	Commelinaceae	8.62	8.82	8.20	6.31	4.99
<i>Conyza stricta</i> Willd.	Asteraceae	0.00	0.00	0.00	8.26	4.84
<i>Craniotome furcata</i> (Link) Kuntze	Lamiaceae	1.79	0.00	0.00	0.00	0.00
<i>Craniotome versicolor</i> Rchb.	Lamiaceae	0.00	2.83	0.00	0.00	1.00
<i>Chrysopogon zizanioides</i> (L.) Robert	Poaceae	0.00	0.00	0.00	4.47	7.87
<i>Cymbalaria muralis</i> G. Gaertn., B. Mey. & Scherb	Plantaginaceae	0.00	0.00	7.80	0.00	5.26
<i>Cynodon dactylon</i> (L.) Pers.	Poaceae	0.00	15.81	0.00	20.82	0.00
<i>Cynoglossum lanceolatum</i> Forssk.	Boraginaceae	0.00	0.00	0.00	3.06	0.00
<i>Cyperus rotundus</i> L.	Cyperaceae	0.00	0.00	0.00	0.00	4.05
<i>Dicliptera bupleuroides</i> Nees	Acanthaceae	0.00	0.00	0.00	0.00	4.57
<i>Dioscorea deltoidea</i> Wall. Ex Griseb.	Dioscoreaceae	0.00	0.00	0.00	0.00	1.57
<i>Erigeron karvinskianus</i> DC.	Asteraceae	22.33	14.44	29.16	13.70	15.52
<i>Eupatorium adenophorum</i> Spreng.	Asteraceae	9.32	6.35	14.59	6.99	7.47
<i>Fagopyrum esculentum</i> Moench.	Polygonaceae	5.01	6.36	0.00	0.00	0.00
<i>Fragaria vesca</i> L.	Rosaceae	5.40	0.00	7.35	0.00	7.05
<i>Galinsoga parviflora</i> Cav.	Asteraceae	5.37	0.00	0.00	0.00	3.36
<i>Galium aparine</i> L.	Rubiaceae	0.00	0.00	0.00	10.64	2.42
<i>Geranium nepalense</i> Sweet.	Asteraceae	0.00	0.00	1.53	6.02	1.57
<i>Gerbera gossypina</i> (Royle) P. Beauv.	Asteraceae	0.00	4.95	4.86	1.26	3.57
<i>Hedera nepalensis</i> C. Koch	Araliaceae	2.85	0.00	1.24	0.00	1.00
<i>Impatiens bicolor</i> L.	Balsamiaceae	6.07	1.77	4.30	0.00	2.78
<i>Justicia simplex</i> D. Don	Acanthaceae	6.85	0.00	0.00	0.00	2.57
<i>Lepidagathis cristata</i> Willd.	Acanthaceae	0.00	0.00	1.24	0.00	0.00
<i>Leucas lanata</i> Benth.	Lamiaceae	1.79	2.83	3.05	0.00	0.00
<i>Micromeria biflora</i> (Buch.-Ham. ex D. Don) Benth	Lamiaceae	8.98	7.06	4.01	4.07	1.21
<i>Myriactis nepalensis</i> Less.	Asteraceae	0.00	0.00	0.00	3.54	0.00
<i>Oplismenus compositus</i> (L.) P. Beauv.	Poaceae	7.89	26.77	11.87	8.30	4.21
<i>Oxalis corniculata</i> L.	Oxilaceae	15.14	9.86	20.46	11.71	10.29
<i>Persicaria capitata</i> (Buch. Ham. Ex D. Don) H. Gross.	Polygonaceae	0.00	2.83	0.00	0.00	0.00
<i>Persicaria nepalensis</i> (Meisner) H. Gross	Polygonaceae	0.00	8.11	10.29	8.54	5.42
<i>Persicaria orientalis</i> (L.) Spach	Polygonaceae	0.00	0.00	0.00	2.04	0.00
<i>Parietaria officinalis</i> L.	Urticaceae	11.86	0.00	0.00	0.00	0.00
<i>Pennisetum clandestinum</i> Hochst.ex Chiov.	Poaceae	3.58	0.00	4.30	7.77	9.08
<i>Pimpinella acuminata</i> (Edgew.) C.B. Clarke	Apiaceae	2.15	0.00	2.49	0.00	2.57
<i>Polygonum chinense</i> L.	Polygonaceae	6.44	10.22	0.00	2.72	2.00
<i>Potentilla fragarioides</i> L.	Rosaceae	0.00	0.00	0.00	0.00	3.21
<i>Roylea cinerea</i> (D. Don) Baill.	Lamiaceae	0.00	0.00	0.00	3.79	0.00
<i>Rubia cordifolia</i> L.	Rubiaceae	0.00	0.00	0.00	0.00	2.78
<i>Rumex hastatus</i> D. Don	Polygonaceae	0.00	6.36	0.00	4.07	2.36
<i>Salvia officinalis</i> L.	Lamiaceae	0.00	2.83	0.00	0.00	1.57
<i>Smilax macrophylla</i> Roxb.	Smilacaceae	0.00	0.00	0.00	0.00	1.00
<i>Sigesbeckia orientalis</i> L.	Asteraceae	0.00	3.18	0.00	0.00	0.00
<i>Sonchus oleraceus</i> L.	Asteraceae	0.00	1.41	0.00	0.00	0.00
<i>Strobilanthes angustifrons</i> C.B. Clarke	Acanthaceae	7.19	6.00	4.30	1.51	2.57
<i>Thalictrum foliolosum</i> DC.	Ranunculaceae	5.37	0.00	7.92	0.00	3.78
<i>Trifolium repens</i> L.	Fabaceae	0.00	0.00	0.00	3.54	0.00
<i>Valeriana hardwickii</i> Wall.	Caprifoliaceae	6.10	0.00	4.58	4.52	2.57
<i>Veronica beccabunga</i> L.	Plantaginaceae	0.00	0.00	0.00	0.00	1.00
<i>Verbascum thapsus</i> L.	Scrophulariaceae	3.58	0.00	0.00	0.00	0.00
<i>Vicia villosa</i> Roth.	Fabaceae	0.00	0.00	1.24	0.00	0.00
<i>Viola canescens</i> Wall. Ex Roxb.	Violaceae	7.95	0.00	8.93	4.52	4.05
<i>Vitis himalayana</i> (Royle) Brandis	Vitaceae	4.28	6.01	2.77	0.00	2.36

