

Structure, composition and diversity of trees within the dry evergreen reserve forest of Kondapalli (Eastern Ghats, southern India)

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Abstract. The dry evergreen forest of Kondapalli (Andhra Pradesh state, India) is declared as a forest reserve, but, despite of this, it is subjected to degradation resulting in loss of biodiversity. Thus, the current study was carried out to investigate the tree diversity of Kondapalli forest. A total of 566 ± 16 trees (≥ 10 cm) representing 46 ± 8 species from 40 genera and 21 families were recorded from the 0.36 ha area of Kondapalli forest. Mimosaceae was the most species rich family, while Rutaceae was the most abundant family. *Atalantia monophylla* was most frequent and abundant species and, with respect to basal area, *Melia azedarach* and *Syzygium cumini* were the dominant taxa. The recorded stem density was 1572 stems ha^{-1} and the mean forest basal area was 47.17 m^2 ha^{-1} . The results of cluster analysis revealed that *Atalantia monophylla*, characterised by a high ecological amplitude, had a wide distribution and was associated with species forming different communities. The study showed that Kondapalli forest is characterised by a fairly high species richness, which provides the baseline data on the floristic structure and diversity of this forest for better management and conservation.

Key words: dry evergreen forest, degradation, diversity, species richness, ecological amplitude

1. Introduction

Currently, tremendous pressure is exerted on natural resources, exploiting them at the maximum limit beyond their regeneration and recover capacity (Pech & Sunada 2008; Timah *et al.* 2008). Globally, forest ecosystems are threatened due to various anthropogenic activities connected with overuse of forest resources or area encroachment for agriculture, settlement and economic development (Iftekhhar & Hoque 2005). Depletion and degradation of forests not only results in the loss of valuable goods and services provided by them, but also have an effect on the climate (Prasad *et al.* 2010). In turn, climate change has a negative feedback on ecosystems by modifying their structure, as well as species composition. A component that is worst affected by the cumulative pressure of anthropogenic and natural factors, is biodiversity. The recent studies on biodiversity revealed the consequences of climate change, such as: sea level rise leading to the loss of bio-

diversity due to the submergence of islands and coastal areas (Bellard *et al.* 2014), rise in temperature resulting in the shifting of species (Donato 2013), replacement of native species by invasive species (Kumari *et al.* 2010), and transformation of forest types (Prasad *et al.* 2010), etc.

Some natural ecosystems across the world have been declared as protected and reserve areas, wildlife sanctuaries and national parks. Also, some areas were tagged as “Biodiversity Hotspots”, due to their high species richness, diversity, and endemism (Marchese 2015). The main purpose of giving such a status to forest ecosystems is to retain their virgin structure and biodiversity and restore them, if they are in degraded condition, through strict protection against human interference. However, in spite of assigning such a legal status to these ecosystems, their exploitation is in some cases still continued, resulting in the destruction, degradation and loss of biodiversity. An example of such a situation is the Kondapalli Reserve Forest (KRF) of

Krishna district, Andhra Pradesh state (AP), India. This forest, in the form of a remnant patch, serves as green lungs for the city of Vijayawada in the Krishna district and is under a serious threat resulting from various human interventions (Salghuna *et al.* 2018). The forest was declared a reserve, with demarcated boundaries, in 1980 under the forest conservation Act. Despite declaring it a reserve area, it is subjected to degradation resulting in the shrinking of forest along the boundaries, as well as within its interior (Anonymous 2001; Prasad *et al.* 2011). Several changes in land use and land cover occurred in the vicinity of KRF, such as: agricultural development, mining activities, and establishment of settlements and industries. All these changes not only deteriorated the forest, but also affected its biodiversity (Pullaiah & Sandhya Rani 1999). Salghuna *et al.* (2018) reported a decrease in the forest cover from 11500 ha (1990) to 10600 ha (2015). In addition, a new threat to the forest comes from the proposal of AP Government to denotify a portion (890.43 ha) of KRF for a new state capital establishment (Anonymous 2017).

Hitherto, some floristic inventories and medicinal plant surveys were conducted in KRF (Venkanna 1990; Reddy *et al.* 2005, 2010), but no attempt has been made to quantify the floristic structure, species richness and tree diversity patterns. A detailed assessment of tree diversity of the reserve is essential in the context of human interference needed for its conservation. Hence, the objectives of this study were: (i) to investigate the floristic structure, composition and diversity of KRF (ii) to check

whether it is apt to compare ecosystems flourishing under dissimilar environmental conditions (iii) to review the diversity patterns of other dry evergreen forests of southern India and assess the status of KRF in relation to those forests, but not to compare their diversity patterns. The study is first of its kind carried out for KRF. The result of this research is expected to provide better insights for formulating and strengthening the conservation measures in KRF and for the sustainability of this reserve in long term perspective.

2. Materials and methods

2.1. Study area

The Eastern Ghats (EG) of the southern India form one of nine floristic zones of the country that stretches across three states: Tamil Nadu, AP and Orissa. They are discontinuous and divided into the southern and northern EG. KRF (situated between 16°37'N and 80°31'E latitudes and 16°45'N and 80°26'E longitudes), with an area of 121 km², is located on the western side of Vijayawada city and forms a connection between these two parts (Rao & Pullaiah 2007) (Fig. 1). The study area is characterised by tropical climatic conditions with an average annual temperature of 28.5°C and rainfall of 1067 mm (climate.org). Geologically, the area is dominated by gabbroic and anorthosite rocks with subordinate ultramafic rocks, plagioclase, orthopyroxene and clinopyroxene (Leelanandam 1994).

