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**GENOTYPE×ENVIRONMENT INTERACTION IN
HEDGE LUCERNE (*Desmanthus virgatus* (L.) Willd.) FOR
YIELD AND QUALITY**

by

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(2016-11-034)

THESIS

**Submitted in partial fulfilment of the
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2018

DECLARATION

I, hereby declare that this thesis entitled "**GENOTYPE×ENVIRONMENT INTERACTION IN HEDGE LUCERNE (*Desmanthus virgatus* (L.) Willd.) FOR YIELD AND QUALITY**" is a bonafide record of research work done by me during the course of research and the thesis has not previously formed the basis for the award to me of any degree, diploma, associateship, fellowship or other similar title, of any other University or Society.

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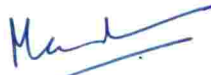
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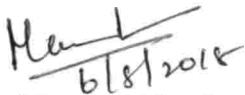
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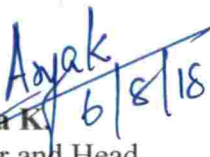
We, the undersigned members of the advisory committee of Mr. Arun Chacko a candidate for the degree of **Master of Science in Agriculture** with major in Plant Breeding and Genetics, agree that the thesis "**GENOTYPE×ENVIRONMENT INTERACTION IN HEDGE LUCERNE (*Desmanthus virgatus* (L.) Willd.) FOR YIELD AND QUALITY**" may be submitted by Mr. Arun Chacko, in partial fulfilment of the requirement for the degree.


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LIST OF ABBREVIATIONS AND SYMBOLS USED

%	Per cent
G x E	Genotype x Environment
CD	Critical Difference
C.V	Coefficient of variation
cm	Centimetre
mg	Milligram
RBD	Randomised Block Design
Cluster ⁻¹	Per cluster
DAS	Days After Sowing
°C	Degree Celsius
g m ⁻² day ⁻¹	gram per meter square per day
g dm ⁻² day ⁻¹	gram per decimetre square per day
<i>et al.</i>	And others
Fig.	Figure
g	Gram
g ⁻¹	Per gram
Kg	Kilo gram
ha ⁻¹	Per hectare
KAU	Kerala Agricultural University
t ha ⁻¹	Tonne per hectare
kg ha ⁻¹	Kilogram per hectare

kg plot ⁻¹	Kilogram per plot
Plant ⁻¹	Per plant
Plot ⁻¹	Per plot
Day ⁻¹	Per day
<i>Via</i>	Through
Mm	Millimetre
No.	Number
CGR	Crop Growth Rate
cv	Cultivar
NAR	Net Assimilation Rate
Sl.	Serial
sp. or spp.	Species (Singular and Plural)
LAI	Leaf Area Index
<i>viz.</i>	Namely
<i>i.e</i>	that is
d.f	Degrees of freedom
S. E	Standard Error
mg g ⁻¹	Milligram per gram

INTRODUCTION

1. INTRODUCTION

Desmanthus virgatus commonly known as hedge lucerne is a perennial shrub legume belonging to the family Fabaceae and subfamily Mimosoideae. It is a native of tropics and subtropics of new world. The fodder can be safely fed to ruminants and non-ruminants as it is palatable, aggressive, persistent and tolerant to grazing. It contains high condensed tannins and is devoid of toxicants like mimosine.

Hedge lucerne is a forage legume which is preferred by cattle for its palatable green fodder and adequate amount of crude protein (Deepthi *et al.*, 2013). Comparative evaluation of chemical composition (dry matter percentage, crude protein, calcium and phosphorous) of hedge lucerne with other tropical and sub-tropical forage legumes reveal it to be a nutritious feed (Johri *et al.*, 1987). The pithy stem of the fodder makes it easier to harvest and frequent cuts can be taken. Hedge lucerne is observed as potential fodder legume that can substitute leucaena for ruminant feed due to its versatile nature.

In Kerala, only 40 per cent of green and dry fodder requirement is roughly met from the available feed resources (NDDB, 2016). To narrow the demand supply gap in feed and fodder, genetic improvement of fodder crops with regard to high yield and quality is essential. Many fodder crops are under-utilized and their cultivation is reduced due to the fluctuant performance of diverse varieties. Being a promising fodder legume with ample advantages, hedge lucerne demands genetic improvement in terms of productivity. Information on adaptability and stability with regard to performance of genotypes can be drawn from the analysis of interaction of genotypes with locations and other agro-ecological conditions.

The phenotypic expression of a character is influenced by the genotype and environmental factors. The correlation between genotype and phenotype get reduced in the presence of G x E interaction which makes it difficult to assess the genetic potential of a particular genotype whose relative ranking will be altered in different environments. This forges the need to determine G x E magnitude in

varying environments to identify the stable genotypes. Several methods were proposed to analyze G×E interaction and to determine the stability in performance of genotypes (Becker and Leon, 1988). The linear regression model proposed by Eberhart and Russell (1966) is the most commonly used method for analysis of genotype x environment interaction. Development of improved varieties showing stable performance across wide environmental conditions is a necessity to increase the productivity. Currently, there is a need to develop and identify hedge lucerne genotypes with higher yield potential.

In this perspective, the present investigation was conducted across four locations in Kerala to study the genotype x environment interaction in hedge lucerne for yield and quality.

The main objective of this study was

- ❖ To identify stable genotypes of *Desmanthus virgatus* in varied environments with respect to yield and quality.

REVIEW OF LITERATURE

2. REVIEW OF LITERATURE

Hedge lucerne (*Desmanthus virgatus* (L.) Willd) is a tropical forage legume native of tropics and sub tropics of new world. The word 'Desmanthus' originates from the Greek terms 'desme' meaning bundle and 'anthos' meaning flower.

Hedge lucerne belongs to the family Fabaceae, subfamily Mimosoideae. It's a small shrub, 2-3 m tall and roughly resembles *Leucaena*. No poisonous principle is observed in the foliage (KAU, 2016).

Being a palatable, aggressive, persistent, grazing tolerant perennial browse shrub free from anti-nutritional factors and from which frequent cuts can be taken, it's a potential forage legume. However the genetic improvement achieved in terms of its productivity is very low. Hence this investigation aims at the study of genotype-environment interaction in hedge lucerne genotypes under varied climatic conditions to identify stable genotypes with respect to yield and quality.

The plant is a herbaceous perennial, branched and suffruticose at the base with a height upto 0.5 to 2.5 m. It posses deep woody tap root. The stems are slender, pithy in center, angular, green turning brown. The leaves are 2-8 cm long, compound, bipinnate, bearing 10-25 pairs of linear-oblong leaflets. The leaves posses nyctinastic leaf movements. The inflorescence bears 9-11 whitish mimosoid flowers. The fruits are linear, dehiscent, 5.5-8.5 cm long pods which contain 11-26 reddish brown or golden brown U-shaped seeds (Gutteridge and Shelton, 1994).

2.1. GROWTH, YIELD AND QUALITY ATTRIBUTES OF HEDGE LUCERNE

The comparison between the lucerne and hedge lucerne revealed that optimum biomass, dry matter and crude protein were obtained at 45 days cutting interval for hedge lucerne. Thus hedge lucerne can be considered as a better substitute for lucerne due to its versatile nature (Khan and Bose, 1994).

Shanthi (1995) studied the genetic variability in forty types of hedge lucerne (*Desmanthus virgatus* (L.) Willd) and observed the maximum plant height of genotypes in first cutting followed by a reduction in the second cutting. A linear increase in number of branches, leaf to stem ratio, crude protein and crude fibre was observed from first cutting to second cutting. A general reduction in green fodder yield from first cutting to the second cutting was also observed.

The effect of height and frequency of cutting on yield, quality and persistence of hedge lucerne (*Desmanthus virgatus* (L.) Willd) accession IRFL 1857 over two years. Plants harvested every two weeks at a stubble height of 100 cm gave the highest leaf yield and total dry matter (Trujillo *et al.*, 1996)

Suksombat and Buakeeree (2006) conducted an experiment to determine the effects of cutting interval and cutting height on the yield and nutrient composition of hedge lucerne (*Desmanthus virgatus* (L.) Willd). They found that significant increase in dry matter and nutrient yield were observed with increased cutting of hedge lucerne stand every 40 to 50 days. The cutting height during harvest had no effect on dry matter or nutrient content.

The biomass yield, palatability, chemical composition and nutritive value of hedge lucerne (*Desmanthus virgatus* (L.) Willd) in sheep was studied and obtained a biomass yield of 39.81 t ha⁻¹. The mean crude fibre and crude protein content were 19.77% and 15.20% respectively. The experiment revealed that hedge lucerne can be used as a potential leguminous fodder for small ruminants (Radhakrishnan *et al.*, 2007)

The plant height of hybrid napier exhibited significant differences between the varieties. Growth was slow during the initial stages as evidenced by low plant height. The maximum plant height was observed during the fourth harvest and a gradual decrease in further harvests. Crude fibre content did not show any significant differences among different cultivars. With respect to leaf area index, green fodder yield, dry matter yield, crude protein content, relative growth rate

and net assimilation rate significant differences were observed between the cultivars. Green fodder yield was comparatively less during the initial harvests and highest yield was observed during the third harvest (Soumya, 2011).

Deepthi *et al.* (2013) conducted a study on genetic divergence and association analysis in hedge lucerne (*Desmanthus virgatus* (L.) Willd) in which twenty mutants along with control of variety TNDV-1 were evaluated for twelve traits. The simple correlation estimates revealed that green fodder yield plant⁻¹ was significant and positively correlated with number of branches plant⁻¹, leaf to stem ratio and dry matter yield plant⁻¹. The path analysis results showed the high positive direct effect of dry matter yield on green fodder yield plant⁻¹ followed by number of branches plant⁻¹, plant height, leaf to stem ratio and pods cluster⁻¹.

An effect of date of sowing and cutting intervals on growth attributes and yield in lucerne (*Medicago sativa* L.) was evaluated under North Gujarat agro-climatic conditions. Significantly higher plant height, mean number of leaves plant⁻¹, mean leaf area plant⁻¹ and higher dry forage yield was observed by 30 days cutting interval after common cut (Kumar and Patel, 2013).

Shashikanth *et al.* (2013) evaluated the performance of eight guinea grass varieties in Southern dry zone of Karnataka during *kharif* season over three years. From the pooled data, significantly higher green fodder yield (1007.04 q ha⁻¹year⁻¹), dry matter yield (147.72 q ha⁻¹year⁻¹), crude protein yield (12.99 q ha⁻¹year⁻¹) along with growth parameters like plant height (78.47 cm) and leaf to stem ratio (0.71) were recorded in the variety JHGG-08-1.

Jindal *et al.* (2015) conducted a varietal evaluation trial on three entries of lucerne (*Medicago sativa*) namely, BAIF Lucerne, Anand-21 and Anand 22 along with two national checks viz., RL-88 and Anand-2 under three different agro-ecological zones for assessing quality parameters and forage yield potential. The green fodder yield, dry fodder yield and crude protein content pooled over three years recorded highest at Coimbatore in South zone.

Ishrath (2016) conducted a study on cutting intervals and additives for quality silage production using hybrid Bajra Napier variety Suguna. The cutting interval 75 days recorded highest green fodder yield. The highest crude protein (10.56%) and the lowest crude fibre content (26.81%) were observed from 45 days cutting interval.

Green fodder yield, chemical and nutrient composition and nutrient uptake potential of fifteen sorghum varieties belonging to sweet sorghum, dual purpose and forage types under double cut system with 75 days interval were studied. Green fodder yield showed significant difference among the sorghum varieties. The highest green fodder yield (71.28 t ha⁻¹) was obtained from sweet sorghum variety CSH 22 SS which was on par with forage varieties HC 308 and CSV 21F. Forage variety HJ 513 was observed with high green fodder yield (69.48 t ha⁻¹). For dry matter yield and crude protein content, non-significant differences were observed among the sorghum varieties. For crude protein, significant differences were obtained among the varieties. SPV 462 recorded the highest crude protein percentage (7.08 %) which was on par with varieties Phule Revati, CSV 19 SS, Pant Chari 5 and CSV 15. Crude fibre percentage recorded the highest in HJ 513 (33.11%) and the lowest in CSV 27 (22.71%). Forage varieties showed significantly higher values in dry matter percentage. Dry matter percentage was observed higher in SSG 898 (33.11 %) and the lowest in CSH 22 SS (25.91 %) (Singh and Chauhan, 2017).

Forage nutritional quality of ten Bajra x Napier hybrids (B x N hybrids) along with their eleven parents and four checks were assessed. Hybrids GB x FD-444 and GB x FD-436 recorded the highest dry matter per cent. Highest crude protein per cent (9.63 %) was recorded from male parent TNCN-011 and check CO -3 (Gate *et al.*, 2018).

2.2. GENOTYPE X ENVIRONMENT INTERACTION AND STABILITY ANALYSIS

Plant breeders aim at developing new varieties with high yield stability over wide range of environments. Genotype-environment interaction is one of the basic reasons for the difference in high performance in genotype for the yield and other essential agronomic traits.

Genotype of an individual is its genetic constitution while phenotype is the observable characteristics and phenotypic value is the observed value of a particular characteristic. Phenotypic value of an individual is affected partly by the genotype and partly by the environment which it grows.

An environment is referred as all the external agencies which determine the performance at the phenotypic level. According to Comstock and Moll (1963), the environment can be micro and macro. A potential environment which changes over locations or time is referred as a macro-environment while micro-environment is the special environment circumstances confined to an individual which have very small contribution to the genotypic expression. Environmental variation can be divided as predictable and unpredictable (Allard and Bradshaw, 1964). All the permanent characters of an environment such as climate, soil type along with characters which show fluctuation in a systematic manner can be referred as predicable environment. Unpredictable environment includes changes that are uncontrollable such as the amount and distribution of rainfall in a given area, temperature and atmosphere. Only the macro-environment and its interaction with genotype can be studied since it has major contribution to the final expression of a genotype.

The performance of a genotype with respect to changing environmental factors over time within a given location is referred to as stability of a genotype whereas adaptability refers to the stability in performance of genotypes with respect to changes across locations. Thus a stable variety is less sensitive to the environmental changes taking place.

Genotype-environment interaction is the interplay of genetic and non-genetic effects causing differential relative genotypic responses in different environments. The presence of genotype-environment interaction can be confirmed from the analysis of variance developed by Sprague and Federer (1951). Other approach makes use of regression technique in which G x E interaction is partitioned into linear and non-linear components. Ecovalence measure is the contribution of a genotype to the interaction sum of squares (Wricke, 1962). The genotype with least ecovalence is considered most stable.

Differential responses of improved genotypes to change in environments were initially interpreted by Finlay and Wilkinson (1963). In this approach the components of a genotype and environment interaction were linearly related to environmental effects when measured on small scale as the genotypic effects. An improvement in Finlay and Wilkinson model was done by Eberhart and Russell (1966) by adding a stability parameter, which shows the deviation from regression. According to Eberhart and Russell (1966), a stable genotype exhibits high mean yield, regression coefficient around unity and deviation from regression near zero. Shukla (1972) used the term 'stability variance' an estimate of variance in terms of the residuals in a two-way classification for indentifying the stability of a particular genotype.

Mini (1989) conducted a comparative study of genotype environment interaction in sesame grown over three different locations. Stability analysis methods of Eberhart and Russell (1966), Perkins and Jinks (1968), Freeman and Perkins (1971), Wricke (1962) and Shukla (1972) were used. By examining the stability parameters, variety IS 284 was found to be more stable for most of the characters.

Moneim *et al.* (1990) evaluated twenty-five types of forage peas (*Pisum sativum* L.) across four locations for two years to study the genotype-environment interaction for herbage and seed yield under rainfed conditions. The

accession 88/335 was identified for wide adaptation and stability for high herbage yield and seed yield.

Jyothi (2002) evaluated eight Kunjukunju rice cultures across three locations of Palghat for identifying the stable culture. For many of the yield and yield related traits, Kunjukunju rice culture K-6 was identified as stable in different environments. For many yield traits, variety Kanchana was found specifically adapted only to favourable environments. Nine rice cultures from F₆ generation of wide crosses were evaluated across three locations to study the stability for characters. For most of the yield contributing traits, Culture C 26T (b) was stable over three locations (Palathingal, 2003).

Fourteen mutants of coleus were evaluated across four locations of Kerala for analysing the stability. Significant differences in most of the economic traits were observed among the selected mutants. Mutants '641' and '352' were identified as high yielding and well adapted over locations for many of the economic traits (Shinoj, 2003).