Table 4. Species density of herbs, soil moisture content and total soil nitrogen at five sites in forests of Central Himalaya.

Sites	Species density (Plant m <sup>-2</sup> )	Soil type		Soil moisture (%)		Total soil nitrogen (%)
		0-15 cm	15-30 cm	0-15 cm	15-30 cm	
RU	110	Clay loam	Sandy clay loam	5.56±1.83	7.09±0.75	0.20±0.04
HG	114	Clay loam	Sandy clay loam	7.39±0.55	9.60±0.94	0.33±0.08
PI	141	Sandy clay loam	Sandy clay loam	6.98±0.73	9.07±0.71	0.25±0.05
BP	164	Clay loam	Sandy clay loam	11.43±0.98	15.40±1.28	0.36±0.34
TA	188	Sandy loam	Sandy clay loam	8.03±1.75	10.75±1.32	0.41±0.07

Table 5. Species diversity and biomass of the herbaceous vegetation at five sites in forests of Central Himalaya.

Sites	β-diversity	Species number	Shannon's index	Evenness	Biomass (kg m <sup>-2</sup> )
RU	2.43	30	4.48	1.316	5.23
HG	2.61	28	4.17	1.251	3.57
PI	2.61	28	4.01	1.204	2.33
BP	2.09	35	4.50	1.266	6.26
TA	1.46	50	5.09	1.301	6.46