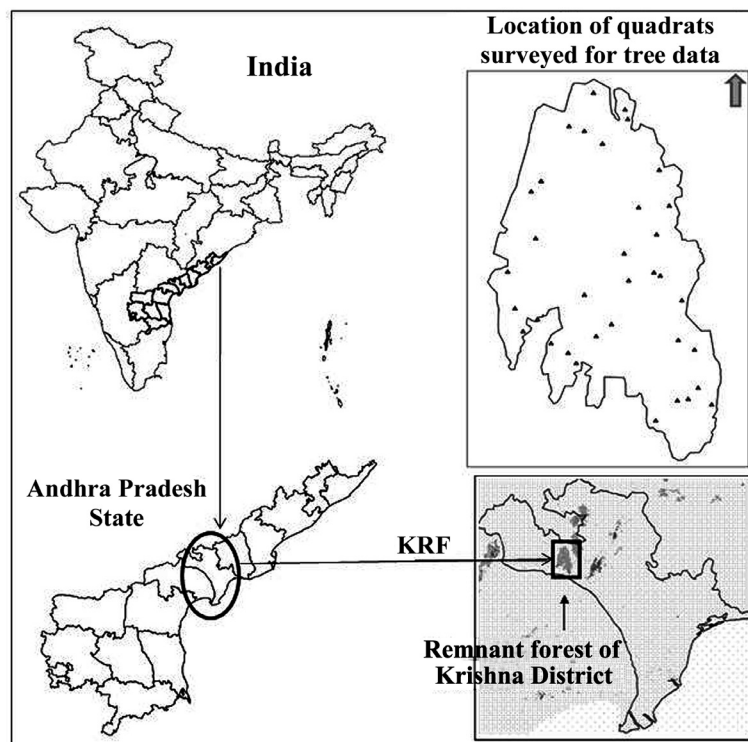


Fig. 1. Location map of the Kondapalli Reserve Forest (KRF)



Fig. 2. Different views of tropical dry evergreen forest vegetation in the Kondapalli Reserve Forest

Champion & Seth (1968) classified KRF as tropical dry evergreen forest-7/CI (Fig. 2). It has some unique floristic elements with bushy habit, making it difficult to enter into the forest. KRF is divided into five forest beats (sections), i.e., Kondapalli, Duggiralapadu, Jujjru, Kanchikacherla and Mullapadu. The dominant trees in this forest include *Atalantia monophylla*, *Strychnos potatorum*, *Gyrocarpus americanus*, *Albizia amara*, *Givotia moluccana* and *Chomelia asiatica*.

2.2. Data collection

The field work on quantitative inventory was carried out between June 2013 and May 2014. Considering the operational forest area chosen for study (10,600 ha) and based on our reconnaissance field survey, the quadrat

size was limited to 10 × 10 m (0.01 ha) area. A total of 36 quadrats were randomly surveyed in the five forest beats based on accessibility and intactness of the forest. Degraded and open forest areas were eliminated during inventory. Geographical coordinates of quadrats were taken using a GPS device. All trees of ≥ 10 cm diameter at breast height (DBH) were measured at 1.3 m from the ground using a diameter tape. Species identification was done while carrying out the inventory with the help of a taxonomist accompanying during field visits. For unknown species, specimens were collected with proper field information (plot number, locality, habitat, etc.) and were identified with the help of literature and flora (Pullaiah & Ramamurthy 2001; Reddy *et al.* 2001, Sandhyarani *et al.* 2007; Pullaiah & Rao 2002).

2.3. Data analysis

Species diversity was calculated using the Shannon-Wiener index with log base 2 (H-Shannon & Wiener 1963), Dominance by Simpson (D-Simpson 1949) index and Evenness by Pielou (1975). The relation between diversity components was calculated using the SHE analysis equation $H = \ln(S) + \ln(E)$ (Buzas & Hayke 2005). Species richness (S) was determined by totaling the number of species in all the quadrats sampled and estimated using Jackknife1, Jackknife2 and Chao2 (Heltsh & Forrester 1983; Smith & Van Belle 1984; Chao 1987 respectively). The Jackknife1 considers the number of unique species, whereas the Jackknife2 and Chao2 use unique species (found in only one quadrat) and duplicate species (found in two quadrats) data and the number of quadrats sampled. To estimate the species richness, we used the EstimateS (Colwell 2004) software default settings of sampling without replacement and sampled quadrats were randomized for 1000 runs. Estimates of species richness were analyzed graphically by plotting the estimator and observed species richness as a function of cumulative number of quadrats sampled. Based on the Raunkiaer (1934) classification, heterogeneity of the forest was determined as the distribution of species in five frequency (%) classes, i.e., A = 1-20%, B = 21-40%, C = 41-60%, D = 61-80% and E = 81-100%.

Family importance value (FIV) was calculated as the sum of relative density, relative dominance and relative diversity (Mori *et al.* 1983). For each species, frequency, density, basal area and abundance were computed. To understand the species share in the forest community,

importance value index (IVI) was calculated as the sum of its relative dominance (RDm), relative density (RD) and relative frequency (RF) (Cottam & Curtis 1956). The spatial distribution of species was measured using the Index of dispersion (D) using calculation of the variance to mean ratio (Selby 1965).

To understand the population structure (Rao *et al.* 1990), the tree data were divided into eight girth classes with 20 cm diameter intervals. In each girth class, species richness, diversity, stem density and basal area were analyzed. To delineate the dominant species communities occurring within the forest, a cluster analysis was performed with PAST software using Ward's method with Euclidean distance. Each cluster was delineated as a different community with dominant and co-dominant/associated species (Caswell 1976).

3. Results and discussion

3.1. Floristic structure, composition and diversity

3.1.1. Diversity, evenness, richness and heterogeneity

A total of 566 ± 16 trees (≥ 10 cm girth) representing 46 ± 8 species from 40 genera and 21 families were recorded from 0.36 ha of KRF. Out of 46 species, 43 were identified as medicinally important species. The recorded Shannon (H), Simpson (D) and Evenness (E) Index values were 3.2, 0.07 and 0.58 respectively. H considers both S and E and is biased by sample size. Calculation of the error using the formula $(S-1)/2N$ (N =sample size) showed a value of 0.04, which is