Seven commercial rice hybrids and two check varieties were used for genotype x environment interaction study across three farming situations of Kerala during *kharif* season. Based on the stability analysis, the hybrid KRH-2 was well suited under poor management conditions as it recorded highest mean value and regression coefficient less than unity. Hybrid NSD-2 was identified well suited for better management conditions (Chandrashekhar, 2004).

Sastry (2005) studied sixteen isabgol genotypes across three environments for analysing the stability using Eberhart and Russell (1966) method. Significance for linear component of G x E interaction was observed in number of effective spikes, seed yield and disease index. For the non-linear components, days to 50 per cent flowering and husk content was found to be significant. Genotypes RI-9809, EC-124345 and DM-4 were identified stable for all environments. Genotypes DM-2 and Niharika were stable for seed yield.

Genotype GI-2 was found to be unstable but having more husk than other genotypes.

To determine genotype-environment interaction and stability, fourteen cotton genotypes were evaluated across four locations. Significant differences in mean yield over the environments were observed. Significance in deviation from regression was found only for four genotypes and regression coefficients ranged between 0.23 to 1.46. Genotypes SG-1001, SG-125 and DLP-5409 which are high yielding were identified as stable genotypes (Killi and Harem, 2006).

Ten varieties of cowpea were raised in four environments to assess the selection techniques in genotype-environment interaction. Based on the environment index best and poorest environment was selected. Significant difference in effects of genotype and environment were obtained. Presence of genotype-environment interaction was confirmed from the joint regression analysis. Regression coefficient, non-parametric statistic, superiority measure can be used together to select genotypes based on yield and environmental response (Aremu *et al.*, 2007).

Marimuthu (2007) studied the genotype x environment interaction in eight selected New Plant Type (NPT) lines of rice along with Jyothi as check variety across three low land rice ecosystems. NPT-7 been had identified with better performance based on performance for yield, agronomic characters, grain characteristics and stability.

For analysing the stability in Black gram, twenty genotypes were evaluated over three environments. Eberhart and Russell (1966) method and genotype grouping technique by Francis and Kannenberg model (1978) were used to analyse the data. Higher mean seed yield, average responsiveness to season and stability was recorded in the genotypes VBG 89, VBG (Bg) 4 and VBG 62 (Shanthi *et al.*, 2007).

Seventeen genotypes of brinjal were evaluated over four environments for stability analysis. Mean squares due to genotypes and environments were significant for all the characters studied. Non-significance in genotype x environment (linear) mean sum of squares for most of the characters showed that variation in performance of genotypes is entirely unpredictable. The genotypes PB-66, PbS, PB-67, PB-60 and PB-4 were found to be stable for number of fruits per plant and yield per hectare (Bora, 2010).

Jagjivan (2010) compared different methods for stability analysis in rice (*Oryza sativa* L.) and observed that Eberhart and Russell model (1966) was on par with latest AMMI model. For predicting the stability in performance of a genotype, Eberhart and Russell model (1966) was found preferable than other methods.

Haridas (2011) evaluated stability of fifty ground nut genotypes having bunch type growth habit over three environments for pod yield per plant and its related characters. Pooled analysis showed significance in mean square values due to genotypes. Mean squares due to G x E interaction were significant for all the characters except oil content. Mean squares due to environments were significant for all the characters in the pooled analysis over environments. Environment E3 was observed with high performance for most of the characters.

Singh (2011) evaluated thirty chilli genotypes along with three checks across three different environmental conditions to study the genotype x environment interaction. Around seventeen genotypes were found stable for yield. Genotype PC-2507 showed the highest yield and identified as the most stable genotype. Indo sem, PC-10 and AC-150 were the genotypes found to be specifically adapted to favourable environment.

The Genotype x Environment study in aromatic rice to analyse the stability over three environments depicted that genotype GT 6 had more adaptable characters over the three environments. More stable grain characters were observed in genotype GT 1 over the two locations (Ram, 2012).

Bikash *et al.* (2013) evaluated 30 hybrids of pearl millet during *kharif* season across four locations for dry fodder yield stability. High significance in mean squares due to genotypes and environments together with G x E interaction from the analysis of variance for stability indicated significant differences among genotypes. The mean squares due to G x E interaction was partitioned into linear and non-linear components and G x E (linear) was predominant for characters like days to 50 percent flowering, plant height and dry fodder yield and the performance of which could be predicted across the environments. The estimation of environmental index helped in selection of the most favourable environment for all the characters as E₁ (irrigated condition). The hybrids found to be stable over the environments are 94111A x 1250 and 96111A x (G73-107 x bsectap 1). For the poor environment, hybrid ICMA97444 x ICMR0/035 was the best suited. The hybrids studied did not exhibit uniform pattern of environment response (linear). For identifying stable hybrid by selection, genotypes with average response for different characters and phenotypic stability of characters should be given importance.

Singh and Arya (2014) evaluated thirty-six genotypes of *Vigna radiata* (L.) to analyse the stability under three environments using methodology of Eberhart and Russell model (1966). The genotype-environment interaction effect was analysed using the additive mean effect and multiplicative interaction effects (AMMI) model. Two genotypes G2 and G36 with high mean seed yield showed non-significant deviation from regression. The genotype G33 with average mean seed yield showed non-significant deviation from regression. Environment A (early sown) and B (timely sown) were favourable for most of the yield component traits, environment C (late sown) was unfavourable for almost all the yield and yield related traits. For seed yield, genotypes G1, G2, G3, G4, G18, G22, G24 and G25 were found to be stable based on AMMI 1. Based on AMMI 2, genotypes G14, G22 and G25 were found to be stable for seed yield. Based on analysis, genotypes and environments were grouped into nine sectors.

Anarase *et al.* (2015) conducted an experiment with five male sterile lines, fourteen testers, resultant seventy hybrids and two checks of rabi sorghum across three different locations for examining the genotype-environment interaction for yield and yield related traits. The study on stability parameters depicted that eight parents were found to be average stable for grain yield.

Ten black gram cultures along with four check varieties were raised during kharif, rabi and summer seasons under open and shaded conditions for analyzing the stability by using Eberhart and Rusell's model. The data analysed as pooled over open condition as well as shaded condition did not show any variation between genotypes under three seasons for number of seeds pod⁻¹ and none of the genotypes showed stability in protein content. The data pooled over six environments depicted that all the traits showed variation between genotypes under the environments studied. Ranking of genotypes were done based on stability, yield and yield contributing characters and the least ranked genotypes T₆, T₅ and T₃ were recommended for cultivation under open and shaded conditions (Bhagwat, 2015).

Stability analysis study in ten F₁ hybrids and two check varieties of okra over four locations obtained significant differences among genotypes and environments for all the characters in the pooled analysis of variance for evaluation of F₁ hybrids over locations and seasons. The hybrids Thirumala local x Mallapalli local, Thirumala local x Kattakada and Thirumala local x Punjab Phalgani local were stable over different locations and seasons (Gogineni, 2015).

To identify stability for green fodder yield in forage maize (*Zea mays* L.) forty five hybrids and fourteen parents were evaluated during kharif (E₁), rabi (E₂) and summer (E₃). For fresh green stem weight plant⁻¹ and green forage yield plant⁻¹ significance was observed in G x E (linear) and G x E (non-linear). The hybrids with average stability such as IC-170121 x GWC-0511 and African Tall x GWC-0401 would be well adapted over a range of environments. Hybrids such

as IC-130726 x GWC-0512, GM-6 x IC-130693 with below average stability were specifically adapted to favourable environment. The hybrid IC-130726 x GWC-9603 with above average stability was specifically adapted to poor environment (Nanavati, 2016).

Preeti *et al.* (2016) conducted a study on effects of changing environments on wheat dry matter yield. Forty-two genotypes were grown across four different locations during rabi season to identify the stable genotypes. Non-linear component accounted a major portion of G x E for dry fodder yield plant⁻¹. Environment E₁ was identified as the most favourable environment for all characters from the estimates of environment additive effects. The genotypes WH 1098, WH1126, PBW 343, WH 1081, WH 542 and HD 2851 were identified as stable for dry matter yield in all environments and were more adaptive.

Pangti (2016) examined the stability parameters in thirty bread wheat (*Triticum aestivum* L. em Thell) genotypes for yield and quality attributes in two environments. More than eleven genotypes were found to be stable for grain yield. For both the environments *viz.*, E1 (timely sown) and E2 (late sown), all the characters exhibited significant differences.

Saranya (2016) studied stability analysis in nine neelamari mutants over four locations for yield and indigotin content. In the pooled analysis of variance, significant differences for all the characters were observed among the genotypes studied. This indicated that the genotypes interacted significantly with environments. Mutants It-1, It-2 and It-8 were identified as stable mutants for favourable environments. The stable mutants identified over different locations during the summer season are It-3, It-6 and It-9. Mutant It-10 was the mutant suitable for unfavourable environment.

Baranda (2017) evaluated thirty genotypes of cowpea (*Vigna unguiculata* (L.) Walp) under three environments for analyzing stability using Eberhart and Russell model (1966). Environment-2 was found to be the most suitable for many of the characters based on the mean performance. High consistent

expression for yield character in all environments was observed in genotype CPD-197. Poor yielding genotypes were CPD-115 (Environment E₁, E₂ and on pooled basis) and CPD-127 (Environment-E₃). Wide adaptability under desired environments was shown by the genotype CPD-201. Genotypes CPD-17, CPD-196, CPD-78 and CPD-200 were observed with desired response moisture stress condition.

The existence of genotype x environment interactions and stability were assessed for yield and quality in four Pineapple varieties over seven locations. From the pooled analysis of variance all the characters showed significant difference in genotypic variances over the seven locations. Varieties Amritha and Mauritius were stable for all quantitative and qualitative traits. The stability type was determined based on regression coefficient and mean values (Manivannan *et al.*, 2017).

Mehraj *et al.* (2017) studied genotype-environment interaction in twelve different genotypes of oats (*Avena sativa* L.) for forage yield and its related traits. The genotypes SKO-90, SKO-96 and Sabzaar were identified as stable across the environments. For favourable environments, genotypes SKO-148, SKO-160, SKO-166 and SKO-167 were the most suited while genotype SKO-20 was found to be suited for poor environments for forage yield.

Patil *et al.* (2017) evaluated thirty-seven entries of okra *viz.*, eight parents, twenty eight F₁'s and one standard check over four season for developing stable hybrids for fruit yield plant⁻¹ and its related traits. From the pooled analysis of variance, high significance for the mean squares for genotypes was obtained which indicated the variability for all the characters among the genotypes. Highly significant differences obtained for environments and genotype x environment except for ascorbic acid indicated the divergence among growing environments and differential response of genotypes to various environments. Environment E₂ was found to be most favourable whereas E₃ most unfavourable. Eight hybrids were identified as stable for fruit yield plant⁻¹ and its components.

AMMI analysis was used to study the stability of twenty genotypes of bio-fortified red kernel rice (*Oryza sativa* L.) across three environments and three locations. First interaction principle axis was favourable for all characters whereas second interaction principle axis are favourable for characters such as spikelet fertility, grain yield plot⁻¹(kg), iron content (ppm), protein and amylose content (%) and grain yield plant⁻¹. For grain yield plot⁻¹, grain yield plant⁻¹ and protein content, the genotype RTN-1211-4-2-1-1 was found to be stable. For all the characters across three environments, the genotype RTN-1201-13-2-2-1-32 was found to be most favourable (Rajaram, 2017).

Assessment of stability for forage yield in ten genotypes of *Cenchrus ciliaris* was carried under rainfed conditions for four years. Except green fodder yield at first cut, all other characters were significant for gxe interaction which depicted the differential behaviour of genotype over the time. For green fodder production, genotypes CAZRI 231 and CAZRI 2177 were found to be stable, whereas for higher dry matter production over wide environmental conditions genotypes CAZRI 231 and CAZRI 2177 were found stable (Rajora *et al.*, 2017).

Ramesh *et al.* (2017) investigated twenty genotypes of Pigeon pea (*Cajanus cajan* (L.) Mill sp.) for analysing stability for yield and its components during kharif season over three years. Among the varieties high significant differences were obtained for all the characters except pod bearing length, number of pods plant⁻¹ and seed yield. Except 100 seed weight, all other characters showed significant differences in different environment. For seed yield, the genotypes ICP-9691 and ICP-12654 were on par with check and the genotype ICP-13270 was stable for pod length across the environments.

For analysing the yield and micronutrient stability in Mung bean (*Vigna radiata* L.), thirty genotypes were raised across six artificial environments. Majority of the genotypes exhibited high significant difference in deviation from regression which depicted the suitability of these genotypes for better environment and their unpredictable response. Genotypes ML-1108, SMH-99-2,

MH-124, PDM-9-249 and ML-759 were recommended for better environment with high seed yield and iron content (Singh *et al.*, 2018).

Ten hybrids along with one check variety in Brinjal (*Solanum melongena* L.) was raised across four locations in Kerala for analyzing the stability and adaptability of yield and yield attributing characters using Eberhart and Russell model. Significant differences among genotypes, environments and genotype x environment interaction for all the characters were obtained from the pooled analysis of variance. Stability parameters like overall mean, regression coefficient (b_i) and deviation from regression (S^2_{di}) were used to identify the promising hybrids. The significance in mean squares due to Environments + (Genotype x Environment) revealed the existence of genotype x environment interaction. In *kharif* season, the stable hybrid identified was Wardha local x Palakurthi local (SMV1 x SMV2), widely adapted to all environments for days to first flowering, number of fruits plant⁻¹, fruit weight, fruit length, calyx length, yield plant⁻¹, yield plot⁻¹ and plant height. The hybrid with regression coefficient lower than unity and non-significant deviation from regression for days to first flowering, Surya x Vellayani local (SMV3 x SMV6) was found to be suited for poor environments. The hybrid NBR-38 x Vellayani local (SMV4 x SMV6) had high mean values, regression coefficient greater than unity and non-significant deviation from regression for days to first harvest and found stable for favourable environments (Vishwanath, 2018).

MATERIALS AND METHODS

3. MATERIALS AND METHODS

The present investigation on “Genotype×Environment interaction in Hedge lucerne (*Desmanthus virgatus* (L.) Willd.) for yield and quality” was conducted in the Department of Plant Breeding and Genetics, College of Agriculture, Vellayani, Thiruvananthapuram during 2016-2018. The field experiment was conducted at four locations of Kerala to determine the stability of eight genotypes of *Desmanthus virgatus* for yield and quality.

The details of materials used and methodologies adopted in the present study are described below.

3.1. EXPERIMENTAL SITE

The present work was carried out in four locations of Kerala.

Location I: College of Agriculture, Vellayani.

Location II: Krishi Vigyan Kendra, Kottarakkara.

Location III: College of Horticulture, Thrissur.

Location IV: Regional Agricultural Research Station, Ambalavayal.

3.2. EXPERIMENTAL DESIGN

The experiment was laid out in Randomized Block Design (RBD) with eight treatments and four replications. The spacing of 50 cm x continuous line sowing with a plot size of 3 m x 1 m was followed in the field experiment.

3.3. CULTURAL OPERATIONS

The land was prepared thoroughly by digging and leveling. The seeds collected were treated with hot water for 3 minutes followed by water soaking over night, shade dried and sown in the field in continuous line with 50 cm spacing between the lines.

3.4. RECORDING OF OBSERVATIONS

The list of hedge lucerne genotypes used as treatments in the study are described in Table 1. From each replication five competitive plants per treatment were randomly selected and tagged. From these plants observations with respect to different characters were recorded and the mean values of five plants were considered for statistical analysis. Observations were recorded on the following characters during 90 (first cut), 140 (second cut), 190 (third cut) and 240 (fourth cut) DAS (Days After Sowing).

3.4.1. Growth Characters

3.4.1.1. *Plant Height (cm)*

The height of the plant was measured from the base of the plant to the tip of the tallest branch at the time of each harvest using measuring scale and their mean values was expressed in centimeter.

3.4.1.2. *Number of Branches Plant⁻¹*

The total number of branches in a plant was counted at each harvest.

3.4.1.3. *Length of Branches (cm)*

The length of the branches was measured from the main stem to the tip of the branch at the time of each harvest using measuring scale and their mean value was expressed in centimeter.

3.4.1.4. *Number of Leaves Plant⁻¹*

The total number of leaves produced in a plant was counted at each harvest.