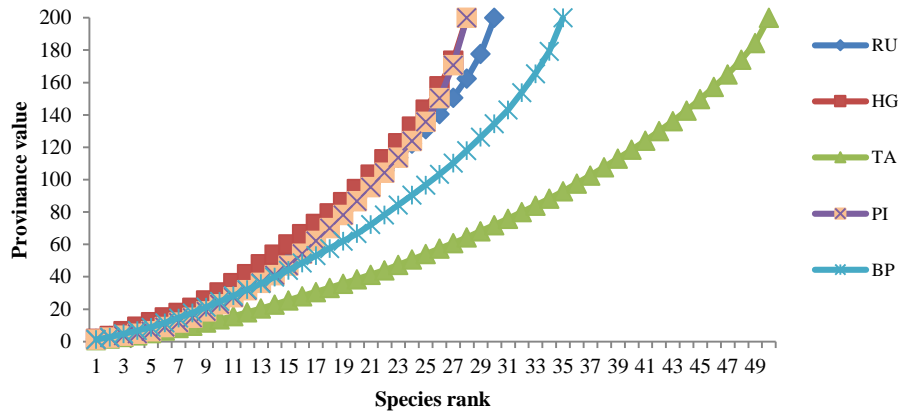


Figure 2. Dominance diversity curve, showing percentage cumulative provenance value plotted against species rank for each site.

Table 6. Index of similarity (%) between selected forest stands.

	RU	HG	PI	BP	TA
RU	100.0	51.72	34.48	46.15	62.50
HG		100.0	50.00	50.79	53.84
PI			100.0	50.70	61.53
BP				100.0	56.47
TA					100.0

Table 7. Species turnover (ST) between selected forest stands.

	RU	HG	PI	BP	TA
RU	1.00	0.48	0.31	0.54	0.38
HG		1.00	0.50	0.49	0.46
PI			1.00	0.48	0.39
BP				1.00	0.44
TA					1.00

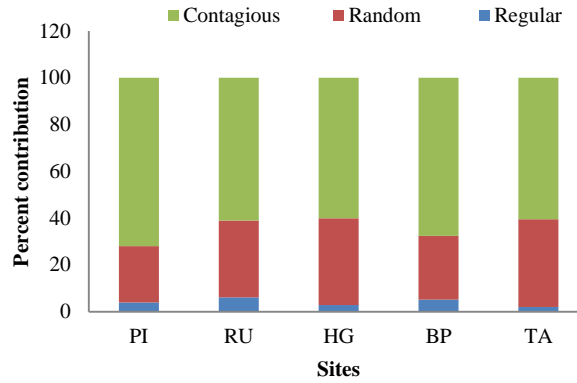
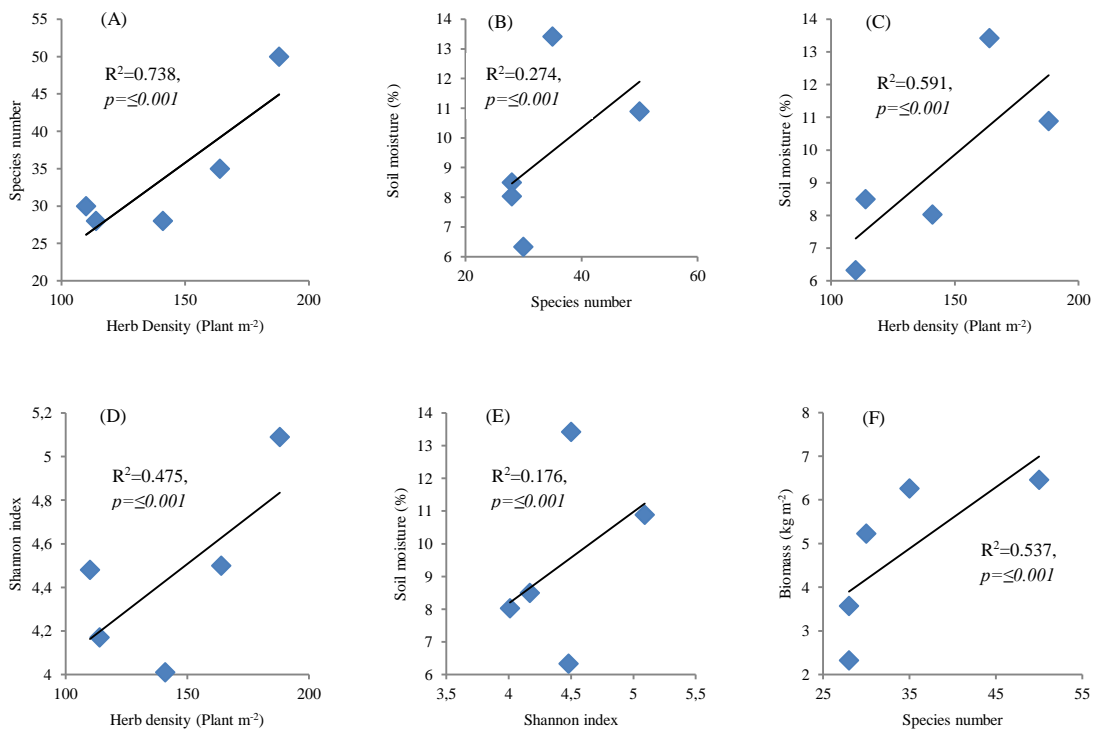