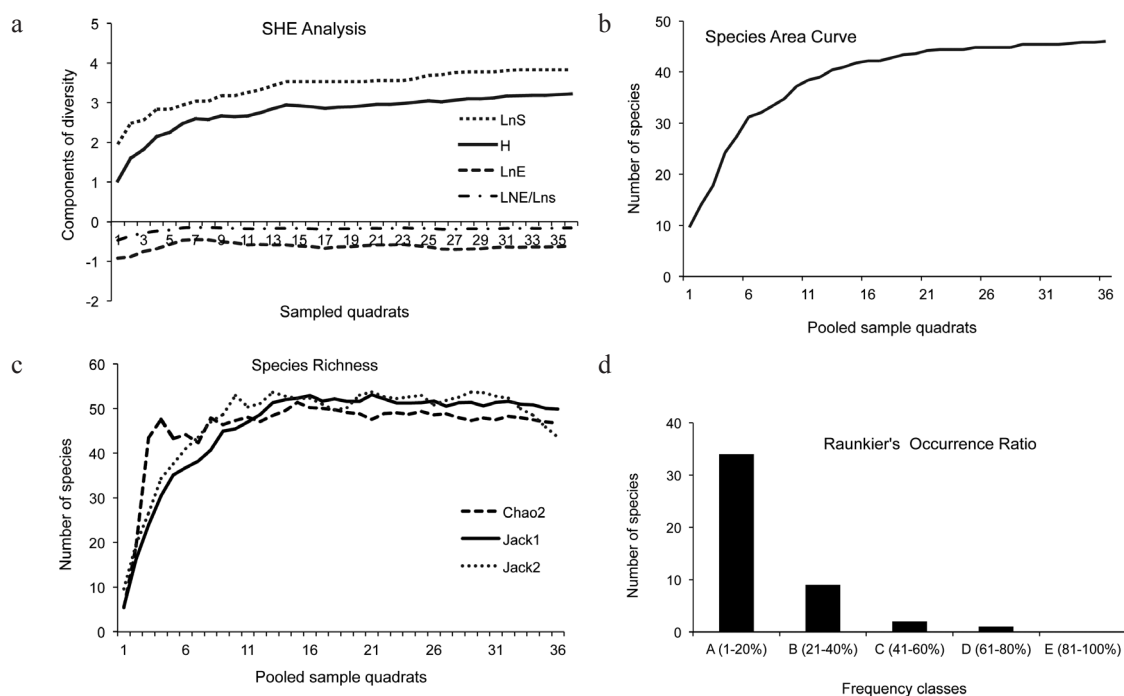


Fig. 3. Structure and diversity indices of the Kondapalli Reserve Forest

smaller than H, indicating that the calculated H is an acceptable estimate. The D value of 0.07 indicates a heterogeneous community and E value (0.58) represents the abundance of a few species in the forest. In addition, the Simpson index of diversity was obtained by subtracting the D value from 1(1-D i.e. 1-0.07= 0.93). For D, value approaching 1 indicates a highly homogenous community, while for the Simpson diversity Index values approaching 1 is a high species diversity community. Simpson diversity is less sensitive to species richness and more to evenness, while H shows a reverse trend (Colwell 2009).

Furthermore, SHE analysis provided a better understanding about the relationship between S, H, and E within the KRF community (De Benedictis 1973; Stirling & Wilsey 2001; Fig. 3a). The diversity indices, S and H showed an increased trend, while E showed negative values or the reverse. As the sample number increased, both S and H showed maximum value > 3, while it was low for E. The constant ln E/S and H with the increasing number of samples indicate the characteristic structure of forest between lognormal and log series distribution, which is a common feature of forest community with the low number of abundant species (Magurran 2004; Reshi *et al.* 2009). The E (0.58) value recorded in the present study also confirms this statement.

The observed species richness curve declined with increased sampling area and gradually reached an asymptote (Fig. 3b). The curve started flattening at an area of 0.29 ha. This indicates that the sampled area (0.36 ha) is sufficient to capture the species richness of KRF. The species richness predicted by estimators Chao2, Jackknife1 and Jackknife2 was 47, 50 and 44 species, respectively (Fig. 3c). The bias observed in species richness estimations either positive (over-47/50 species) or negative (under 44) is smaller compared to actual observed species (46) obtained from the field data. Among the three estimators, Jackknife1 was found to be best as it reached S_{max} quickly at 0.14 ha and remained relatively constant. In conclusion, the predicted species richness number in KRF is 50 compared to observed richness of 46. However, as suggested by Kotz & Johnson (1982-1988) and Stuart & Ord (1991), a good estimator is the one that estimates the values nearer to the true values. In such case, Chao2 can be considered a better estimator with the predicted species number of 47 that is closest to the true value observed from the inventory (46).

The frequency of Raunkiaer's classes followed the pattern A>B>C>>D>E. As per Raunkiaer, when classes A, B, C, D are high, the community is considered to be heterogeneous; on the other hand, if the class E is greater than another, it is a uniform or homogenous community.

Table 1. Family importance values found for the Kondapalli Reserve Forest in decreasing order

No.	Family	Individuals	BA m ² ha ⁻¹	Species	Genera	RD	RDm	RDv	FIV
1	Mimosaceae	76	1.30	9	4	13	8	19.6	40.6
2	Rutaceae	122	1.84	3	3	22	11	6.5	38.9
3	Rubiaceae	61	1.66	5		11	10	10.9	31.4
4	Loganiaceae	58	1.43	2	1	10	8	4.3	23.0
5	Meliaceae	10	2.26	2	2	2	13	4.3	19.4
6	Euphorbiaceae	38	1.33	2	2	7	8	4.3	18.9
7	Burseraceae	26	1.58	1	1	5	9	2.2	16.1
8	Hernandiaceae	38	1.07	1	1	7	6	2.2	15.2
9	Myrtaceae	5	1.88	1	1	1	11	2.2	14.1
10	Papilionaceae	20	0.29	4	3	4	2	8.7	13.9
11	Apocynaceae	21	0.40	3	3	4	2	6.5	12.6
12	Caesalpiniaceae	12	0.35	2	2	2	2	4.3	8.5
13	Sapindaceae	15	0.15	2	2	3	1	4.3	7.9
14	Combretaceae	11	0.23	2	2	2	1	4.3	7.6
15	Fabaceae	7	0.52	1	1	1	3	2.2	6.5
16	Anacardiaceae	13	0.20	1	1	2	1	2.2	5.6
17	Santalaceae	12	0.11	1	1	2	1	2.2	4.9
18	Moringaceae	7	0.23	1	1	1	1	2.2	4.8
19	Sterculiaceae	7	0.07	1	1	1	0	2.2	3.8
20	Lecythidaceae	5	0.06	1	1	1	0	2.2	3.4
21	Sapotaceae	2	0.02	1	1	0	0	2.2	2.6
	Total	566	16.98	46	40	100	100	100	300

Explanations: BA – basal area, RD – relative density, RDm – relative dominance, RDv – relative diversity, FIV – family importance value

As observed (Fig. 3d), the majority of species are found in the lower frequency class A=1-20% (34), representing the heterogeneous nature of the forest community in concurrence with D (Simpson) values. Overall, the value of D, 1-D and H indicates KRF as heterogeneous and diverse community in relation to its size.