3.4.1.5. *Leaf to Stem Ratio*

The sample plants were cut at the base. The leaves and the stem were separated and oven dried for 5 days till constant weight was obtained. The dry weight of leaves and stem of individual plants were recorded. The ratio was computed by dividing leaf dry weight by the stem dry weight.

Table 1. List of hedge lucerne (*Desmanthus virgatus* L. (Willd)) genotypes used in the study

Sl. No.	Genotypes	Genotype name	Source
1.	T ₁	IC 345276	IGFRI, Jhansi
2.	T ₂	IC 343710	IGFRI, Jhansi
3.	T ₃	IC 89910	IGFRI, Jhansi
4.	T ₄	IC 261839	IGFRI, Jhansi
5.	T ₅	IC 90934	IGFRI, Jhansi
6.	T ₆	IC 421199	IGFRI, Jhansi
7.	T ₇	TNDV 1	Tamil Nadu Agricultural University
8.	T ₈	Thumburmuzhi local	Cattle Breeding Farm, Thumburmuzhi

Plate 1. General field view

a) COA, Vellayani



b) KVK, Kottarakkara



Plate 2. General field view

c) COH, Thrissur



d) RARS, Ambalavayal



3.4.1.6. Number of Pods Cluster⁻¹

The number of pods in each cluster was counted in a plant at harvest and mean values were measured.

3.4.2. Yield Attributes

3.4.2.1. Green Fodder Yield (g plant⁻¹)

At regular cutting intervals the crop was harvested and fresh weight of the plants in the net plot was recorded and expressed in g plant⁻¹.

3.4.2.2. Dry Fodder Yield (g plant⁻¹)

From each harvest crop samples were collected and dried in a hot air oven at 70°C to a constant weight and expressed in g plant⁻¹.

3.4.3. Quality Aspects

3.4.3.1. Crude Protein Content (%)

The crude protein content was calculated by multiplying the nitrogen content of the plant by the factor 6.25 (Simpson *et al.*, 1965) and expressed in percentage (%).

3.4.3.2. Crude Fibre Content (%)

The crude fibre content was determined by AOAC method (AOAC, 1975) and expressed in percentage (%).

3.4.4. Physiological Characters

3.4.4.1. Dry Matter Production (g plant⁻¹)

During the seed maturation stage five observational plants from each replication were uprooted. Shoot, leaves and roots were separated and dried to a constant weight at 70°C in a hot air oven. The sum of these individual components gave total dry matter production.

3.4.4.2. Leaf Area Index

Leaf area of observational plants was measured using Leaf area meter LI-COR 3100 available at Department of Crop Physiology, Tamil Nadu Agricultural University, Coimbatore. The leaf area index was obtained from the formula according to Watson (1952).

$$\text{LAI} = \frac{\text{Leaf area of the plant (cm}^2\text{)}}{\text{Ground area occupied (cm}^2\text{)}}$$

3.4.4.3. Crop Growth Rate ($\text{g m}^{-2} \text{ day}^{-1}$)

CGR was computed using the formula of Watson (1958) and expressed as $\text{g m}^{-2} \text{ day}^{-1}$.

$$\text{CGR} = \frac{W_2 - W_1}{P(t_2 - t_1)}$$

Where,

W_1 and W_2 = Plant dry weights at times t_1 and t_2

t_1 and t_2 = time interval in days

P = ground area on which W_1 and W_2 have been estimated.

3.4.4.4. Net Assimilation Rate ($\text{g dm}^{-2} \text{ day}^{-1}$)

Net Assimilation Rate (NAR) refers to the change in dry weight of the plant per unit leaf area per unit time. The procedure given by Gregory (1917) and modified by Williams (1946) was followed for calculating NAR.

$$\text{NAR} = \frac{w_2 - w_1}{t_2 - t_1} \times \frac{\log_e L_2 - \log_e L_1}{L_2 - L_1}$$

Where, W_1 - dry weight of plant (g) at time t_1 , L_1 - leaf area (dm^2) at t_1

W_2 - dry weight of plant (g) at time t_2 , L_2 - leaf area (dm^2) at t_2

$t_2 - t_1$:- time interval in days

3.5 STATISTICAL ANALYSIS

The data recorded on different traits were subjected to the following statistical analysis.

1. Analysis of Variance
2. Stability Analysis

3.5.1 Analysis of Variance

3.5.1.1 Analysis in Randomized Block Design (RBD)

The Randomized Block Design (RBD) was adopted with four replications. As per the method outlined by Panse and Sukhatme (1985) the analysis of variance was carried out.

$$Y_{ij} = m + g_i + r_j + e_{ij}$$

Where, Y_{ij} = Phenotypic observation of i^{th} genotype in j^{th} replication

M = General mean

g_i = True effect of i^{th} genotype

r_j = True effect of j^{th} replication

e_{ij} = Random error associated with i^{th} genotype and j^{th} replication

For each character Analysis of Variance (ANOVA) was carried out as indicated below:

Source of variation	d.f.	SS	MSS	F-ratio
Replications	$r-1$	RSS	M_r	M_r/M_e
Genotypes	$g-1$	TSS	M_g	M_g/M_e
Error	$(r-1)(g-1)$	ESS	M_e	
Total	$rg-1$			

Where,

r = Number of replications

t = Number of treatments (genotypes)

M_r = Mean sum of squares of replications

M_g = Mean sum of squares of treatments

M_e = Mean sum of squares of error

d.f = Degrees of freedom

The significance of mean sum of squares for each character was tested against the corresponding error degrees of freedom using 'F' test (Fisher and Yates, 1967).

$$\text{Standard Error Mean (SE(m))} = (M_e/r)^{1/2}$$

Where,

M_e = Error mean of squares

r = Number of replications

C.D = S.E (d) x t

Where,

$$\text{S.E (d)} = (2 M_e/r)^{1/2}$$

't' = t Table value at error degrees of freedom

$$\text{C.V} = (\text{S.D}/\bar{X}) \times 100$$

Where, S.D = Standard deviation of the population

\bar{X} = Population mean

3.5.2 Stability Analysis

3.5.2.1 Methods to Measure Stable Performance of Genotypes

Analysis of variance of genotypic mean was computed for each agronomic variable in each environment. The data were pooled over environments as the coefficient of variation values in each environment was generally low.

3.5.2.2 Eberhart and Russell's model (1966)

To study the stability of genotypes under different environments the methodology of Eberhart and Russell's model is used. The parameters estimated are the following (i) overall mean genotype over a range of environments, (ii) the regression of each genotype on the environmental index and (iii) a function of squared deviation from the regression.

$$Y_{ij} = m + B_i I_j + \delta_{ij} \quad (i = 1, 2, \dots, g \text{ and } j = 1, 2, \dots, e)$$

Where, Y_{ij} = mean of i^{th} genotype in j^{th} environment

m = mean of all genotypes over all the environments

B_i = regression coefficient of the i^{th} genotype on the environmental index which measures the response of this genotype to varying environments

I_j = environmental index which is defined as the deviation of the mean of all the genotypes at a given location from overall mean

$$I_j = \frac{\sum_i Y_{ij}}{t} - \frac{\sum_i \sum_j Y_{ij}}{ge}$$

$$\text{With } \sum_j I_j = 0$$

δ_{ij} = The deviation from regression of the i^{th} genotype at j^{th} environment

3.5.2.3 Analysis of Variance for Stability

The analysis of variance proposed by Eberhart and Russell (1966) is given below.

ANOVA to estimate stability parameters (Eberhart and Russell, 1966)

Source	d.f	Sum of squares	Mean sum of squares
Total	(ge-1)	$\sum \sum Y_{ij}^2 - CF$	
Genotype	(g-1)	$\frac{\sum Y_i^2}{e} - CF$	MS ₁
Environment + (Genotype x Environment)	g(e-1)	$\sum \sum Y_{ij}^2 - \frac{\sum Y_i^2}{e}$	
Environment Linear	1	$\frac{\sum (Y_{ij} I_j)^2}{g(\sum I_j^2)}$	
Genotype + Environment (Linear)	(g-1)	$\sum \left[\frac{(\sum_j Y_{ij} I_j)^2}{\sum_j I_j^2} \right] - \frac{(\sum_j Y_j I_j)^2}{g \sum_j I_j^2}$	MS ₂
Pooled deviation	g(e-2)	$\sum \sum \delta_{ij}^2$	MS ₃
Deviation due to Genotype 1 Genotype 2 : Genotype g	(e-2)	$\left[\sum Y_{ij}^2 - \frac{Y_i^2}{e} \right] - \left[\frac{(\sum Y_{ij} I_j)^2}{\sum I_j^2} \right]$	
Pooled error	ge(r-1)		S _e ²

g = No. of genotypes = 8, r = No. of replications = 4

e = No. of environments = 4

3.5.2.4 Estimation of stability parameters

The two stability parameters, regression coefficient (b_i) and deviation from regression (S_{di}^2) were estimated as follows :

3.5.2.5 Computation of regression coefficient (b_i) for each genotype

$$b_i = \frac{\sum_j Y_{ij} I_j}{\sum_j I_j^2}$$

Where,

b_i = regression coefficient of i^{th} genotype

$\sum_j I_j^2$ = The sum of squares of environmental indices (I_j) which are common to each value of b_i

$\sum_j Y_{ij} I_j$ = (for each genotype) = The sum of products of environmental index (I_j) and the corresponding means of that genotypes t for each environment (Y_{ij}).

3.5.2.6 Computation of Mean Square Deviation S_{di}^2 from Linear Regression

In regression analysis, the variance of dependent variable (Y) is partitioned into two parts, the one which explains the linearity between dependent and independent variables (variance due to regression) and the other which explains the variance due to deviations from linearity symbolically.

$$\sigma^2 Y = \sigma^2(\text{regression}) + \sigma^2 (\text{deviation from regression})$$

For estimating S_{di}^2 values, the variance due to deviation from regression can be obtained by subtracting the variance due to regression from $\sigma^2 Y$. The variance of means of individual genotypes over different locations can be obtained by,

$$\sigma_i^2 = \sum_j Y_{ij}^2 - \left(\frac{Y_i^2}{g}\right)$$

Where, Y_{ij} and Y_i are the mean values of genotypes in each location and total value of a variety in all the locations respectively.

The variance due to deviations from regression $(\sum_j \delta_{ij}^2)$ for a genotype being:

$$\sum_j \delta_{ij}^2 = \left[\sum_j Y_{ij}^2 - \frac{Y_i^2}{g} \right] - \frac{(\sum_j Y_{ij} l_j)^2}{\sum_j l_j^2}$$

Where,

$$\left[\sum_j Y_{ij}^2 - \frac{Y_i^2}{g} \right] = \text{The variance due to dependent variable}$$

$$\frac{(\sum_j Y_{ij} l_j)^2}{\sum_j l_j^2} = \text{The variance due to regression}$$

From which it can be obtained as

$$S^2 d_i = \left[\frac{\sum_j \delta_{ij}^2}{e-2} \right] - \left(\frac{S_e^2}{r} \right)$$

3.5.2.7 Test of Significance

The mean sum of squares due to genotypes and environments were tested against pooled deviation. Whereas, mean sum of squares due to G x E interaction was tested against pooled error. Environment (linear) and G x E (linear) were tested against pooled deviation. If pooled deviation is non-significant both these linear components were tested against pooled error. Mean sum of squares due to pooled deviations were tested against pooled error.

The following tests of significance were carried out:

1. To test the significance of difference among genotypes means i.e.,

$$H_0 = \mu_1 = \mu_2 = \mu_3 \dots = \mu_g$$

$$F = \frac{MS_1}{MS_3}$$

2. To test that the genotypes did not differ for their regression on environmental index, i.e., $H_0 = b_1 = b_2 = b_3 \dots = B_e$

$$F = \frac{MS_2}{MS_3}$$

3. Individual deviation from linear regression was tested as follows:

$$F = [(\sum_j \delta_{ij}^2)/(e - 2)] / \text{pooled error}$$

Against F table value at (e-2), ge (r-1), at 5% or 1% probability level.

3.5.2.8 Stable Genotype

A stable genotype with unit response was the genotype with unit regression coefficient ($b_i = 1$) and deviation not significantly different from zero ($S_{di}^2 = 0$).

Mean and standard error of 'b'

$$\text{Mean of } b = \bar{b} = \sum_i \frac{b_i}{v}$$

$$\text{S.E. (b)} = \sqrt{\frac{\text{Mean Sum of square due to pooled deviation}}{\sum_j I_j^2}}$$

$$\text{S.E. } b_i = \sqrt{\sum_j \delta_{ij}^2 / (e - 2) / \sum_j I_j^2}$$

3.5.2.9 Population mean

Population mean (μ) and standard error was calculated as

Population mean (μ) = Grand total/Number of observations

$$\text{S.E. (mean)} = \sqrt{\frac{\text{Mean sum of square due to pooled deviation}}{\text{Number of environments} - 1}}$$

RESULTS

4. RESULTS

The present study was conducted to evaluate the performance of eight hedge lucerne genotypes (*Desmanthus virgatus* (L.) Willd.) over four locations viz., College of Agriculture, Vellayani, College of Horticulture, Thrissur, Krishi Vigyan Kendra, Kottarakkara and Regional Agricultural Research Station, Ambalavayal, Wayanad in Kerala with an objective to identify stable genotypes of *Desmanthus virgatus* in varied environments with respect to yield and quality. The results obtained from the study are presented below under the following titles.

1. Analysis of variance
2. Mean performance
3. Stability parameters (Eberhart and Russell, 1966)

4.1 ANALYSIS OF VARIANCE

The analysis of variance showed significant differences among the eight hedge lucerne genotypes for all the characters studied across four environments (Table 2.1 – 2.4).

4.2 MEAN PERFORMANCE

4.2.1 Mean performance of hedge lucerne at College of Agriculture (COA), Vellayani

The mean performances of eight hedge lucerne genotypes for different parameters of growth, yield, quality and physiological characters at COA, Vellayani were recorded and presented below.

4.2.1.1. Growth characters

The mean performance of growth characters of hedge lucerne for different cuttings at COA, Vellayani are given in the Table 3.