Figure 3. Distribution pattern of herb species at selected forest stands

Table 8. One-way ANOVA for species density (SD), soil moisture (SM) and total soil nitrogen with respect to the selected forest stands

Selected traits		Sum of Squares	df	Mean Square	F	Significance level (P<0.001)
SD	Between Groups	26395.200	4	6598.800	-	-
	Within Groups	0.000	25	0.000		
	Total	26395.200	29			
SM	Between Groups	144.438	4	36.109	7.973	0.000
	Within Groups	113.228	25	4.529		
	Total	257.666	29			
TSN	Between Groups	0.157	4	0.039	17.627	0.000
	Within Groups	0.056	25	0.002		
	Total	0.212	29			





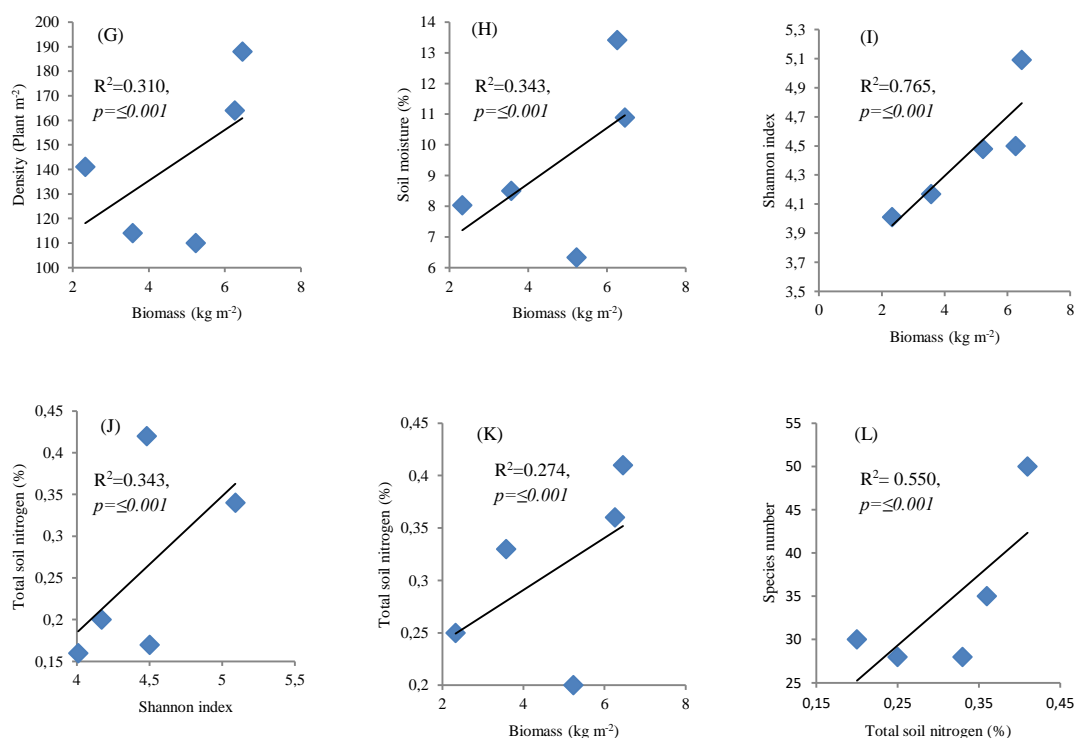


Figure 4. Relationship between different traits of herbs and soil.