3.1.2. Family and species dominance

Mimosaceae was the most species rich family with nine species, followed by Rubiaceae (five species), Papilionaceae (four species), Rutaceae and Apocynaceae (3 species each). Out of 21 families, 10 families were represented by only one species. Within Mimosaceae, the genus *Acacia* was represented by four species, *Albizia* by three and *Samania* and *Xylia* by one species, respectively. Of the recorded families, Rutaceae was the most abundant family with 122 individuals, while Sapotaceae was represented by only two individuals belonging to *Manilkara hexandra*. The family Rutaceae accounted for 22% of tree density, followed by Mimosaceae (13%) and Rubiaceae (11%), which summed up to 46% of the total tree density. About 13% of the basal area was contributed to the family Meliaceae, followed by Myrtaceae (11%) and Rutaceae (11%). These three families together constituted 35% of the total forest basal area. The maximum FIV value was observed for Mimosaceae (40.6) followed by Rutaceae (38.9), Rubiaceae (31.4), Loganiaceae (23.0) and Meliaceae (19.4), together accounting for 50% of total FIV (Table 1).

Atalantia monophylla was the most frequent and abundant species represented by 113 individuals out of 566 and provided 20% of the total tree density, followed by *Strychnos potatorum* (41). With respect to basal area, *Melia azedarach* and *Syzygium cumini* were the dominant species contributing to 13 and 11% of the total basal area. In terms of species importance based on IVI, *Atalantia monophylla* recorded a high value of 38.10, followed by *Strychnos potatorum* (19.33), *Commiphora*

caudate (19.02), *Gyrocarpus americanus* (18.86), and *Givotia moluccana* (16.36), together contributing 37% to overall IVI (Table 2). Low value of IVI was recorded for *Pongamia pinnata*. The observed dominance of particular species may be due to their optimal resource use, dispersal mechanism, function of stress and disturbances that create more space with high competition and exclusive growth (Wisheu & Keddy 1992; Richards 1996; Allison & Vitousek 2004).

3.1.3. Stem density and basal area

The recorded stem density is 1572 stems ha⁻¹. Out of 46 species, 39 species were represented by < 17 individuals and 7 species > 26. About 21% of the species were represented as single and two individuals (8) falling under the category of rare species. Probably they may be the victims of local extinction owing to their small populations (Primack & Hall 1992). About 45.4% of the stem density is contributed by top five species, i.e., *Atalantia monophylla* (314), *Strychnos potatorum* (114), *Gyrocarpus americanus* (105), *Albizia amara* (97) and *Givotia moluccana* (83). The mean basal area of the forest is 47.17 m² ha⁻¹. Maximum DBH was recorded for *Melia azedarach* (450 cm) followed by *Syzygium cumini* (361 cm). The top five species contributing 50% of basal area include: *Melia azedarach* (6%), *Syzygium cumini* (5.2%), *Atalantia monophylla* (4.8%), *Commiphora caudate* (4.4%), and *Myrtagyna parviflora* (3.4%).

3.1.4. Species spatial pattern

Out of 46 species, 25 species showed clumped and 21 random spatial distribution. The dispersion pattern of species is mainly governed by their interaction with the physical environment/microclimatic conditions (Diggle 1983; Armesto *et al.* 1986), the availability of resources and competition among species, mainly in their seed dispersal mechanism (Seidler & Plotkin 2006). Populations of species with cluster dispersion

Table 2. Importance value indices of species recorded in the Kondapalli Reserve Forest

No.	Species Name	Individuals	Abundance	Frequency	Density	BA m ² ha ⁻¹	RF	RD	RDm	IVI
1	<i>Atalantia monophylla</i> (L.) DC.	113	514	61.11	313.89	1.716	8.03	19.96	10.11	38.1
2	<i>Strychnos potatorum</i> L.f.	41	293	38.89	113.89	1.185	5.11	7.24	6.98	19.33
3	<i>Commiphora caudata</i> (Wight & Arn.) Engl.	26	186	38.89	72.22	1.584	5.11	4.59	9.33	19.03
4	<i>Gyrocarpus americanus</i> Jacq.	38	238	44.44	105.56	1.072	5.84	6.71	6.31	18.86
5	<i>Givotia moluccana</i> (L.) Sreem.	30	273	30.56	83.33	1.198	4.01	5.3	7.05	16.37
6	<i>Melia azedarach</i> L.	8	133	16.67	22.22	2.146	2.19	1.41	12.64	16.24