Table 2. Analysis of variance (mean square) for individual locations

Table 2.1. Location-I (COA, Vellayani)

Source of variation	df	Plant height	Number of branches plant ⁻¹	Length of branches plant ⁻¹	Number of leaves plant ⁻¹	Leaf to stem ratio	Number of pods cluster ⁻¹	Green fodder yield	Dry fodder yield	Dry matter production	Crude protein	Crude fibre
Replication	3	3.48	0.21	2.73	3.16	0.0035	0.0067	37.45	1.51	0.26	3.07	2.36
Genotypes	7	973.25**	26.21**	1226.21**	1138.22**	0.032**	3.68**	340.78**	48.42**	78.79**	84.53**	28.79**
Error	21	1.81	0.59	0.18	6.29	0.0028	0.84	8.66	0.33	0.07	0.26	1.01

Table 2.2. Location-II (KVK, Kottarakkara)

Source of variation	df	Plant height	Number of branches plant ⁻¹	Length of branches plant ⁻¹	Number of leaves plant ⁻¹	Leaf to stem ratio	Number of pods cluster ⁻¹	Green fodder yield	Dry fodder yield	Dry matter production	Crude protein	Crude fibre
Replication	3	1.16	1.48	2.55	14.80	0.0094	0.0056	21.81	0.07	0.03	0.34	2.99
Genotypes	7	1075.01**	29.20**	1184.41**	1274.32**	0.034**	3.07**	259.97**	46.32**	120.02**	76.98**	34.25**
Error	21	1.11	0.45	0.17	19.99	0.0011	0.0071	19.17	0.09	0.15	0.26	0.57

** Significant at 1%, * Significant at 5%

Table 2.3. Location-III (COH, Thrissur)

Source of variation	df	Plant height	Number of branches plant ⁻¹	Length of branches plant ⁻¹	Number of leaves plant ⁻¹	Leaf to stem ratio	Number of pods cluster ⁻¹	Green fodder yield	Dry fodder yield	Dry matter production	Crude protein	Crude fibre
Replication	3	0.87	0.95	0.28	7.27	0.0027	0.035	8.52	0.13	0.27	0.24	0.16
Genotypes	7	1054.30**	31.30**	1155.50**	1502.46**	0.044**	3.04**	398.24**	54.11**	93.07**	81.96**	26.65**
Error	21	1.03	0.21	0.20	12.39	0.0016	0.046	9.27	0.12	0.15	0.13	0.11

Table 2.4. Location-IV (RARS, Ambalavayal)

Source of variation	df	Plant height	Number of branches plant ⁻¹	Length of branches plant ⁻¹	Number of leaves plant ⁻¹	Leaf to stem ratio	Number of pods cluster ⁻¹	Green fodder yield	Dry fodder yield	Dry matter production	Crude protein	Crude fibre
Replication	3	1.22	0.43	0.30	3.74	0.0016	0.02	2.41	0.09	1.68	0.47	0.15
Genotypes	7	893.12**	22.08**	1242.09**	1026.62**	0.031**	2.89**	308.83**	60.83**	84.41**	90.13**	27.71**
Error	21	1.08	0.38	0.43	16.56	0.0016	0.01	8.72	0.14	0.45	0.35	0.33

** Significant at 1%, * Significant at 5%

Table. 3 Mean performance of biometric characters of different cuttings of hedge lucerne at COA, Vellayani

Genotypes	Plant Height (cm)				Number of Branches Plant ⁻¹				Length of Branches (cm)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	104.3	110.9	113.7	115.9	3.758	4.158	4.955	5.705	15.38	19.64	20.39	23.66
T ₂	75.42	79.46	82.69	89.12	5.190	5.710	6.230	7.258	26.66	30.94	35.97	38.07
T ₃	60.35	66.34	70.95	75.31	8.640	9.232	10.94	12.26	22.94	25.91	28.64	32.80
T ₄	89.29	90.84	95.39	98.16	3.680	4.120	4.900	5.503	44.21	46.52	50.29	54.26
T ₅	91.23	96.00	100.9	102.8	5.490	6.039	6.940	7.273	53.33	56.62	60.75	63.82
T ₆	101.9	105.9	110.9	112.2	4.120	4.743	4.950	5.395	36.28	40.92	42.87	48.34
T ₇	77.59	80.79	83.75	85.71	3.990	4.150	4.830	5.465	24.91	27.34	30.85	33.38
T ₈	102.9	111.1	114.3	118.7	9.100	9.610	10.94	11.63	64.55	67.62	70.94	75.32
S.E (m)	1.038	0.725	0.937	0.671	0.043	0.070	0.070	0.134	0.304	0.381	0.479	0.217
C.D (0.05)	3.072	2.146	2.775	1.988	0.127	0.206	0.206	0.280	0.900	1.128	1.419	0.642

Genotypes	Number of Leaves Plant ⁻¹				Leaf to Stem Ratio				Number of Pods Cluster ⁻¹			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	112.3	124.6	139.5	146.2	0.910	0.914	0.921	0.948	0.000	1.208	1.230	1.758
T ₂	86.31	95.64	100.9	113.1	0.650	0.690	0.719	0.765	1.250	1.354	1.550	1.640
T ₃	75.61	89.61	98.62	107.9	0.810	0.820	0.880	0.898	1.053	1.183	1.430	1.538
T ₄	72.51	85.62	106.3	128.2	0.670	0.700	0.730	0.748	1.853	2.190	2.640	3.083
T ₅	88.59	102.9	132.9	136.5	0.630	0.650	0.690	0.685	1.560	1.970	2.148	2.335
T ₆	106.3	125.9	139.6	142.7	0.850	0.880	0.900	0.890	0.000	0.000	0.000	0.000
T ₇	84.21	91.08	101.6	111.0	0.740	0.790	0.800	0.885	2.150	2.338	2.590	2.718
T ₈	99.99	114.9	134.6	146.1	0.780	0.830	0.850	0.823	1.540	1.678	1.958	2.515
S.E (m)	1.062	1.107	1.085	2.031	0.007	0.007	0.009	0.038	0.012	0.017	0.017	0.058
C.D (0.05)	3.144	3.277	3.213	4.225	0.021	0.020	0.028	0.078	0.036	0.050	0.050	0.172

4.2.1.1.1. *Plant height (cm)*

The plant height showed significant difference in all the genotypes. The genotype T₁ recorded the maximum (104.3cm) plant height, which was on par with T₈ (102.9 cm) and T₆ (101.9 cm) during first cut of hedge lucerne. The plant height of genotypes increased in successive cuttings with the highest plant height recorded for the genotype T₈ during second, third and fourth cuttings (111.1 cm, 114.3 cm and 118.3 cm). The genotype T₈ was on par with the genotype T₁ for plant height at second (110.9 cm) and third (113.7 cm) cuttings of hedge lucerne. The genotype T₃ showed the minimum plant height at four successive cuttings (60.35 cm, 66.34 cm, 70.95 cm and 75.31 cm).

4.2.1.1.2 *Number of branches plant⁻¹*

The highest number of branches plant⁻¹ was observed in genotype T₈ for the successive three cuttings (9.100, 9.610 and 10.94) and the genotype T₃ showed maximum number of branches plant⁻¹ during third (10.94) and fourth (12.26) cut of hedge lucerne. The genotype T₄ recorded the minimum number of branches plant⁻¹ during first (3.684) and second cut (4.120) of hedge lucerne. The genotype T₇ showed the lowest number of branches plant⁻¹ during third cut and the genotype T₆ during fourth cut (5.395).

4.2.1.1.3 *Length of branches (cm)*

The highest length of branches was observed for the genotype T₈ during first (64.55 cm), second (67.62 cm), third (70.92 cm) and fourth (75.32 cm) cuttings. The genotype T₃ showed the minimum length of branches during all the four cuttings (15.38 cm, 19.64 cm, 20.39 cm and 23.66 cm) of hedge lucerne.

4.2.1.1.4 *Number of leaves plant⁻¹*

The highest number of leaves plant⁻¹ was observed for the genotype T₁ during first (112.3) and fourth (146.2) cut of hedge lucerne. During second (125.9) and third (139.6) cutting, the genotype T₆ showed the maximum number of leaves plant⁻¹. The genotype T₁ was on par with T₆ during second (124.6) and

third (139.5) cut and T₈ (146.1) and T₆ (142.7) were on par with the genotype T₁ during fourth cut of hedge lucerne in number of leaves plant⁻¹. The lowest number of leaves plant⁻¹ was observed for the genotype T₄ during first (72.51) and second (85.62) cut and the genotype T₃ (75.61) was on par with T₄ during first cut. The minimum number of leaves plant⁻¹ was observed for the genotype T₃ during third (98.62) and fourth (107.9) cut and this was on par with T₇ (101.6) and T₂ (100.9) during third cut and with T₇ (111.0) during fourth cut of hedge lucerne.

4.2.1.1.5 Leaf to stem ratio

The genotype T₁ recorded highest leaf to stem ratio (0.910, 0.914, 0.924 and 0.948) for all the cuttings. The genotype T₆ (0.900) was on par with the genotype T₁ during third cut and T₃ (0.898), T₆ (0.890) and T₇ (0.885) were on par with T₁ during fourth cut. The minimum leaf to stem ratio was observed for the genotype T₅ (0.630, 0.650, 0.690 and 0.685) during all the four cuttings. The genotype T₂ (0.650) was on par with the genotype T₅ during first cut.

4.2.1.1.6 Number of pods cluster⁻¹

The genotype T₇ showed the maximum number of pods cluster⁻¹ during first (2.150) and second (2.338) cut whereas, the genotype T₄ recorded the highest number of pods cluster⁻¹ during third (2.640) and fourth (3.083) cut. The genotype T₇ (2.590) was on par with T₄ at the third cut for the character number of pods cluster⁻¹.

4.2.1.2 Yield attributes

The mean performances of yield attributes of hedge lucerne for different cuttings at COA, Vellayani are given in the Table 4.

4.2.1.2.1 Green fodder yield (g plant⁻¹)

Green fodder yield showed significant variation among the genotypes for different cut in hedge lucerne. Superior yield was reported in the genotype T₆ during first (96.39 g) and second (100.69 g) cut. The genotype T₁ was on par with

Table. 4 Mean performance of yield attributes of different cuttings of hedge lucerne at COA, Vellayani

Genotypes	Green Fodder Yield (g)				Dry Fodder Yield (g)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	91.230	98.610	108.94	116.55	25.390	27.638	30.278	34.313
T ₂	80.970	84.608	98.640	103.09	18.625	21.368	24.190	28.603
T ₃	76.340	81.290	90.283	96.785	18.553	20.818	23.080	26.005
T ₄	90.283	95.288	100.94	105.30	23.613	24.320	31.293	32.805
T ₅	84.290	90.840	94.860	93.928	21.503	23.220	25.908	27.183
T ₆	96.390	100.69	108.89	112.03	24.680	28.880	30.288	31.940
T ₇	85.673	91.062	100.79	109.09	26.383	27.340	28.940	30.363
T ₈	92.190	97.060	109.75	119.58	27.768	30.820	33.520	35.905
S.E (m)	0.936	0.931	0.849	1.971	0.198	0.299	0.307	0.285
C.D (0.05)	2.773	2.757	2.513	4.100	0.587	0.885	0.908	0.844

T₆ (98.61 g) for green fodder yield at the second cut. During the third cut, the genotype T₈ recorded the maximum (109.75 g) yield which was on par with T₆ (108.89 g) and T₁ (108.94 g). The highest green fodder yield was recorded for the genotype T₈ during fourth cut which was on par with T₁ (116.55 g). The lowest yield was recorded for the genotype T₃ during first (76.34 g), second (81.29 g) and third (90.283 g) cut whereas, the minimum yield during fourth cut was observed for the genotype T₅ (93.928 g) which was on par with T₃ (96.785 g).

4.2.1.2.2 Dry fodder yield (g plant⁻¹)

The highest dry fodder yield was observed for the genotype T₈ at all the four cuttings (27.768 g, 30.820 g, 33.520 g and 35.905 g). The lowest dry fodder yield was observed for the genotype T₂ (18.625 g) during first cut which was on par with T₃ (18.553 g). The genotype T₃ showed the minimum dry fodder yield during second (20.818 g), third (23.080 g) and fourth (26.005 g) cut. The genotype T₂ was on par with the genotype T₃ (21.368 g) during the second cut of harvest.

4.2.1.3 Quality aspects

The mean performances of quality aspects of hedge lucerne for different cuttings at COA, Vellayani are given in Table 5.

4.2.1.3.1 Crude protein content (%)

Crude protein content was the highest for the genotype T₈ for all the four cuttings i.e., 25.00%, 25.02%, 25.29% and 25.46%. The genotype T₇ was on par with the genotype T₈ (24.58%) during first cut. The lowest crude protein content was observed for the genotype T₄ during first, second, third and fourth cut with 13.95%, 14.00%, 14.02% and 14.16% respectively.

4.2.1.3.2 Crude fibre content (%)

The crude fibre content was the highest for the genotype T₈ for all the four cuttings in hedge lucerne (30.24%, 30.25%, 30.75% and 30.99%) while the

Table. 5 Mean performance of quality aspects of different cuttings of hedge lucerne at COA, Vellayani

Genotypes	Crude Protein Content (%)				Crude Fibre Content (%)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	22.44	22.48	22.85	23.61	23.12	24.13	24.50	24.52
T ₂	16.01	16.03	16.31	16.93	24.15	24.25	24.59	24.64
T ₃	16.02	16.08	16.14	16.16	28.09	28.30	28.31	28.49
T ₄	13.95	14.00	14.02	14.16	24.03	24.11	24.19	24.25
T ₅	20.43	20.48	14.56	14.99	29.00	29.10	29.22	29.55
T ₆	23.49	23.81	21.59	21.66	23.19	24.99	25.03	25.06
T ₇	24.58	24.98	24.11	24.44	28.12	28.23	28.59	28.64
T ₈	25.00	25.02	25.29	25.46	30.24	30.25	30.75	30.99
S.E (m)	0.202	0.205	0.178	0.258	0.232	0.278	0.293	0.504
C.D (0.05)	0.598	0.607	0.526	0.765	0.687	0.824	0.868	1.492

Fig.1 Mean performance of yield attributes of hedge lucerne at COA, Vellayani

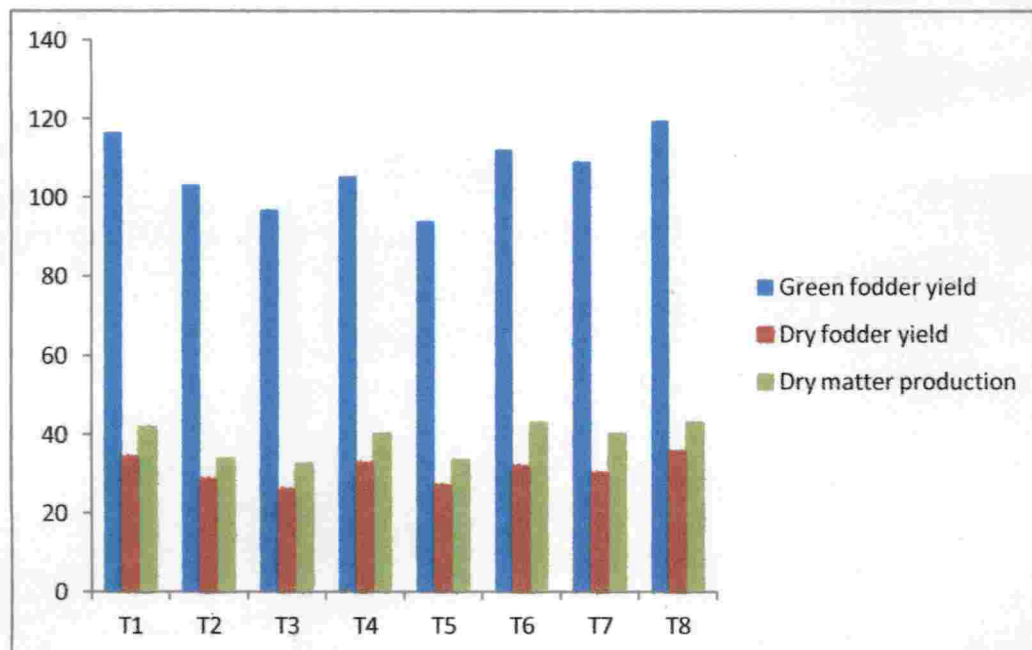
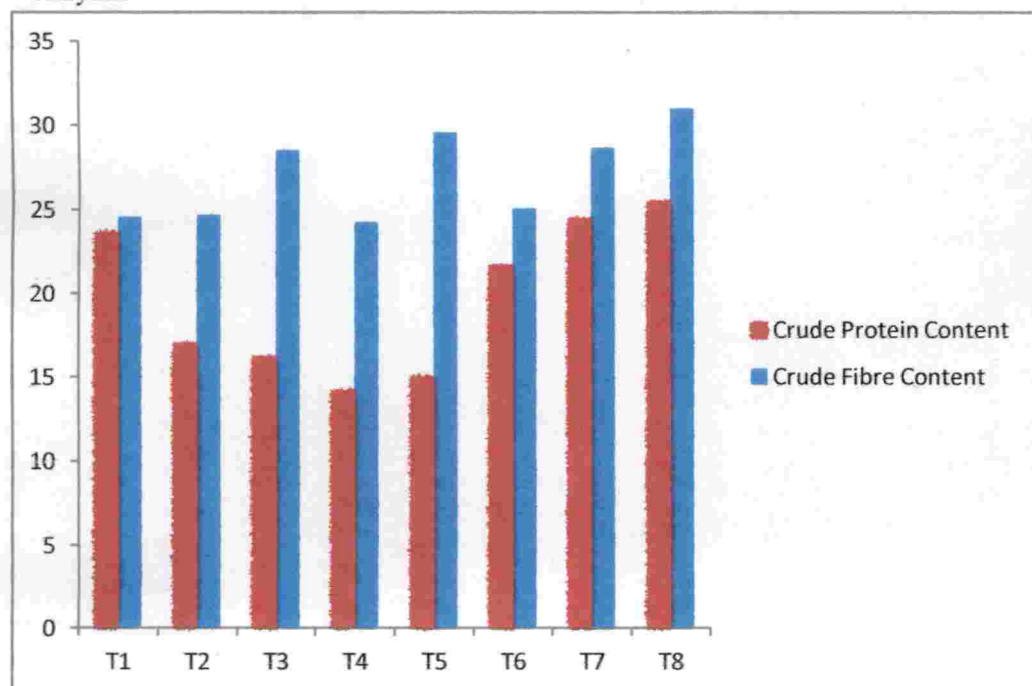


Fig 2. Mean performance of quality characters of hedge lucerne at COA, Vellayani



genotype T₅ (29.55%) was on par with the genotype T₈ for the crude fibre content during fourth of cut. During the first cut, the lowest crude protein content was observed for the genotype T₁ (23.12%) which was on par with the genotype T₆ (23.19%). The lowest crude protein content was observed for the genotype T₄ during second (24.11%), third (24.19%) and fourth (24.25%) stage of cuttings which was on par with T₅ (24.99%, 25.03% and 25.06%), T₁ (24.13%, 24.50% and 24.52%) and T₂ (24.25%, 24.59% and 24.64%) during second, third and fourth cuttings.