Table 9. Relationship of herbaceous biomass with  $\alpha$ -diversity (Shannon index) and its component (species richness) in Central Himalaya forest, India.

Regression models	$R^2$	Standard error of estimate (Sy)	$p$
Relationships between species richness (X) and biomass (Y)			
Quadratic	0.865	0.924	0.135
Logarithmic	0.596	1.306	0.126
Power	0.509	0.350	0.176
Linear	0.538	1.397	0.159
Relationships between Shannon index (X) and biomass (Y)			
Quadratic	0.964	0.481	0.036
Logarithmic	0.794	0.932	0.042
Power	0.750	0.250	0.058
Linear	0.765	0.996	0.052

## Discussion

In specific time and space, diversity of vegetation is determined by one of the many abiotic factor soil. Small and McCarthy (2005) showed that biomass, composition and diversity of herbaceous vegetation responded sensitively to change in available soil nitrogen. Soil moisture and soil nitrogen favour growth of annuals or short-lived perennials (Bargali et al. 2015b), the highest soil nitrogen and optimum soil moisture recorded at site-TA resulted in maximum number of species. While at site-PI and HG, comparatively low soil moisture and soil nitrogen decreased species richness possibly due to competition between species for resources.

In this study, the herbaceous vegetation of Central Himalayan forest was identified and then the relationship of those plant species was developed with the soil moisture and nitrogen. Maximum number of herb species was reported at site -TA (50) and minimum number was reported at sites PI and HG (28). In contrast, according to Gilliam (2006) increase in N availability caused loss of many species that are efficient under low N-condition and resulted in decrease in species richness of herbaceous

vegetation. He proposed N homogeneity hypothesis, which predicts that the excess N inputs reduce the naturally high spatial heterogeneity in soil N availability (*i.e.* patchiness) that helps to maintain the species diversity of the herbaceous layer, the biodiversity of affected forests will decline (Gilliam, 2006).

Availability of soil moisture and nitrogen in ample amount play very vital role in flourishing herbaceous vegetation and this study showed significant effect of soil moisture and soil nitrogen on herb density as also demonstrated by some researchers (Lv and Dong, 2011; Maestre and Escudero, 2010; Garduno et al. 2010; Legate et al. 2010; Taf et al. 2017). Lowest herb density was recorded at site - RU with lowest percentage of soil moisture and nitrogen (Table 3) and maximum herb density was recorded at site- TA with highest soil nitrogen and soil moisture. It has been also recorded in studies demonstrating significant effect between soil moisture and plant diversity (Xu et al. 2015). However, plant community theory assumed interactions between different factors (Grime, 1979; Huston, 1979; Tilman, 1982; Keddy, 1989; Xiong et al. 2003) therefore; many experiments have combined more than two factors to test such effects.

Total density values of herbs in the present study are much higher than the total herb density (15.9–33.3 plant m<sup>-2</sup>) in the mid elevation forest of Central Himalayas, India (Khera et al. 2001) as well total herb density (68-114 plant m<sup>-2</sup>) in the forest stands in the Eastern ghat of India (Behera and Misra, 2006). Maximum species diversity (Shannon–Wiener index) was recorded in the TA forest stand (5.09), which showed maximum species number, may be due to decreased competition and increased resource availability (Sagar et al. 2003). The diversity index values (4.01-5.09) are positively correlated with soil moisture and soil N (Fig. 4). The evenness values ranged between 1.20 (PI) and 1.32 (RU) and showed an inconsistent pattern with soil moisture and soil N. According to Gilliam (2006) decrease in species evenness with increasing soil N is caused by the increasing dominance of relatively few species that require high N availability. Studies conducted on the influence of vegetation on soil moisture and suggested that different plant species can affect the temporal and spatial characteristics of the soil moisture in different manner (Zhuang et al. 2015; Musa et al. 2014; Yang et al. 2014; Chen et al. 2007; Zhang et al. 2013) and their distribution also affects the spatial and temporal changes in soil moisture patterns (Wang et al. 2015; Kong et al. 2009; Xu et al. 2015; García-Baquero et al. 2016). Apart from above noteworthy studies, it was determined that soil moisture did not have a significant effect on diversity unless it was part of an interaction with another variable (Xiong et al. 2003), such soil nitrogen of the research site (Smith et al. 2016).