7	<i>Albizia amara</i> (Roxb.) B. Boivin	35	269	36.11	97.22	0.684	4.74	6.18	4.03	14.96
8	<i>Syzygium cumini</i> (L.) Skeels	5	167	8.33	13.89	1.881	1.09	0.88	11.08	13.06
9	<i>Mitragyna parviflora</i> (Roxb.) Korth.	12	171	19.44	33.33	1.241	2.55	2.12	7.31	11.98
10	<i>Chomelia asiatica</i> (L.) Kuntze	28	175	44.44	77.78	0.107	5.84	4.95	0.63	11.41
11	<i>Strychnos nux-vomica</i> L.	17	155	30.56	47.22	0.249	4.01	3	1.46	8.48
12	<i>Tamarindus indica</i> L.	7	117	16.67	19.44	0.518	2.19	1.24	3.05	6.48
13	<i>Acacia auriculiformis</i> Benth.	11	110	27.78	30.56	0.109	3.65	1.94	0.64	6.23
14	<i>Dalbergia paniculata</i> (Roxb.) Thoth.	12	150	22.22	33.33	0.149	2.92	2.12	0.88	5.92
15	<i>Schleichera oleosa</i> (Lour.) Oken.	12	133	25	33.33	0.073	3.28	2.12	0.43	5.83
16	<i>Santalum album</i> L.	12	150	22.22	33.33	0.11	2.92	2.12	0.65	5.69
17	<i>Wrightia tinctoria</i> R. Br.	13	325	11.11	36.11	0.314	1.46	2.3	1.85	5.61
18	<i>Cassia fistula</i> L.	10	143	19.44	27.78	0.172	2.55	1.77	1.01	5.34
19	<i>Canthium dicoccum</i> (Gaertn.) Merr.	9	129	19.44	25	0.139	2.55	1.59	0.82	4.96
20	<i>Lannea coromandelica</i> (Houtt.) Merr.	13	325	11.11	36.11	0.195	1.46	2.3	1.15	4.91
21	<i>Acacia chundra</i> (Rottler) Willd.	11	183	16.67	30.56	0.11	2.19	1.94	0.65	4.78
22	<i>Morinda pubescens</i> Sm.	8	114	19.44	22.22	0.125	2.55	1.41	0.73	4.7
23	<i>Anogeissus latifolia</i> (DC.) Wallich ex Guill. & Perr.	9	225	11.11	25	0.197	1.46	1.59	1.16	4.21
24	<i>Moringa concanensis</i> Nimmo	7	175	11.11	19.44	0.23	1.46	1.24	1.35	4.05
25	<i>Phyllanthus reticulatus</i> Poir.	8	200	11.11	22.22	0.129	1.46	1.41	0.76	3.63
26	<i>Holarrhena pubescens</i> Wall. ex G. Don	7	140	13.89	19.44	0.082	1.82	1.24	0.48	3.54
27	<i>Albizia lebbek</i> (L.) Benth.	6	120	13.89	16.67	0.066	1.82	1.06	0.39	3.28
28	<i>Aegle marmelos</i> (L.) Correa.	6	120	13.89	16.67	0.059	1.82	1.06	0.35	3.23
29	<i>Pterospermum canescens</i> (Roxb.)	7	175	11.11	19.44	0.073	1.46	1.24	0.43	3.12
30	<i>Sesbania grandiflora</i> (L.) Poiret	5	125	11.11	13.89	0.108	1.46	0.88	0.63	2.98
31	<i>Ixora pavetta</i> Andr.	4	133	8.33	11.11	0.053	1.09	0.71	0.31	2.11
32	<i>Samanea saman</i> F. Muell.	2	100	5.56	5.56	0.153	0.73	0.35	0.9	1.98
33	<i>Careya arborea</i> Roxb.	5	250	5.56	13.89	0.056	0.73	0.88	0.33	1.94
34	<i>Bauhinia racemosa</i> Lam.	2	200	2.78	5.56	0.181	0.36	0.35	1.07	1.78
35	<i>Azadirachta indica</i> A. Juss.	2	100	5.56	5.56	0.117	0.73	0.35	0.69	1.77
36	<i>Xylia xylocarpa</i> (Roxb.) Taub.	4	200	5.56	11.11	0.056	0.73	0.71	0.33	1.77
37	<i>Sapindus emarginatus</i> Vahl	3	150	5.56	8.33	0.073	0.73	0.53	0.43	1.69
38	<i>Chloroxylon swietenia</i> (Roxb.) DC.	3	150	5.56	8.33	0.068	0.73	0.53	0.4	1.66
39	<i>Acacia ferruginea</i> DC.	2	100	5.56	5.56	0.089	0.73	0.35	0.52	1.61
40	<i>Acacia concinna</i> (Willd.) DC.	3	150	5.56	8.33	0.02	0.73	0.53	0.12	1.38
41	<i>Dalbergia latifolia</i> (Roxb.)	2	100	5.56	5.56	0.031	0.73	0.35	0.18	1.26
42	<i>Terminalia arjuna</i> (Roxb.) Wight & Arn.	2	100	5.56	5.56	0.029	0.73	0.35	0.17	1.26
43	<i>Manilkara hexandra</i> (Roxb.) Dubard	2	100	5.56	5.56	0.018	0.73	0.35	0.11	1.19
44	<i>Albizia odoratissima</i> (L.f.) Benth.	2	100	5.56	5.56	0.012	0.73	0.35	0.07	1.15
45	<i>Plumeria alba</i> L.	1	100	2.78	2.78	0.004	0.36	0.18	0.02	0.56
46	<i>Pongamia pinnata</i> Pierre	1	100	2.78	2.78	0.002	0.36	0.18	0.01	0.55

Explanations: BA – basal area, RF – relative frequency, RD – relative density, RDm – relative dominance, IVI – importance value index