4.2.1.4 Physiological characters

The mean performances of physiological characters of hedge lucerne for different cuttings at COA, Vellayani are given in Table 6.

4.2.1.4.1 Dry matter production (g plant⁻¹)

T₈ genotype recorded highest dry matter production (35.64 g, 38.69 g, 41.29 g) during the first, second and third cuttings of hedge lucerne. The genotype T₆ (34.82 g) and T₇ (34.62 g) was on par with the genotype T₈ in the first cutting and the genotype T₆ (37.88 g) was on par with the genotype T₈ in the second cutting. During the third and fourth cutting, the maximum dry matter production (41.29 g and 43.08 g) was recorded from T₆ genotype. The lowest dry matter production was recorded from the genotype T₅ (25.64 g), T₂ (28.55 g), T₂ (30.89 g) and T₃ (32.65 g) during first, second, third and fourth cutting respectively. The genotype T₃ (31.19 g) was on par with the genotype T₂ during the third cutting.

4.2.1.4.2 Leaf Area Index

The leaf area index showed significant variation in each genotype. The highest leaf area index was observed for the genotype T₁ (7.418, 8.028, 8.058 and 8.168) and the lowest for the genotype T₃ (2.740, 3.410, 3.430 and 3.453) in all the four cuts of hedge lucerne.

Table. 6 Mean performances of physiological characters of hedge lucerne at COA, Vellayani

Genotypes	Dry Matter Production(g)				Leaf Area Index				Crop Growth Rate (g m ⁻² day ⁻¹)				Net Assimilation Rate (g dm ⁻² day ⁻¹)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	33.59	34.29	36.29	41.96	7.418	8.028	8.058	8.168	2.97	1.50	1.76	2.69	0.02	-0.32	-0.01	0.01
T ₂	26.35	28.55	30.89	34.02	3.308	4.110	4.120	4.440	1.65	1.83	1.88	1.59	0.02	-0.63	0.05	-0.01
T ₃	28.45	30.15	31.19	32.65	2.740	3.410	3.430	3.453	0.80	1.51	1.51	1.95	0.01	-0.92	-0.05	0.03
T ₄	33.91	35.49	38.56	40.22	5.033	5.960	6.003	6.020	0.58	0.47	1.80	1.01	0.02	0.57	0.04	0.05
T ₅	25.64	28.61	32.09	33.67	6.230	5.920	5.950	5.958	1.58	1.15	1.79	0.85	0.02	-0.63	-0.05	0.08
T ₆	34.82	37.88	41.29	43.08	6.938	7.050	7.123	7.390	1.44	2.03	0.94	1.10	0.03	-0.54	0.03	0.02
T ₇	34.62	35.49	38.69	40.48	4.110	4.328	4.353	4.478	1.54	0.64	1.07	0.95	0.02	-0.80	0.37	0.02
T ₈	35.64	38.69	41.29	43.04	5.810	6.113	6.310	6.405	3.67	2.80	4.65	2.94	0.03	0.63	1.21	0.08
S.E (m)	0.366	0.268	0.363	0.133	0.059	0.089	0.067	0.236	0.341	0.234	0.325	0.292	0.002	0.037	0.019	0.023
C.D (0.05)	1.085	0.793	1.074	0.394	0.125	0.187	0.142	0.490	0.973	0.732	0.936	0.837	0.007	0.109	0.057	NS

4.2.1.4.3 Crop Growth Rate ($\text{g cm}^{-2} \text{ day}^{-1}$)

Crop growth rate was highest in the genotype T₈ ($3.67 \text{ g m}^{-2} \text{ day}^{-1}$) in first, second, third and fourth cutting. The genotype T₁ was on par ($2.97 \text{ g m}^{-2} \text{ day}^{-1}$ and $2.69 \text{ g m}^{-2} \text{ day}^{-1}$) with the genotype T₈ in the first and fourth cutting. The lowest crop growth rate was recorded from the genotype T₄ ($0.58 \text{ g m}^{-2} \text{ day}^{-1}$) in the first and second cutting followed by the genotypes T₆ ($0.94 \text{ g m}^{-2} \text{ day}^{-1}$) and T₅ ($0.85 \text{ g m}^{-2} \text{ day}^{-1}$) in the third and fourth cutting respectively. The genotypes T₃ ($0.80 \text{ g m}^{-2} \text{ day}^{-1}$), T₆ ($1.44 \text{ g m}^{-2} \text{ day}^{-1}$) and T₇ ($1.54 \text{ g m}^{-2} \text{ day}^{-1}$) were on par with the genotype T₄ in the first cutting. During the third cutting the genotypes T₁ ($1.76 \text{ g m}^{-2} \text{ day}^{-1}$), T₃ ($1.51 \text{ g m}^{-2} \text{ day}^{-1}$), T₄ ($1.80 \text{ g m}^{-2} \text{ day}^{-1}$), T₅ ($1.79 \text{ g m}^{-2} \text{ day}^{-1}$) and T₇ ($1.07 \text{ g m}^{-2} \text{ day}^{-1}$) was on par with the genotype T₆. The genotypes T₂ ($1.59 \text{ g m}^{-2} \text{ day}^{-1}$), T₄ ($1.01 \text{ g m}^{-2} \text{ day}^{-1}$), T₆ ($1.10 \text{ g m}^{-2} \text{ day}^{-1}$) and T₇ ($0.95 \text{ g m}^{-2} \text{ day}^{-1}$) were on par with the genotype T₅ during the fourth cutting.

4.2.1.4.4 Net Assimilation Rate ($\text{g dm}^{-2} \text{ day}^{-1}$)

The maximum net assimilation rate was recorded in the genotype T₈ ($0.03 \text{ g dm}^{-2} \text{ day}^{-1}$, $0.63 \text{ g dm}^{-2} \text{ day}^{-1}$ and $1.21 \text{ g dm}^{-2} \text{ day}^{-1}$) for first, second and third cutting. The genotype T₄ ($0.57 \text{ g dm}^{-2} \text{ day}^{-1}$) was on par with the genotype T₈. The lowest net assimilation rate was recorded in the genotype T₃ ($0.01 \text{ g dm}^{-2} \text{ day}^{-1}$, $-0.92 \text{ g dm}^{-2} \text{ day}^{-1}$ and $-0.05 \text{ g dm}^{-2} \text{ day}^{-1}$) during first, second and third cutting.

4.2.2 Mean performance of hedge lucerne at KVK, Kottarakkara

The mean performances of eight hedge lucerne genotypes for different parameters of growth, yield, quality and physiological characters at KVK, Kottarakkara were studied.

4.2.2.1 Growth characters

The mean performances of growth characters of hedge lucerne genotypes for different cuttings at KVK, Kottarakkara are given in Table 7.

Table. 7 Mean performances of biometric characters of different cuttings of hedge lucerne at KVK, Kottarakkara

Genotypes	Plant Height (cm)				Number of Branches Plant ⁻¹				Length of Branches (cm)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	100.7	108.6	110.4	112.6	4.130	4.950	5.125	5.728	15.30	18.34	20.53	22.73
T ₂	74.12	80.94	82.01	86.04	4.868	5.110	5.943	6.203	22.41	25.96	27.64	32.62
T ₃	54.55	60.88	65.97	68.05	7.150	8.940	9.763	10.11	24.85	28.64	32.19	38.13
T ₄	82.96	86.92	90.38	92.00	2.193	2.840	3.213	3.675	43.67	48.21	50.94	55.75
T ₅	89.64	95.01	98.64	100.3	4.668	5.128	5.860	6.428	50.21	54.19	56.34	61.13
T ₆	97.45	103.9	107.1	109.9	2.990	3.213	3.870	4.138	34.92	39.61	43.85	47.40
T ₇	77.91	82.53	85.91	89.40	2.190	2.940	3.160	3.843	20.91	25.95	29.64	34.15
T ₈	104.4	110.8	115.9	117.5	8.610	9.430	10.94	11.26	65.29	68.66	70.28	74.70
S.E (m)	1.067	0.914	1.001	0.527	0.046	0.059	0.061	0.087	0.240	0.390	0.432	0.211
C.D (0.05)	3.160	2.706	2.964	1.559	0.135	0.174	0.181	0.178	0.710	1.153	1.280	0.626

Genotypes	Number of Leaves Plant ⁻¹				Leaf to Stem Ratio				Number of Pods Cluster ⁻¹			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	109.6	125.9	140.9	147.5	0.840	0.870	0.900	0.890	0.000	1.123	1.250	1.325
T ₂	95.21	112.6	135.2	143.9	0.620	0.630	0.650	0.680	1.200	1.233	1.290	1.485
T ₃	71.02	89.64	95.32	99.37	0.880	0.900	0.930	0.910	0.953	0.960	1.100	1.353
T ₄	75.61	91.20	100.6	124.7	0.850	0.870	0.710	0.770	1.140	1.230	2.490	2.883
T ₅	86.38	102.4	128.9	144.1	0.590	0.610	0.650	0.675	1.420	1.850	1.968	2.080
T ₆	95.64	119.6	125.9	143.3	0.820	0.830	0.840	0.843	0.000	0.000	0.000	0.000
T ₇	81.84	90.26	96.34	114.8	0.750	0.760	0.770	0.790	1.518	1.960	2.150	2.333
T ₈	96.39	105.7	132.1	144.9	0.750	0.790	0.801	0.868	1.250	1.450	1.890	2.183
S.E (m)	1.090	1.044	1.186	3.162	0.007	0.006	0.009	0.005	0.010	0.013	0.017	0.042
C.D (0.05)	3.226	3.092	3.512	6.577	0.022	0.018	0.019	0.022	0.031	0.039	0.052	0.125

4.2.2.1.1 Plant height (cm)

The maximum plant height for hedge lucerne was recorded from genotype T₈ (104.4 cm, 110.8 cm, 115.9 cm and 117.5 cm) and the lowest plant height was recorded from the genotype T₃ during all the four cuttings. The genotype T₁ (108.6 cm) was on par with T₈ during the second cutting.

4.2.2.1.2 Number of branches plant⁻¹

The highest number of branches plant⁻¹ was obtained in the genotype T₈ (8.61, 9.43, 10.94 and 11.26) in all the four cuttings. The lowest number of branches was observed in the genotype T₇ (2.19 and 3.16) during first and third cutting. The genotype T₄ (2.19 and 3.12) was on par with T₇ for the first and third cutting. The genotype T₄ showed the lowest number of branches (2.84 and 3.68) during second and fourth cutting. The genotype T₇ was on par with T₄ (2.94 and 3.84) for the second and fourth cutting.

4.2.2.1.3 Length of branches (cm)

For length of branches, the highest and lowest value was observed in the genotype T₈ (65.29 cm, 68.66 cm, 70.28 cm and 74.7 cm) and the genotype T₁ (15.3 cm, 18.34 cm, 20.53 cm and 22.73 cm) respectively for the four cuttings in hedge lucerne.

4.2.2.1.4 Number of leaves plant⁻¹

The number of leaves was highest in the genotype T₁ (109.6, 125.9, 140.9 and 147.5) in four cuttings. During the fourth cutting genotypes T₂, T₅, T₆ and T₈ were on par with T₁. Lowest number of leaves was recorded in the genotype T₃ for the four cuttings which was on par with T₇ (90.26 and 96.34) during second and fourth cutting and also with T₄ during the second cut.

4.2.2.1.5 Leaf to stem ratio

Among the genotypes evaluated, highest (0.88, 0.99, 0.93 and 0.91) and lowest (0.59, 0.61, 0.65 and 0.67) leaf to stem ratio was observed in the genotype

T₃ and T₅ respectively for all cuttings. T₁ and T₆ were on par with T₃ and T₂ was on par with T₅ during the fourth cutting.

4.2.2.1.6 Number of pods cluster⁻¹

The maximum number of pods per cluster was obtained from the genotype T₇ (1.52 and 1.96) for the first and second cutting and the genotype T₄ (2.49 and 2.88) during the third and fourth cutting. The genotype T₆ did not produce any pods at the time of four cuttings.

4.2.2.2 Yield attributes

The mean performance of yield attributes of hedge lucerne genotypes for different cuttings at KVK, Kottarakkara are given in Table 8.

4.2.2.2.1 Green fodder yield (g plant⁻¹)

The green fodder yield plant⁻¹ varied from 70.58 g to 88.89 g in the genotype T₃. The highest green fodder yield was obtained for the genotype T₈ (112.99 g) during the fourth cutting and 110.89 g, 102.97 g and 98.61 g for the third, second and first cutting respectively.

4.2.2.2.2 Dry fodder yield (g plant⁻¹)

For the four cuttings, the highest dry fodder yield was obtained from the genotype T₈ (28.51 g, 30.95 g, 32.09 g and 33.13 g) and lowest from the genotype T₃ (18.62 g, 19.63 g, 20.95 g and 22.85 g).

4.2.2.3 Quality aspects

The mean performances of quality aspects of hedge lucerne genotypes for different cuttings at KVK, Kottarakkara are given in Table 9.

4.2.2.3.1 Crude protein content (%)

Among the genotypes, the highest crude protein content was recorded from the genotype T₈ (24.20 %, 24.21 %, 24.35 % and 24.46 %) and the genotype T₄

Table. 8 Mean performance of yield attributes of different cuttings of hedge lucerne at KVK, Kottarakkara

Genotypes	Green Fodder Yield (g)				Dry Fodder Yield (g)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	90.280	93.288	102.94	110.69	26.350	28.560	30.818	31.973
T ₂	83.238	85.908	89.640	101.99	21.583	23.153	23.608	27.250
T ₃	70.588	74.120	80.290	88.893	18.620	19.630	20.950	22.850
T ₄	88.670	90.280	95.670	97.420	25.690	27.123	28.640	31.113
T ₅	82.910	87.617	92.310	103.91	20.638	23.283	24.923	28.058
T ₆	96.578	99.990	105.82	109.93	26.330	28.508	30.620	32.230
T ₇	97.380	100.87	106.38	108.37	23.283	25.153	28.510	30.163
T ₈	98.610	102.97	110.89	112.99	28.508	30.950	32.090	33.130
S.E (m)	0.710	0.845	0.718	3.095	0.254	0.262	0.223	0.148
C.D (0.05)	2.103	2.502	2.125	6.439	0.752	0.777	0.660	0.440

Table. 9 Mean performance of quality aspects of different cuttings of hedge lucerne at KVK, Kottarakkara

Genotypes	Crude Protein Content (%)				Crude Fibre Content (%)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	21.00	21.01	21.05	21.17	22.14	22.54	22.67	22.71
T ₂	15.62	15.96	16.03	16.02	22.00	22.09	22.15	22.25
T ₃	14.20	14.22	14.28	14.54	26.31	26.45	26.52	26.66
T ₄	12.94	13.01	13.06	13.14	23.00	23.12	23.23	23.44
T ₅	13.00	14.02	14.09	14.15	28.09	28.35	28.41	28.52
T ₆	13.99	19.55	19.62	19.93	23.42	23.51	23.58	24.03
T ₇	19.52	22.42	22.95	23.06	27.51	27.52	27.64	27.64
T ₈	24.20	24.21	24.35	24.46	29.45	29.46	29.58	29.58
S.E (m)	0.134	0.173	0.174	0.260	0.209	0.306	0.254	0.256
C.D (0.05)	0.397	0.513	0.515	0.769	0.620	0.905	0.753	0.755

Fig 3. Mean performance of yield attributes of hedge lucerne at KVK, Kottarakkara