Analysis of PV provides information about social status of a herb species and can be used to recognize the pattern of association of dominant species in a community. Analysis of PV indicated that the three forest stands (PI, RU and TA) represented similar combinations *Erigeron karvinskianus* (dominant) and *Oxalis corniculata* (co-dominant) species. The reason that these species grow together is usually because they have similar requirement and their dominance at these sites could be possible due to availability of optimum conditions for their growth. *Cynodon dactylon* was the dominant species in the BP forest stand while the co- dominant species (*Erigeron karvinskianus*) is a common species of the ground vegetation of the natural forest in the area, and was recorded from all the five forest stands. *Oplismenus compositus* the dominant species in the HG forest stand is a member of family poaceae and was recorded from all the five forest stands while *Cynodon dactylon* was the co- dominant species at this site. Therefore, it is concluded that plant species occupied specific climatic and edaphic factors experience variability in their composition.

When the provenance values (PV) of the herbaceous species of the five forest stands were ordinated against the species sequence, the dominant–diversity curves (Whittaker, 1965) followed lognormal distribution in all the stands (Fig. 2). These dominance–diversity curves in the stands indicate the heterogeneity of the species (Bahera and Misra, 2006, Pandey et al. 2018) and suggest that species

importance is governed by a large number of factors for success in the niche hyperspace (Whittaker, 1970). Higher values of PV in any individual species indicate that all the available resources are being utilized by that species and left over being utilized by other species as the competitor and associates. Similar types of curves have been reported by Sagar et al. (2008a) for the dry-tropical forest in Northern India.

The herbaceous layer significantly affects the structure and functioning of forest ecosystems. Though it represents less than 1% of the biomass of the forest but contains 90% or more of the plant species of the forest vegetation and contributes up to 20% of the foliar litter to the forest floor (Gilliam, 2007). The highest herbaceous biomass in the TA forest stand may be attributed to higher diversity and density of herbaceous species. In addition, more moisture and soil nitrogen was available at this site as compared to other sites. Decline in biomass in other forest stands was due to competition among species for the resources, which increased with decrease in resource availability. In this study, herbaceous biomass showed significant positive correlation with species number, density, soil moisture and nitrogen (Fig. 3). The herbaceous standing crop biomass reported in this study (2.36-6.46 kg m<sup>-2</sup>) was higher than the values (0.31-0.44 kg m<sup>-2</sup>) reported for the herbaceous layer of mixed conifer forest in Central Himalaya (Bargali et al. 2015b; Mourya et al. 2019) and 0.008 kg m<sup>-2</sup> to 0.071 kg m<sup>-2</sup> reported for the fire affected dry forest ecosystem of Bhoramdeo wildlife sanctuary, India (Jhariya and Singh, 2020a).

Biodiversity is being studied based on species number and evenness (Harper et al. 1994) but diversity is not just based on the number of individuals, but it is also very important that to which species they belong (Peng et al. 2018) because type of species diversity is somewhere directly affected by type of soil characteristics (Migala et al. 2014). In comparison to other studies, smaller differences in soil moisture between sites may be due to shorter (one year) study period. For example, soil moisture data were analysed over 40 years by Song et al. (2013) and monitored soil moisture and temperature were monitored for 3 years by Morecroft et al. (1998). Microclimatic and soil moisture conditions over 6 year period under different gap sizes (canopy cover) were studied by Gray et al. (2002) and a significant effect of gap size on soil moisture and temperature was recorded.

This study demonstrated that the levels of soil moisture and nitrogen significantly affected the patterns of herbaceous floral composition. The individual forest stands harboured 28-50 species, with only nine species being common to all the five forest stands, however, collectively they harboured 73 species. For the management of herbaceous diversity, soil moisture and nutrients can indeed be an important tool. Therefore, it is important to understand how soil moisture impacts plant diversity, because in future climate change will change the soil moisture due to change in pattern of rainfall and increase in temperature. Having a better understanding of how an ecosystem is affected by changes in soil moisture will be helpful to cope with the decreased soil moisture condition. Due to the importance of herbaceous vegetation in forest ecosystems, it is vital that there should be sufficient data that will lead to the proper conservation management.

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