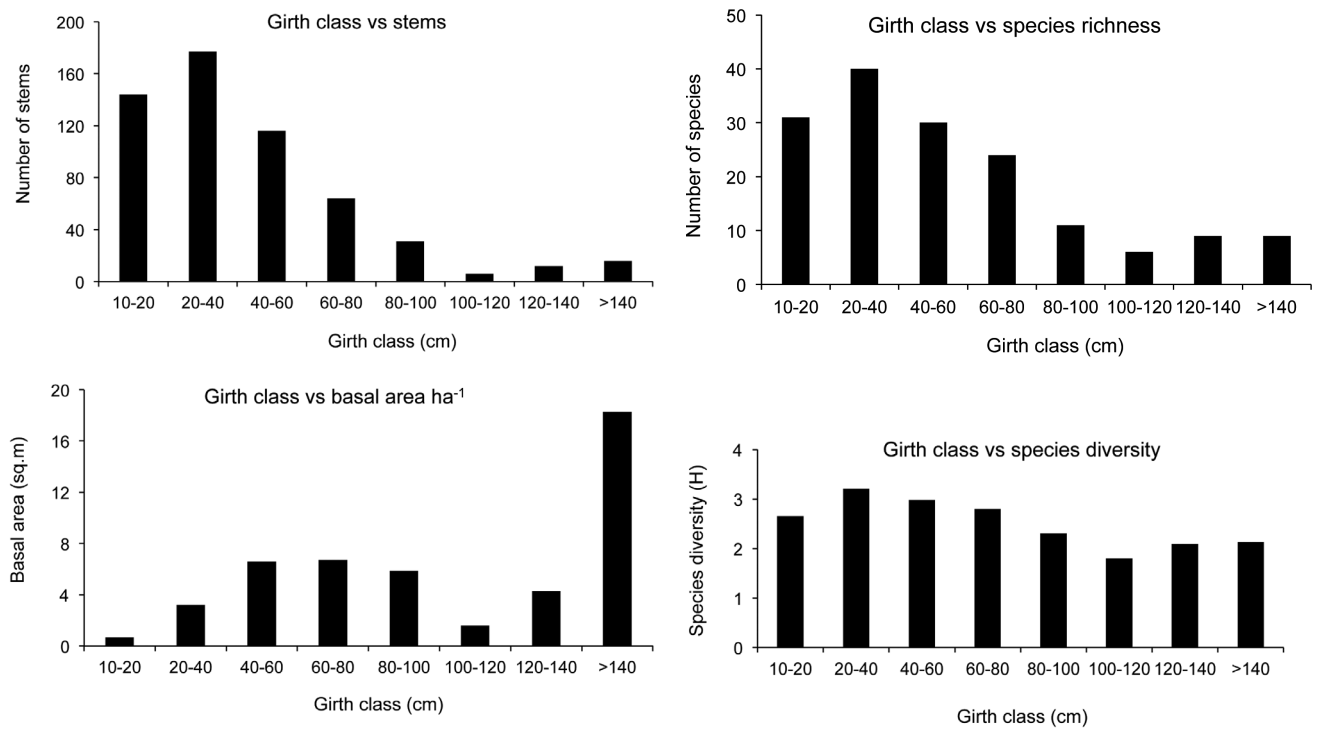


Fig. 4. Stem density, basal area, species richness and diversity among the girth classes in the Kondapalli Reserve Forest

show the negative binomial distribution, while these with random pattern follow a Poisson distribution. Generally, clustered distribution is common in nature (Odum 1971), compared to random, and is also a function of spatial scale (Hurlbert 1990). According to Leps & Kindlmann (1987), a random pattern is mostly exhibited by mature plants, though they tend to show a cluster pattern at their seedling stage. It is assumed that the neighborhood competition among the seedlings finally changes a dispersion pattern from cluster to random. This indicates that the spatial patterns of individuals are dynamic and change at varying spatial scales, as well as at different developmental stages (Yi *et al.* 2008).

3.1.5. Tree girth analysis

An assessment of tree diameter distribution often reflects the disturbing effect (Denslow 1995; Ramirez-Marcial *et al.* 2001), as well as resource utilization by species within a forest (Hitimana *et al.* 2004). In the studied area, species richness, diversity and stem density decreased with increasing girth class, except in 10-20 cm, 100-120 cm and 120-140 cm girth classes (Fig. 4). This could be due to preferential logging of these girth class woods by local people. Such type of logging modifies forest structure, species composition and diversity (Smiet 1992; Cazzolla *et al.* 2015). About 77% of the recorded stems were from lower girth classes of 10-60 cm. The tree girth exhibited positively skewed asymmetrical distribution within the studied population.

This represents the forest as mature and expanding type with a high contribution of trees from the lower girth classes. Relatively high species richness (87%) and diversity (3.2) were found in 20-40 cm girth class (Fig. 4). Interestingly the girth class of 100-120 cm, comprised six species represented by six individuals. In terms of basal area, no particular trend was observed, indicating low values of girth classes similar to stem density. Overall, this analysis highlights the signs of anthropogenic disturbances in the study area by selective logging of wood.

3.1.6. Cluster analysis

Overall 36 quadrats were clustered into 7 groups with varied species composition based on their species similarity distance (Fig. 5). Cluster 1 and 5 are represented by a single quadrat, while 13 quadrats were grouped under cluster 3, forming a large group. Cluster C1 is represented by *Atalantia monophylla* and *Lanea coromandelica*, C2: *Atalantia monophylla* *Strychnos potatorum*, *Wrightia tinctoria*, C3: *Gyrocarpus americanus*, *Commiphora caudate*, C4: *Albizia amara* and *Givotia moluccana*, C5: *Givotia moluccana* C6: *Atalantia monophylla*, and *Chomelia asiatica*, and C7: *Albizia amara*, *Atalantia monophylla* and *Strychnos potatorum*. Finally, it can be interpreted that *Atalantia monophylla* has high ecological amplitude with a wide distribution in the study area, associated with different species in KRF.

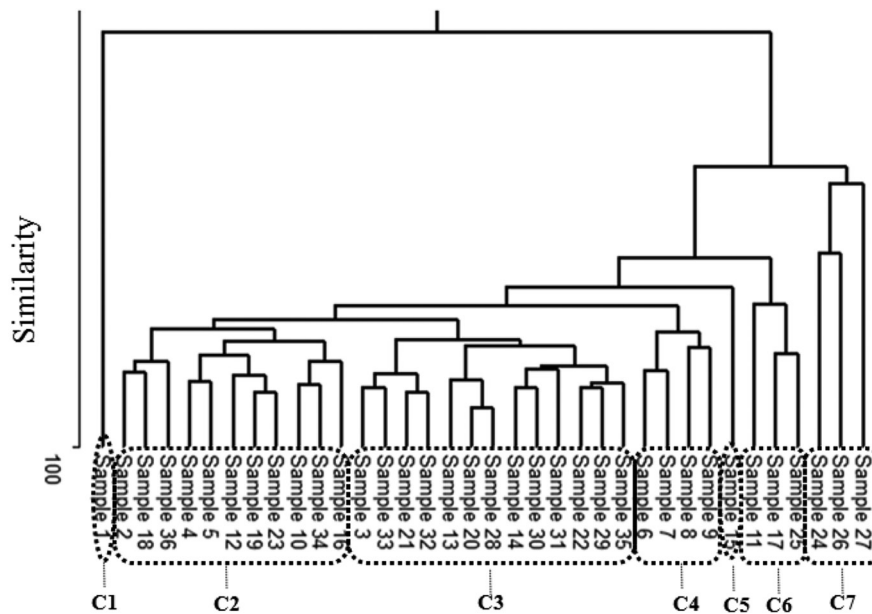


Fig. 5. Demarcating dominant species communities of the Kondapalli Reserve Forest using cluster analysis

3.2. Is it apt to compare ecosystems growing under dissimilar environmental conditions?

Edaphic and climatic factors, along with topography, influence species richness, diversity and dominance of an ecosystem beside human interferences (Huang *et al.* 2003; Prasad *et al.* 2007; Amissah *et al.* 2014). Also, as stated by Denslow & Hughes (2004), species richness varies depending on the dominant species in a community. Every ecosystem on the earth is unique by itself and the current climax communities observed globally are the summation of natural and anthropogenic interactions. All the ecosystems across the world have been subjected to such interferences, and the only difference is the level of interactions, some might have encountered high and others low. Ultimately, such interactions stabilize the species richness and diversity of a given ecosystem.