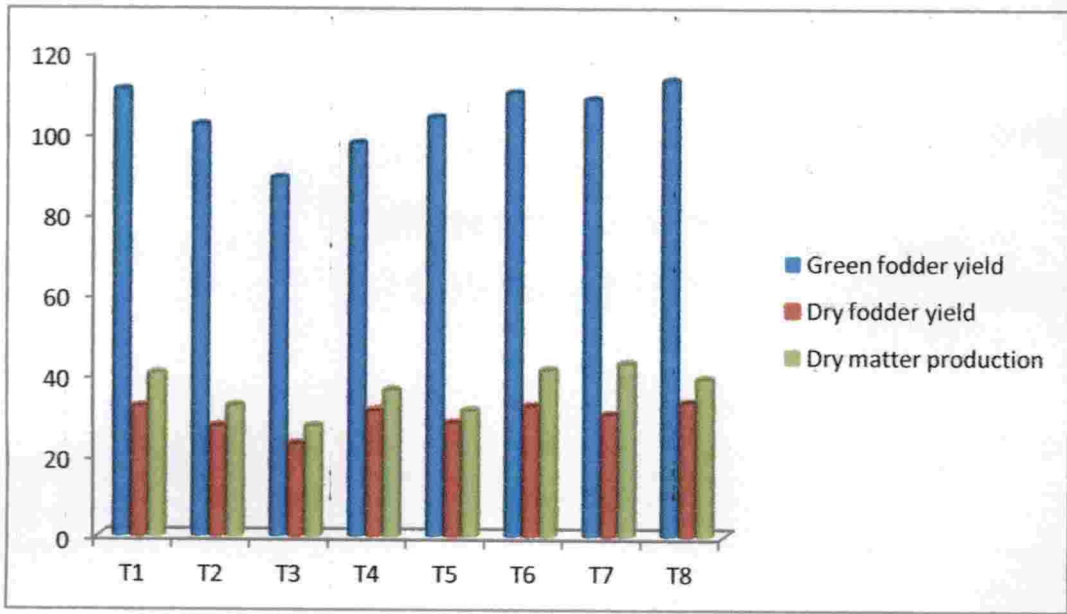
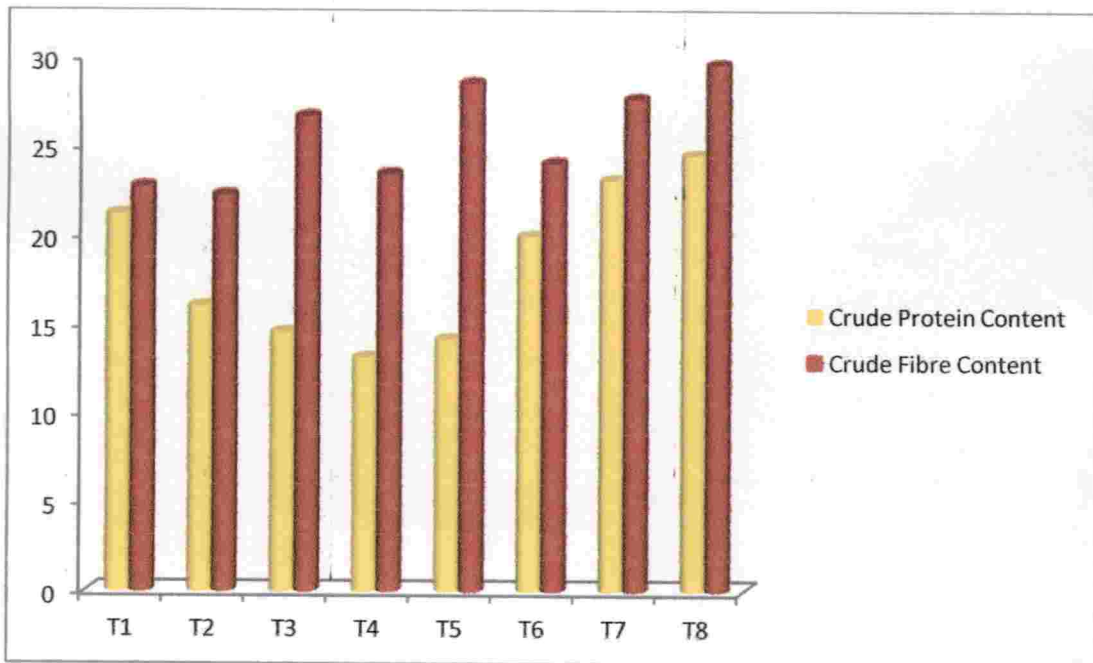


Fig 4. Mean performance of quality characters of hedge lucerne at KVK, Kottarakkara



recorded the lowest crude protein content (12.94 %, 13.01%, 13.06 % and 13.14 %) during the four cuttings. The genotype T₅ was on par with T₄ during the first cutting (13 %).

4.2.2.3.2 Crude fibre content (%)

The highest crude fibre content was obtained from the genotype T₈ (29.45 %, 29.46 %, 29.58% and 29.58%) and lowest crude fibre content was recorded in the genotype T₂ (22%, 22.09%, 22.15% and 22.25%) which was on par with the genotype T₁ (22.14%, 22.54%, 22.67% and 22.71%) during all the four cuttings.

4.2.2.4 Physiological characters

The mean performances of physiological characters of hedge lucerne genotypes for different cuttings at KVK, Kottarakkara are given in Table 10

4.2.2.4.1 Dry matter production ($g\ plant^{-1}$)

The genotype T₇ recorded the highest dry matter production $plant^{-1}$ in four cuttings (34.29 g, 37.24 g, 40.27 g and 42.69 g). The lowest dry matter production was obtained from the genotype T₃ (21.02 g, 23.95 g, 25.94 g and 27.25 g) in all the four cuttings of hedge lucerne.

4.2.2.4.2 Leaf Area Index

Leaf area index measured the highest values (7.42, 7.54, 7.61 and 7.71) in the genotype T₁ and the lowest values (2.74, 2.75, 2.81 and 2.89) in the genotype T₃ for all four harvest.

4.2.2.4.3 Crop Growth Rate ($g\ m^{-2}\ day^{-1}$)

Crop growth rate in the genotype T₈(6.07 $g\ m^{-2}\ day^{-1}$), T₅(1.76 $g\ m^{-2}\ day^{-1}$), T₈ (2.24 $g\ m^{-2}\ day^{-1}$ and 2.43 $g\ m^{-2}\ day^{-1}$) recorded the highest among the genotypes first, second, third and fourth cut respectively. The genotypes T₁ (1.47 $g\ m^{-2}\ day^{-1}$, T₆(1.45 $g\ m^{-2}\ day^{-1}$), T₇ (1.25 $g\ m^{-2}\ day^{-1}$) and T₈ (1.63 $g\ m^{-2}\ day^{-1}$)

Table.10. Mean performances of physiological characters of hedge lucerne at KVK, Kottarakkara

Genotypes	Dry Matter Production (g)				Leaf Area Index				Crop Growth Rate (g m ⁻² day ⁻¹)				Net Assimilation Rate (g dm ⁻² day ⁻¹)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	34.26	36.28	39.21	40.19	7.418	7.540	7.618	7.715	2.90	1.47	1.51	0.77	0.22	0.024	0.011	0.407
T ₂	24.52	26.34	30.64	32.29	3.308	3.440	3.540	3.995	3.44	1.05	0.88	1.27	0.21	0.031	0.012	-0.172
T ₃	21.02	23.95	25.94	27.25	2.740	2.753	2.813	2.895	2.71	0.67	0.31	0.69	0.26	0.017	0.010	-0.743
T ₄	24.39	28.63	30.65	36.12	5.033	5.120	5.240	5.470	3.77	0.95	1.01	1.65	0.22	0.017	0.015	-5.651
T ₅	23.95	27.91	29.34	31.17	6.230	6.418	6.510	6.650	1.90	1.76	1.09	2.09	0.17	0.018	0.011	-1.918
T ₆	33.92	35.65	38.66	41.12	6.938	7.013	7.090	7.098	4.15	1.45	1.41	1.07	0.18	0.035	0.018	0.313
T ₇	34.29	37.24	40.27	42.69	4.110	4.120	4.220	4.230	3.12	1.25	0.76	1.10	0.24	0.027	0.016	-1.044
T ₈	28.64	31.92	34.59	38.87	5.810	5.940	6.000	6.020	6.07	1.63	2.24	2.43	0.31	0.039	0.019	1.512
S.E (m)	0.247	0.382	0.300	0.191	0.059	0.077	0.092	0.195	0.614	0.225	0.182	0.214	0.029	0.002	0.001	0.246
C.D (0.05)	0.731	1.131	0.889	0.566	0.125	0.159	0.191	0.404	1.762	0.643	0.524	0.582	NS	0.005	0.002	0.728

was on par with the genotype T₅ during second cut. The genotype T₅ (2.09 g m⁻² day⁻¹) was on par with the genotype T₈ during the fourth cutting. The lowest crop growth rate was observed in the genotype T₅ (1.9 g m⁻² day⁻¹) in the first cutting which was on par with the genotypes T₁ (2.9 g m⁻² day⁻¹), T₃ (2.71 g m⁻² day⁻¹) and T₇ (3.12 g m⁻² day⁻¹). During the second cutting, the genotype T₃ (0.67 g m⁻² day⁻¹) recorded the lowest crop growth rate and was on par with the genotypes T₂ (1.05 g m⁻² day⁻¹) and T₄ (0.95 g m⁻² day⁻¹). In the third cutting, the genotype T₃ (0.31 g m⁻² day⁻¹) was recorded lowest CGR and was on par with the genotype T₇ (0.76 g m⁻² day⁻¹). The genotype T₃ (0.69 g m⁻² day⁻¹) recorded the lowest crop growth rate which was on par with the genotypes T₁ (0.77 g m⁻² day⁻¹), T₆ (1.07 g m⁻² day⁻¹) and T₇ (1.1 g m⁻² day⁻¹).

4.2.2.4.4 Net Assimilation Rate (g dm⁻² day⁻¹)

The highest net assimilation rate was observed in the genotype T₈ (0.039 g dm⁻² day⁻¹, 0.019 g dm⁻² day⁻¹ and 1.512 g dm⁻² day⁻¹) during the second third and fourth cutting which was on par with the genotypes T₆ (0.035 g dm⁻² day⁻¹ and 2.9 g dm⁻² day⁻¹) in the second and third cutting. The lowest net assimilation rate was recorded from the genotypes T₃ (0.017 g dm⁻² day⁻¹ and 0.010 g dm⁻² day⁻¹) and T₄ (-5.651 g m⁻² day⁻¹) during the second, third and fourth cuttings. The genotypes T₅ (0.018 g dm⁻² day⁻¹ and 0.011 g m⁻² day⁻¹) and T₁ (0.011 g m⁻² day⁻¹) was on par with the genotype T₃ in the second and third cutting.

4.2.3 Mean performance of hedge lucerne at College of Horticulture (COH), Thrissur

The mean performances of eight hedge lucerne genotypes for different parameters of growth, yield, quality and physiological characters at COH, Thrissur were studied.

4.2.3.1 Growth characters

The mean performances of growth characters of hedge lucerne genotypes for different cuttings at COH, Thrissur are presented in Table 11.

4.2.3.1.1 Plant height (cm)

The genotype T₈ showed the maximum plant height during all the four cuttings (99.46 cm, 104.8 cm, 110.7 cm and 114.1 cm). The minimum plant height was observed for the genotypes T₃ for the first (55.12 cm), second (59.16 cm), third (63.59 cm) and fourth (66.13 cm) cuttings.

4.2.3.1.2 Number of branches plant⁻¹

The number of branches plant⁻¹ showed significant difference in all the eight genotypes of hedge lucerne. The maximum branches were observed for the genotype T₈ at all the four cuttings (7.910, 8.610, 9.468 and 10.38). The lowest branches were obtained for the genotype T₇ during first (1.878), second (2.104), third (2.760) and fourth (3.200) cut and it was on par with T₄ during first (1.943) and second (2.188) cut of hedge lucerne.

4.2.3.1.3 Length of branches (cm)

The length of branches varied significantly in all the genotypes. The longest branch was observed for the genotype T₈ (61.09 cm, 64.86 cm, 69.37 cm and 73.49 cm) while the genotype T₁ showed the shortest branch (14.26 cm, 15.62 cm, 19.64 cm and 22.44 cm) for all the four cuttings.

4.2.3.1.4 Number of leaves plant⁻¹

For the number of leaves plant⁻¹, the genotype T₆ (108.9) showed maximum number of leaves at first cut, T₁ (115.2 and 125.6) showed more number of leaves during second and third cut, while T₅ (143.3) recorded maximum number of leaves during fourth cut of hedge lucerne. The genotype T₆ was on par with T₁ during second (113.2) and third (123.9) cut, while T₈ (124.6) was on par with T₁ during third cut of hedge lucerne. The genotypes T₁ (142.2)

Table. 11 Mean performances of biometric characters of different cuttings of hedge lucerne at COH, Thrissur

Genotypes	Plant Height (cm)				Number of Branches Plant ⁻¹				Length of Branches (cm)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	93.29	100.8	104.1	108.4	3.090	3.810	4.123	4.620	14.26	15.62	19.64	22.44
T ₂	66.26	71.19	78.64	81.12	3.638	4.090	4.895	5.135	21.05	25.34	28.91	32.23
T ₃	55.12	59.16	63.59	66.13	7.208	7.943	8.753	9.675	23.22	26.84	31.29	36.46
T ₄	74.59	80.64	84.31	88.18	1.943	2.188	2.945	3.375	40.95	45.20	47.61	53.10
T ₅	87.22	91.62	95.82	98.24	4.120	4.818	5.178	5.663	46.38	51.61	55.94	60.87
T ₆	94.92	99.81	104.2	106.7	2.100	2.940	3.160	3.633	34.12	38.24	42.09	46.74
T ₇	70.66	76.23	80.99	84.92	1.878	2.140	2.760	3.200	21.85	25.64	28.68	32.94
T ₈	99.64	104.8	110.7	114.1	7.910	8.610	9.468	10.38	61.09	64.86	69.37	73.49
S.E (m)	0.840	0.961	0.851	0.507	0.030	0.057	0.045	0.081	0.349	0.480	0.521	0.225
C.D (0.05)	2.488	2.844	2.519	1.501	0.090	0.169	0.133	0.170	1.032	1.421	1.542	0.665

Genotypes	Number of Leaves Plant ⁻¹				Leaf to Stem Ratio				Number of Pods Cluster ⁻¹			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	98.98	115.6	125.6	142.2	0.810	0.850	0.860	0.868	0.000	0.890	1.100	1.200
T ₂	76.38	90.65	98.63	106.6	0.620	0.638	0.642	0.630	1.150	1.283	1.490	1.625
T ₃	69.56	75.12	82.39	91.70	0.840	0.872	0.915	0.930	0.980	1.010	1.090	1.180
T ₄	94.28	100.6	112.7	118.7	0.650	0.670	0.702	0.715	1.250	1.950	2.120	2.370
T ₅	84.95	95.64	109.6	143.3	0.570	0.600	0.630	0.653	1.150	1.490	1.670	1.963
T ₆	108.9	113.2	123.9	135.7	0.760	0.780	0.810	0.840	0.000	0.000	0.000	0.000
T ₇	74.91	86.31	91.19	110.3	0.680	0.710	0.730	0.810	1.890	2.010	2.130	2.553
T ₈	98.65	106.3	124.6	139.2	0.730	0.750	0.790	0.793	1.450	1.850	1.950	2.515
S.E (m)	0.945	0.959	0.998	2.489	0.008	0.007	0.007	0.032	0.013	0.009	0.015	0.153
C.D (0.05)	2.799	2.838	2.956	5.177	0.023	0.020	0.020	0.059	0.037	0.027	0.044	0.320



and T₈ (139.2) were on par with the genotype T₅ at fourth cut of hedge lucerne. The genotype T₃ recorded the minimum number of leaves during first (69.56), second (75.12), third (82.39) and fourth (91.70) cuts.

4.2.3.1.5 Leaf to stem ratio

During all the four cuts the genotype T₃ recorded the highest (0.840, 0.872, 0.915 and 0.930) leaf to stem ratio compared to other genotypes. The lowest leaf to stem ratio was observed for the genotype T₅ at first (0.570), second (0.600) and third (0.630) cut, whereas, T₂ showed the lowest (0.630) ratio during fourth cut. The genotype T₅ (0.653) was on par with T₂ during fourth cut.

4.2.3.1.6 Number of pods cluster⁻¹

The highest pods cluster⁻¹ was noticed in the genotype T₇ in all the four cuttings (1.890, 2.010, 2.130 and 2.553). The genotype T₄ was on par with T₇ during third (2.102) and fourth cut (2.370) while the genotype T₈ was on par with the genotype T₇ during fourth cut. No pods were noticed in the genotype T₆ during all the different stage of cuttings.