So far, in the traditional ecological studies, researchers worldwide, due to an oversight, often compare species diversity parameters of one forest with another forest. In some cases, the comparison is made between evergreen and deciduous systems that are totally different both in terms of their growing conditions as well as species composition. There is also variation in the method of sampling (transect/quadrat; random or contiguous, temporary or permanent plot), sampled area, tree girth size measured, as well as the season, but despite these differences, these ecosystems are compared. The crucial comparison involves species diversity. Most of researchers globally adopt the Shannon-Wiener Index for calculation of species diversity.

$$H = - \sum_{i=1}^S p_i \ln(p_i) \quad \text{Or} \quad H = - \sum_{i=1}^S (ni/N) \ln(ni/N)$$

However, when comparing the results, sufficient attention is not always given to whether the diversity was calculated using natural logarithm (\ln), \log_2 or \log_{10} (Prasad & Rajan 2014). Sometimes, it is even more confusing, e.g. Padalia *et al.* (2004) described the Shannon-Wiener index formula as:

$$H = - \sum [(ni/N) \log_2 (ni/N)] \quad (\log \text{ implies to log base 10})$$

This formula does not provide any clue whether authors used \log_2 or \log_{10} . The results of this study were compared by Shruthakeerthiraja & Kumar (2012), who used \ln for diversity calculation (see Prasad & Rajan 2014 for more details). Depending on the \ln and the log base (2, 10) value diversity values differ for the same region. For example, in the present study diversity value is 3.2 (using \ln) and 4.6 (using \log_2).

Similarly, proper attention should be paid when comparing stem density and basal area – these attributes mostly depend on the availability of resources, dispersal capacity of the species, their spatial dispersion patterns, topography and, more specifically, logging of stems by humans or any other disturbance factor. In a region where there is high pressure of anthropogenic interference for timber products, the stem density will be obviously low. So while making comparison with other forest systems, it is essential to understand the disturbance factor that actually gives low or high values.

This type of comparative evaluations does not seem to be appropriate, because environmental conditions are different for different forest types. However, despite such a variation, it has become a custom in the diversity/ecological analysis. Even with the same forest

Table 3. Species richness and diversity parameters in different tropical dry evergreen forests of southern India

No.	Location	MAT (°C)	MAR (mm)	Plot size	Girth (cm)	S	H	Stems ha ⁻¹	BA m ² ha ⁻¹
1	*Kondapalli RF	28.5	1067	36 plots of 0.01 ha (10×10 m)	≥ 10	46	3.2	1572	47.17
2	^a Guindy National Park	---	---	200x200m quadrat area In each 80 quadrats of 5×5m	≥ 20	31	2.94	---	---
3	^b Marakkanam RF	---	1254	Three 0.1 ha (20×50 m)	≥ 20	---	0.83- 2.43	280	11
4	^b Puthupet SG	---	"	Two 0.1 ha	"	---	1.47- 1.59	1130	36
5	^c Suriyampettai	---	---	---	---	28	1.61	---	21.54
6	^d Puthupet	---	---	Four 0.5 ha plots. 100×50 m (10×10 m)	≥ 10	51	2.28	1338	32.78
4	^e Kuzhanthaikuppam	28.5	1378	Two 1-ha (100×100 m) plots, divided into 10x10 m	≥ 10	42	2.35	1367	15.44
5	^e Thirumanikkuzhi	"	"		"	38	2.57	974	29.48
6	^f Oorani	---	1373	25 plots of 20×20 m adding to 1 ha in each site	≥ 20	30	2.08	1070	25.55
7	^f Olagapuram	---	"		"	21	2.42	953	4.31
8	^f Oorani	28.5	1311	One ha plot (100×100 m) subdivided into 100, 10×10 m quadrats	"	31	1.82	2815	17.63
9	^f Olagapuram	"	"	Two 200×25m plots, divided into 40 (10×10 m) and 20 (5×10 m) quadrats	"	30	2.33	1286	27.3
10	^h Araiypatti (LD)	28.5	1378		≥ 10	35	2.44	8.7	19.1
11	^h Karisakkadu (MD)	"	"		"	30	2.24	596	21.6
12	^h Maramadakki (HD)	"	"		"	28	2.01	724	15.5
13	^h Shanmuganathapuram (MD)	"	"		"	26	1.29	1663	22.1
14	^h Rayapatti (MD)	"	"	1 ha permanent plot (100×100 m) in each site, divided into (10×10 m)	"	19	1.84	886	12.4
15	^h Puthupet (HD)	"	"		"	30	1.64	1567	36.5
16	^h Oorani (MD)	"	"		"	29	2.33	1284	27.3
17	^h Arasodikuppam (MD)	"	"		"	30	1.82	2813	17.6
18	^h Kuzhanthaikuppam (MD)	"	"		"	28	2.02	1349	16.9
19	^h Thirumanikkuzhi (MD)	"	"		"	22	2.06	1077	29.3
20	ⁱ Kuzhanthaikuppam (1995-2005)	28.5	1378	Two 1-ha (100×100 m) plots, subdivided into 10×10 m subplots	≥ 10	26- 24	2.07- 2.14	1229- 1032	14.6- 14.9
21	ⁱ Thirumanikkuzhi (1995-2005)	"	"		"	26- 22	2.23- 2.11	832- 978	28.9-27
22	^j Araiypatti (HD)	29.5	1033		≥ 10	37	2.56	705	19.43
23	^j Karisakkadu (MD)	"	"	Four 1 ha permanent plots (100×100 m); 10×10 m	"	31	2.52	596	20.26
24	^j Maramadakki (MD)	"	"		"	27	2.11	750	18.63
25	^j Shanmuganathapuram (HD)	"	"		"	29	1.54	1182	20.38
26	^k Puducherry, Villupuram, Cuddalore, Pudukottai				≥ 10				
27	RUD	32.9	1282		"	69	3.13	193 - 274	2.82 - 7.62
28	MD	---	---	100 plots of size 10×20 m in four disturbance classes	"	57	2.8	124 - 297	1.73 - 9.76
29	MU	33.6	1079		"	54	2.99	90 - 208	1.75 - 5.42
30	HD	33.4	1033		"	46	2.24	62 - 109	0.55 - 3.98
31	^l Suryanpet	29.5	1141		≥ 10	26	2.13	771	32.55
32	^l Velleripet	"	"	One ha plot in each site, divided into 10×10 m	"	18	1.24	1144	47.84
33	^l S. Pudhoor	"	"		"	25	1.3	1145	17.74
34	^l Chinna Kumatti	"	"		"	27	2.2	1285	36.7
35	^m Puthupet	"	"	Four 0.5 ha (100×50 m; total 2 ha)	≥ 10	23	---	1329	37.5

36	^a Vidathudaiyar SG	40	2043		≥ 30	65	1.9	14	7.72
37	^a Koodaiyakkaruppar SG	"	"		"	55	1.69	8	6.55
38	^a Thiruparkkadal Chellayae Amman SG	"	"	1 ha plot	"	68	2.33	15	12.31
39	^a Aakkamudaiyar SG	"	"		"	68	2.28	12	6.87
40	^o Northern Eastern Ghats	---	---	17 plots of 0.1 ha randomly selected	≥ 10	135	5.2	266	8.65
41	^o Southern Eastern Ghats	---	---	47 plots of 0.1 ha randomly selected	"	365	6.2	334	11.1

Explanations: * – current study area, a – Rajarathinam (1990), b – Visalakshi (1995), c – King (1997), d – Parthasarathy & Sethi (1997), e – Parthasarathy & Karthikeyan (1997), f – Ramanujam & Kadamaban (2001), g – Venkateswaran & Parthasarathy (2003), h – Mani & Parthasarathy (2006), i – Mani & Parthasarathy (2009), j – Pandian & Parthasarathy (2013), k – Anbarashan & Parthasarathy (2012), l – Anbarashan & Parthasarathy (2013), m – Baithalu *et al.* (2013), n – Sundarapandian & Subbiah (2015), o – Babar *et al.* (2011) (except the current study area and Babar *et al.* (2011), all other studies have been carried out along the Coromandel coast of southern India – Tamil Nadu and Pondicherry), MAT – mean annual temperature, MAR – mean annual rainfall, S – species richness, H – Shannon diversity index, RF – reserve forest, SG – sacred groves, RD – relatively undisturbed, LD – least disturbed, MD – moderately disturbed, MU – mostly disturbed, HD – highly disturbed, BA – basal area

type, the floristic elements differ based on the regional eco-climatic conditions. Currently, there is a need for a system of research that exclusively works on providing in-depth insights about richness, diversity and other floristic elements that can show a clear picture of forest under study, together with its current condition. This will help to analyze and comprehend the relationship between diversity parameters in relation to environmental conditions and, further, for conservation implications that may be the set goal of a study.

3.3. Diversity patterns of other dry evergreen forests of southern India – status of KRF

In view of above discussion, in the current study, we did not compare our results with any other forest types. However, we made an attempt to show diversity parameters of only dry evergreen forests (similar type) exclusively restricted to southern India, which has, to a certain extent analogous climate and environmental factors (Table 3). Compared to other forest types, dry evergreen forests are least studied within the country. They are found as scattered and patchy structures of the southern India, particularly towards the eastern coast, Tamil Nadu (Meher-Homji 1974; Parthasarathy *et al.* 2008).

As shown in Table 3, there are several discrepancies when comparing diversity and species richness between different dry evergreen forests, such as: (1) temperature and precipitation among sites ranged between 28.5 to 40 °C and 1033 to 2043 mm; (2) the sampled area and its segmentation for inventory varied – most researchers adopted a contiguous plot of diverse dimensions, except Babar *et al.* (2011) and the current study. In comparison to contiguous sampling, a random quadrat survey conducted in heterogeneous areas captures good species richness (Prasad *et al.* 2007). This also has an impact on the species diversity, stem density (ha^{-1}) and basal area ($\text{m}^2 \text{ha}^{-1}$); (3) in most of the studies, there were sampled

trees with $\text{DBH} \geq 10$ cm, while in few with $\text{DBH} \geq 20$ and ≥ 30 cm; (4) only few studies have been carried out in disturbed forests; (5) in the study of Babar *et al.* (2011), \log_2 was used for diversity calculation, hence, their values are high compared to other studies that used natural logarithm.

Based on these variations, it is not appropriate to compare results and conclude that an area's diversity is low to one study and high to another study. For example, if species richness is considered, out of 41 case studies shown in Table 3, KRF richness is high compared to 31 sites, and low with respect to the rest of the sites, but there is variation in the area sampled, as well as the girth size considered during sampling. With respect to the number of stems ha^{-1} , KRF stood third after the study sites 8 and 17, and with reference to basal area ($\text{m}^2 \text{ha}^{-1}$) was second after the site 32, showing high values for both the parameters compared to the site 41, where large area was sampled. The diversity values across sites cannot be compared due to variation in the log values used in different studies. There are a number of factors that actually influence the diversity parameters of an ecosystem and, thus, it is not appropriate to normalize the sites and compare. However, it is apt to compare diversity parameters of the same area at temporal intervals, like in the studies of Mani & Parthasarathy (2009), Baithalu *et al.* (2013) and Pandian & Parthasarathy (2016). In these studies, they showed the differences in species richness, diversity, and other parameters at an interval of 10 years. Such type of studies helps in assessing the species recruitment, loss, succession and other ecological processes concurrent with human pressure and in proposing future conservative measurements.

4. Conclusions

The KRF, a tropical dry evergreen forest of southern India, is one of the remnant forest patches in the vicinity

of Vijayawada city that is prone to severe anthropogenic disturbances. The current study is first of a kind for the KRF with respect to species diversity analysis. This study provides baseline data about floristic structure and diversity of KRF for future temporal analysis and, further, for the reserve protection and conservation measurements. The study also argues not to compare diverse forest ecosystems with respect to their species richness and diversity patterns. It is suggested that researchers should come out of traditional way of comparing dissimilar ecosystems and should design a new approach of describing and understanding diversity of forest ecosystems.

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