4.2.3.2. Yield attributes

The mean performances of yield attributes of hedge lucerne genotypes for different cuttings at COH, Thrissur are given in Table 12.

4.2.3.2.1 Green fodder yield (g plant⁻¹)

The variation in green fodder yield was noticed during each cut in hedge lucerne. During first cut, the highest yield was observed in case of the genotype T₁ (90.480 g) which was on par with the genotype T₈. For the rest three cuttings, the genotype T₈ showed the maximum green fodder yield of 96.367 g, 100.89 g and 110.54 g respectively, which was on par with the genotype T₁ (95.308 g, 100.58 g and 108.58 g). The lowest yield was performed by the genotype T₃ during all the four cuttings.

Table. 12 Mean performance of yield attributes of different cuttings of hedge lucerne at COH, Thrissur

Genotypes	Green Fodder Yield (g)				Dry Fodder Yield (g)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	90.840	95.308	100.58	108.76	22.343	24.250	26.323	29.365
T ₂	79.610	84.910	90.673	95.505	17.560	19.880	21.053	23.878
T ₃	69.398	75.123	79.420	81.748	17.420	19.120	19.338	19.620
T ₄	72.100	82.087	88.640	95.263	20.368	23.580	24.960	27.615
T ₅	81.690	89.980	94.940	102.04	19.658	22.150	24.393	26.530
T ₆	76.310	84.288	93.293	105.76	20.150	24.560	27.640	29.448
T ₇	75.280	81.073	88.642	110.43	18.250	20.330	22.940	27.608
T ₈	90.373	96.367	100.89	110.54	21.208	25.220	29.548	31.240
S.E (m)	0.808	0.936	0.731	2.153	0.208	0.257	0.212	0.174
C.D (0.05)	2.392	2.772	2.165	4.477	0.617	0.760	0.629	0.514

4.2.3.2.2 Dry fodder yield ($g\ plant^{-1}$)

The highest dry fodder yield was obtained for the genotype T₁ (22.43 g) during first cut of crop while, the genotype T₈ showed the maximum yield during remaining three cuttings. The lowest dry fodder yield was recorded for the genotype T₃ for all the four cuttings.

4.2.3.3 Quality aspects

The mean performances of quality aspects of hedge lucerne genotypes for different cuttings at COH, Thrissur are presented in Table 13.

4.2.3.3.1 Crude protein content (%)

The crude protein content was highest for the genotype T₈ during first (22.41%), second (22.43%), third (22.58%) and fourth (22.67%) cuttings. Whereas the lowest protein content was observed for the genotype T₄ in all the stages of cuttings (11.00%, 11.00%, 11.19% and 11.26%).

4.2.3.3.2 Crude fibre content (%)

During the four cuttings of hedge lucerne, the highest crude fibre content was observed for the genotype T₈ (27.31%, 27.34%, 27.65% and 28.37% respectively). The lowest crude fibre content was recorded for the genotype T₃ for all the four cuttings (20.42%, 20.45%, 20.69% and 20.76% respectively).

4.2.3.4 Physiological characters

The mean performances of physiological characters of hedge lucerne genotypes for different cuttings at COH, Thrissur was listed in Table 14.

4.2.3.4.1 Dry matter production ($g\ plant^{-1}$)

The genotype T₈ recorded maximum dry matter production (31.25 g, 33.62 g, 36.52 g and 38.65 g) during first, second, third and fourth cuttings. The lowest dry matter production was obtained from the genotype

Table. 13 Mean performances of quality aspects of different cuttings of hedge lucerne at COH, Thrissur

Genotypes	Crude Protein Content (%)				Crude Fibre Content (%)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	18.20	18.24	18.59	18.96	21.32	21.32	21.45	21.83
T ₂	13.42	13.54	13.68	13.78	20.42	20.45	20.69	20.76
T ₃	12.00	12.09	12.12	12.37	24.00	24.01	24.21	24.39
T ₄	11.00	11.00	11.19	11.26	22.51	22.63	22.82	22.86
T ₅	12.20	12.21	12.39	12.65	26.22	26.23	26.38	26.50
T ₆	18.12	18.14	18.65	18.78	22.00	22.09	22.13	22.36
T ₇	21.51	21.52	21.64	21.74	23.24	23.28	23.42	25.41
T ₈	22.41	22.43	22.58	22.67	27.31	27.34	27.65	28.37
S.E (m)	0.204	0.158	0.158	0.185	0.196	0.202	0.174	0.172
C.D (0.05)	0.604	0.467	0.469	0.549	0.581	0.597	0.516	0.510

Fig 5. Mean performance of yield attributes of hedge lucerne at COH, Thrissur

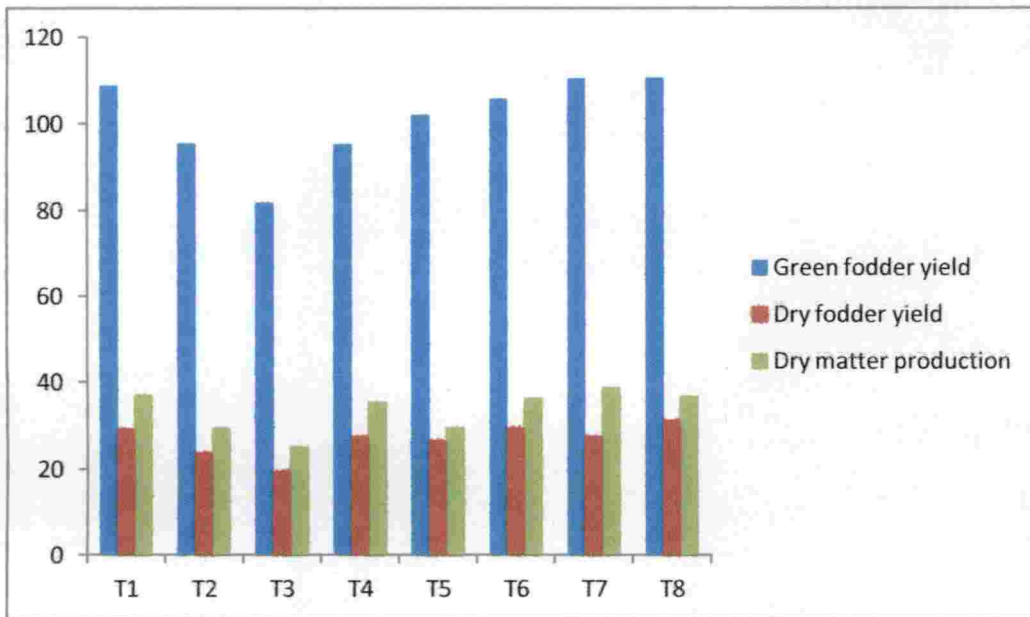


Fig 6. Mean performance of quality attributes of hedge lucerne at COH, Thrissur

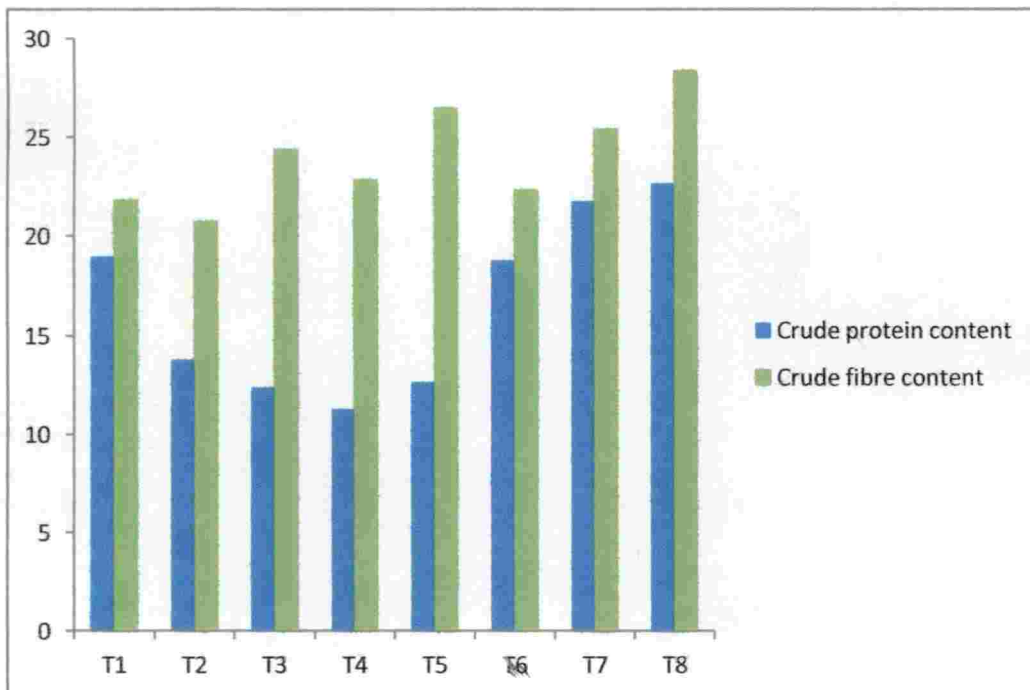


Table. 14 Mean performances of physiological characters of different cuttings of hedge lucerne at COH, Thrissur

Genotypes	Dry Matter Production (g)				Leaf Area Index				Crop Growth Rate (g m ⁻² day ⁻¹)				Net Assimilation Rate (g dm ⁻² day ⁻¹)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	28.62	31.29	34.59	36.91	6.418	6.530	6.620	6.950	3.32	1.27	1.38	2.03	0.056	0.020	0.303	0.088
T ₂	20.33	22.45	25.33	29.46	3.228	3.410	3.563	3.813	2.91	1.55	0.78	1.88	0.071	0.018	1.139	0.115
T ₃	18.61	20.96	23.01	25.05	2.013	2.123	2.750	2.858	3.60	1.13	0.15	0.19	0.056	0.022	0.061	0.793
T ₄	25.96	28.61	23.56	35.37	4.278	4.520	4.650	4.978	3.83	2.14	0.92	1.77	0.076	0.029	0.101	0.499
T ₅	20.37	21.09	25.28	29.59	5.010	5.160	5.760	5.835	2.84	1.66	1.41	1.43	0.057	0.060	0.562	0.196
T ₆	29.66	31.29	34.52	36.29	6.120	6.240	6.308	6.438	3.01	2.67	1.97	1.20	0.043	0.012	0.408	0.059
T ₇	31.25	33.62	36.52	38.65	4.558	4.820	4.860	4.973	3.54	1.39	1.59	1.13	0.039	0.028	0.714	0.819
T ₈	28.64	31.24	35.11	36.73	5.120	5.250	5.410	5.535	3.89	2.94	2.89	3.11	0.121	0.064	1.411	0.253
S.E (m)	0.254	0.230	0.264	0.193	0.059	0.067	0.081	0.230	0.393	0.256	0.215	0.223	0.003	0.004	0.090	0.060
C.D (0.05)	0.752	0.680	0.781	0.570	0.126	0.138	0.170	0.478	0.112	0.734	0.594	0.641	0.010	0.011	0.266	0.178

T₃ (31.25 g, 33.62 g, 36.52 g, 38.65g) during first, second, third and fourth cuttings.

4.2.3.4.2 Leaf Area Index

The genotypes showed significant variation for the character leaf area index. The genotype T₁ showed the maximum leaf area index at all the four cuttings (6.418, 6.530, 6.620 and 6.950 respectively). The genotype T₃ recorded the minimum leaf area index of 2.013, 2.123, 2.750 and 2.858 during first, second, third and fourth cut respectively.

4.2.3.4.3 Crop Growth Rate ($\text{g m}^{-2} \text{day}^{-1}$)

Among the hedge lucerne genotypes, the maximum crop growth rate was observed in the genotype T₈ (3.89 $\text{g m}^{-2} \text{day}^{-1}$, 2.94 $\text{g m}^{-2} \text{day}^{-1}$, 2.89 $\text{g m}^{-2} \text{day}^{-1}$ and 3.11 $\text{g m}^{-2} \text{day}^{-1}$) during the first, second, third and fourth cuttings. The genotypes T₄ (3.83 $\text{g m}^{-2} \text{day}^{-1}$) and T₆ (2.67 $\text{g m}^{-2} \text{day}^{-1}$) were on par with the genotype T₈ during first and second cuttings. The lowest crop growth rate was observed in the genotypes T₅ (2.84 $\text{g m}^{-2} \text{day}^{-1}$), T₃ (1.13 $\text{g m}^{-2} \text{day}^{-1}$, 0.15 $\text{g m}^{-2} \text{day}^{-1}$ and 0.19 $\text{g m}^{-2} \text{day}^{-1}$) during the first, second, third and fourth cuttings which was on par with T₂ (2.91 $\text{g m}^{-2} \text{day}^{-1}$) in the first cutting, T₁ (1.27 $\text{g m}^{-2} \text{day}^{-1}$), T₂ (1.55 $\text{g m}^{-2} \text{day}^{-1}$), T₅ (1.66 $\text{g m}^{-2} \text{day}^{-1}$) and T₇ (1.39 $\text{g m}^{-2} \text{day}^{-1}$) during the second cutting.

4.2.3.4.4 Net Assimilation Rate ($\text{g dm}^{-2} \text{day}^{-1}$)

The maximum net assimilation rate was recorded in the genotype T₈ (0.121 $\text{g dm}^{-2} \text{day}^{-1}$, 0.064 $\text{g dm}^{-2} \text{day}^{-1}$, 1.411 $\text{g dm}^{-2} \text{day}^{-1}$) and T₇ (0.819 $\text{g dm}^{-2} \text{day}^{-1}$) during the first, second, third and fourth cuttings. The genotypes T₅ (0.06 $\text{g dm}^{-2} \text{day}^{-1}$) and T₃ (0.793 $\text{g dm}^{-2} \text{day}^{-1}$) was on par with the genotypes T₈ and T₇ during the second and fourth cuttings respectively. The lowest net assimilation rate was observed in the genotypes T₇ (0.039 $\text{g dm}^{-2} \text{day}^{-1}$), T₆ (0.012 $\text{g dm}^{-2} \text{day}^{-1}$), T₃ (0.061 $\text{g dm}^{-2} \text{day}^{-1}$) and

T₆ (0.059 g dm⁻² day⁻¹) during the first, second, third and fourth cuttings respectively.

4.2.4 Mean performance of hedge lucerne at RARS, Ambalavayal

The mean performances of eight hedge lucerne genotypes for different parameters of growth, yield, quality and physiological characters at Ambalavayal were studied.

4.2.4.1 Growth characters

The mean performances of growth characters of hedge lucerne genotypes for different cuttings at RARS, Ambalavayal are presented in Table 15.

4.2.4.1.1 Plant height (cm)

The highest plant height was observed for the genotype T₈ during first (100.9 cm) and second (105.3 cm) cut, T₁ during third cut (111.0 cm) and T₈ at fourth cut in hedge lucerne. The genotype T₈ (110.9 cm) was on par with T₁ during third cut, whereas T₃ recorded the lowest plant height during all the cuttings (60.22 cm, 64.12 cm, 70.95 cm and 74.63 cm).

4.2.4.1.2 Number of branches plant⁻¹

During first and second cut, the number of branches was more for the genotype T₈ (8.070 and 8.638), but during third and fourth cut the genotype T₃ (9.845 and 11.35) reported the maximum branches plant⁻¹. The lowest number of branches was recorded for the genotype T₄ at first (2.933) and third (4.573) cut, whereas, during second and fourth cut the minimum branches were reported for the genotype T₇ (3.310 and 5.783).

4.2.4.1.3 Length of branches (cm)

The length of branches showed significant difference among the genotypes. The genotype T₈ reported the maximum length of branches during all the four cut (60.28 cm, 64.25 cm, 70.91 cm and 75.09 cm), while, the genotype T₁ reported

Table. 15 Mean performances of biometric characters of different cuttings of hedge lucerne at RARS, Ambalavayal

Genotypes	Plant Height (cm)				Number of Branches Plant ⁻¹				Length of Branches (cm)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	90.37	96.52	111.0	114.5	3.750	4.290	5.943	6.325	13.59	16.52	19.34	23.02
T ₂	76.33	81.09	85.67	90.96	5.190	5.890	6.315	7.973	24.39	28.68	31.28	36.24
T ₃	60.22	64.12	70.95	74.63	7.010	7.690	9.845	11.35	20.94	25.66	30.91	34.76
T ₄	87.54	90.85	94.66	99.77	2.933	3.580	4.573	5.855	43.28	46.38	51.82	54.87
T ₅	88.64	93.66	97.89	102.9	4.110	5.760	6.818	7.090	50.96	54.44	60.29	64.15
T ₆	90.22	94.28	99.09	112.3	3.610	4.670	5.183	5.833	29.64	34.92	41.09	48.19
T ₇	74.31	80.94	83.94	86.21	3.090	3.310	4.820	5.783	17.49	21.09	28.55	33.16
T ₈	100.9	105.3	110.9	116.7	8.070	8.638	9.410	10.61	60.28	64.25	70.91	75.09
S.E (m)	0.609	0.896	0.939	0.521	0.026	0.047	0.054	0.081	0.317	0.378	0.290	0.332
C.D (0.05)	1.804	2.653	2.779	1.544	0.086	0.139	0.161	0.166	0.938	1.120	0.859	0.982

Genotypes	Number of Leaves Plant ⁻¹				Leaf to Stem Ratio				Number of Pods Cluster ⁻¹			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	112.6	124.3	130.2	144.3	0.850	0.900	0.905	0.893	0.000	1.423	1.590	1.754
T ₂	72.61	86.34	98.64	115.0	0.690	0.710	0.730	0.780	1.030	1.120	1.218	1.558
T ₃	75.94	88.61	94.26	102.8	0.850	0.890	0.910	0.925	1.130	1.380	1.448	1.648
T ₄	94.61	104.7	112.9	127.9	0.730	0.750	0.780	0.810	1.850	2.090	2.190	2.733
T ₅	88.64	100.3	119.6	135.4	0.600	0.630	0.650	0.675	1.550	1.893	2.103	2.183
T ₆	106.5	125.6	139.6	142.5	0.840	0.880	0.900	0.943	0.000	0.000	0.000	0.000
T ₇	76.31	86.31	95.62	112.5	0.780	0.810	0.840	0.858	1.413	1.868	2.190	2.483
T ₈	101.3	114.6	134.2	143.0	0.750	0.790	0.810	0.858	1.948	2.000	2.010	2.310
S.E (m)	0.951	0.993	0.832	2.878	0.007	0.009	0.008	0.032	0.011	0.015	0.017	0.051
C.D (0.05)	2.815	2.939	2.464	5.987	0.020	0.028	0.023	0.058	0.032	0.044	0.049	0.150

the minimum length of branches at all the four cut (13.59 cm, 16.52 cm, 19.34 cm and 23.02 cm).

4.2.4.1.4 Number of leaves plant⁻¹

The number of leaves was highest for the genotype T₁ at first (112.6), second (124.3) and fourth (144.3) cut and during third cut the genotype T₆ (139.6) recorded the maximum number of leaves. The genotypes T₈ (143.0) and T₆ (142.5) was on par with T₁ during fourth cut of hedge lucerne. The lowest number of leaves was observed in the genotype T₂ (72.61) during first cut, T₇ (86.31) during second cut, and T₃ during third (94.26) and fourth cut (102.8).

4.2.4.1.5 Leaf to stem ratio

The leaf to stem ratio was highest for the genotype T₁ (0.850) and T₃ (0.850) during first cut and T₁ at second (0.900) cut, T₃ at third (0.910) and T₆ at fourth (0.943) cut. The lowest leaf to stem ratio was observed for the genotype T₅ for all the four cuttings in hedge lucerne (0.600, 0.630, 0.650 and 0.675).

4.2.4.1.6 Number of pods cluster⁻¹

The number of pods was highest in the genotype T₈ (1.948) of first cut, in the genotype T₄ (2.090, 2.190 and 2.733) of second, third and fourth cut respectively. No pods were observed in case of the genotype T₆ during all the four cuts and in T₁ during first cut.

4.2.4.2 Yield attributes

The mean performances of yield attributes of hedge lucerne genotypes for different cuttings at RARS, Ambalavayal are given in Table 16.

4.2.4.2.1 Green fodder yield (g plant⁻¹)

The green fodder yield was highest for the genotype T₈ at first (95.120 g), second (97.690 g), third (104.81 g) and fourth (118.56 g) cuts, T₁ and T₆ were on par during second (96.340g and 69.373 g) and third (102.85 g and 100.27 g) cut.

Table. 16 Mean performance of yield attributes of different cuttings of hedge lucerne at RARS, Ambalavayal

Genotypes	Green Fodder Yield (g)				Dry Fodder Yield (g)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	89.340	96.340	102.85	113.39	28.540	30.290	31.318	34.175
T ₂	81.270	90.270	96.380	104.53	21.008	24.810	28.583	30.133
T ₃	75.930	80.910	85.910	94.368	18.638	21.580	22.610	25.628
T ₄	89.373	95.668	98.620	104.70	24.290	29.993	30.540	33.415
T ₅	80.670	84.290	87.460	95.115	21.590	23.558	24.510	26.360
T ₆	89.640	96.373	100.27	112.18	24.950	28.540	30.940	33.483
T ₇	84.390	94.120	97.613	112.21	24.120	27.693	29.330	32.665
T ₈	95.120	97.690	104.81	118.56	30.890	31.850	34.223	36.753
S.E (m)	0.976	1.088	1.148	2.087	0.237	0.298	0.309	0.188
C.D (0.05)	2.891	3.222	3.400	4.341	0.700	0.882	0.914	0.556

The lowest green fodder yield was reported for the genotype T₃ at all the four cuttings (75.930 g, 80.910 g, 85.910 g and 94.368 g).

4.2.4.2.1 Dry fodder yield (g plant⁻¹)

The highest and lowest dry fodder yield was recorded for the genotype T₈ (30.890 g, 31.850 g, 34.223 g and 36.753 g) and T₃ (18.638 g, 21.580 g, 22.610 g and 25.628 g) during all the cuts in hedge lucerne.

4.2.4.3 Quality aspects

The mean performances of quality aspects of hedge lucerne genotypes for different cuttings at RARS, Ambalavayal are given in Table 17.

4.2.4.3.1 Crude protein content (%)

The highest crude protein content was reported for the genotype T₇ during first three cuttings (24.00%, 24.10% and 24.28% respectively) which was on par with the genotype T₈ (23.94%, 24.00% and 24.16% respectively). During fourth cut the genotype T₈ showed the maximum crude protein content of 24.98 %. The lowest crude protein content was reported for the genotype T₅ during all the cuts (13.00%, 13.49%, 13.15% and 13.88% accordingly) which was on par with the genotype T₄ (13.40%, 13.49%, 13.58% and 13.95% accordingly) during all the four cuttings in hedge lucerne.

4.2.4.3.2 Crude fibre content (%)

For all the four cuts, the genotype T₈ showed the maximum crude fibre content (30.00%, 30.02%, 30.19% and 31.52% respectively). The lowest crude fibre content was reported by the genotype T₂ for all the four cuts (23.00%, 23.00%, 23.01% and 23.23% respectively), which was on par with the genotype T₄ (23.45%, 23.54%, 23.69% and 23.79% respectively).

Table. 17 Mean performances of quality aspects of different cuttings of hedge lucerne at RARS, Ambalavayal

Genotypes	Crude Protein Content (%)				Crude Fibre Content (%)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	21.24	21.26	21.45	21.82	24.11	24.21	24.26	24.35
T ₂	15.00	15.00	15.06	15.31	23.00	23.00	23.01	23.23
T ₃	15.01	15.03	15.12	15.32	27.13	27.25	27.45	27.78
T ₄	13.40	13.49	13.58	13.95	23.45	23.54	23.69	23.79
T ₅	13.00	13.11	13.15	13.88	28.00	28.11	28.21	28.28
T ₆	20.24	20.41	20.46	20.69	24.11	24.12	24.32	24.79
T ₇	24.00	24.10	24.28	24.71	27.12	27.29	27.54	27.97
T ₈	23.94	24.00	24.16	24.98	30.00	30.02	30.19	31.52
S.E (m)	0.228	0.169	0.165	0.299	0.195	0.256	0.267	0.288
C.D (0.05)	0.674	0.501	0.489	0.885	0.577	0.758	0.791	0.854

Fig 7. Mean performance of yield characters of hedge lucerne at RARS, Ambalavayal

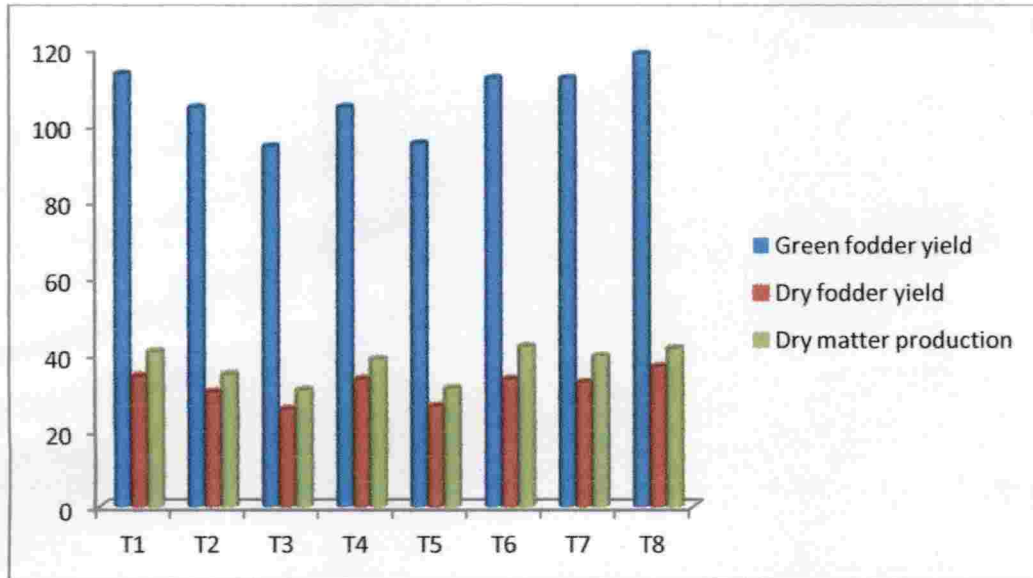
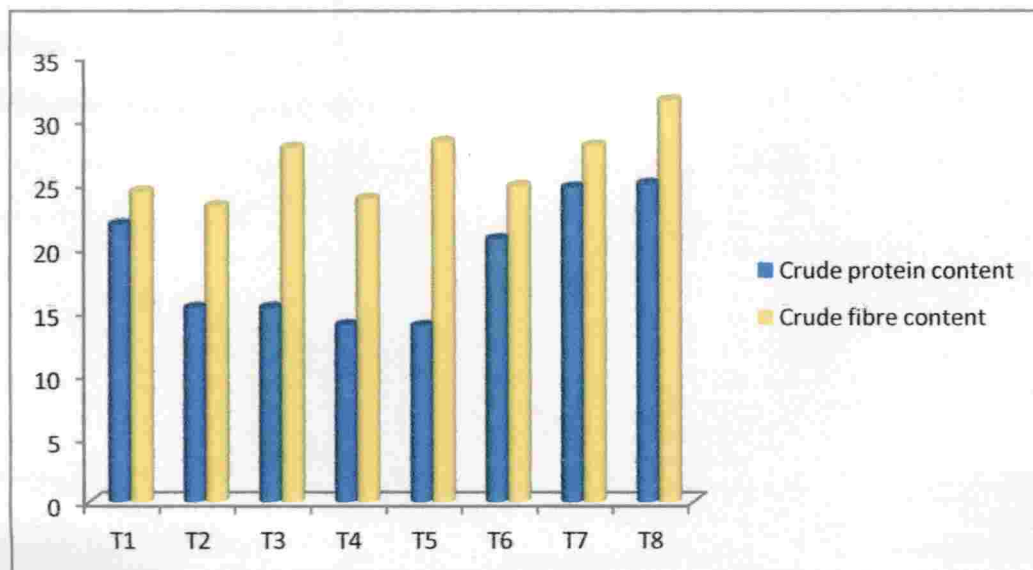


Fig 8. Mean performance of quality characters of hedge lucerne at RARS, Ambalavayal



4.3.4.4 Physiological characters

The mean performances of physiological characters of hedge lucerne genotypes for different cuttings at RARS, Ambalavayal are given in Table 18.

4.2.4.4.1 Dry matter production ($g\ plant^{-1}$)

Dry matter production recorded maximum in the genotypes T₆ (34.93 g), T₈ (37.26 g and 40.12 g) and T₆ (41.99 g). The genotype T₈ (41.53 g) was on par with the genotype T₆ during the fourth cutting. The lowest dry matter production was obtained in the genotypes T₅ (20.67 g), T₃ (24.92 g), T₅ (28.33 g) and T₃ (30.64 g).

4.2.4.4.2 Leaf Area Index

The leaf area index was highest for the genotype T₁ (7.250, 7.440, 7.540 and 7.660) during all the four cuts whereas, the genotype T₃ showed the minimum leaf area index for first (3.240), second (3.358), third (3.490) and fourth (3.585) cuts.

4.2.4.4.3 Crop Growth Rate ($g\ m^{-2}\ day^{-1}$)

The highest crop growth rate was obtained from the genotypes T₂ ($4.52\ g\ m^{-2}\ day^{-1}$), T₈ ($3.80\ g\ m^{-2}\ day^{-1}$, $2.51\ g\ m^{-2}\ day^{-1}$ and $2.22\ g\ m^{-2}\ day^{-1}$) during first, second, third and fourth cuttings respectively. The genotypes T₅ ($3.54\ g\ m^{-2}\ day^{-1}$), T₆ ($3.87\ g\ m^{-2}\ day^{-1}$) and T₈ ($3.96\ g\ m^{-2}\ day^{-1}$) were on par with the genotype T₂ during the first cutting. The genotype T₂ ($1.58\ g\ m^{-2}\ day^{-1}$) and the genotype T₆ ($1.6\ g\ m^{-2}\ day^{-1}$) were on par with the genotype T₈. During the fourth cut, the genotypes T₁ ($1.9\ g\ m^{-2}\ day^{-1}$), T₂ ($2.01\ g\ m^{-2}\ day^{-1}$), T₄ ($1.92\ g\ m^{-2}\ day^{-1}$), T₆ ($1.7\ g\ m^{-2}\ day^{-1}$) and T₇ ($1.69\ g\ m^{-2}\ day^{-1}$) were on par with the genotype T₈.

4.2.4.4.4 Net Assimilation Rate ($g\ dm^{-2}\ day^{-1}$)

The highest net assimilation rate was recorded in the genotype T₈ ($0.059\ g\ dm^{-2}\ day^{-1}$ and $0.021\ g\ dm^{-2}\ day^{-1}$) during first and second cutting

Table. 18 Mean performances of physiological characters of different cuttings of hedge lucerne at RARS Ambalavayal

Genotypes	Dry Matter Production (g)				Leaf Area Index				Crop Growth Rate (g m ⁻² day ⁻¹)				Net Assimilation Rate (g dm ⁻² day ⁻¹)			
	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut	1 st cut	2 nd cut	3 rd cut	4 th cut
T ₁	31.64	34.52	38.51	40.64	7.250	7.440	7.540	7.660	4.30	1.17	0.69	1.90	0.028	0.016	0.021	0.012
T ₂	28.55	30.56	32.15	34.69	4.163	4.260	4.308	4.468	4.52	2.48	1.58	2.01	0.045	0.014	0.067	0.011
T ₃	21.94	24.92	28.62	30.64	3.240	3.358	3.490	3.585	2.47	0.64	0.37	1.03	0.027	0.015	0.013	0.014
T ₄	28.54	32.06	34.62	38.64	5.978	6.098	6.130	6.273	3.19	1.96	0.69	1.92	0.026	0.011	0.027	0.011
T ₅	20.67	25.95	28.33	31.06	5.840	5.970	6.010	6.240	3.54	1.31	0.63	1.23	0.026	0.015	0.037	0.012
T ₆	34.93	36.36	38.64	41.99	7.060	7.120	7.280	7.488	3.87	2.39	1.60	1.70	0.024	0.011	0.034	0.010
T ₇	27.59	31.25	35.26	39.48	4.990	5.100	5.118	5.215	2.93	2.38	1.09	1.69	0.058	0.011	0.018	0.012
T ₈	34.22	37.26	40.12	41.53	6.480	6.520	6.820	6.908	3.96	3.80	2.51	2.22	0.059	0.021	0.031	0.013
S.E (m)	0.327	0.291	0.314	0.337	0.089	0.074	0.067	0.294	0.342	0.315	0.321	0.265	0.001	0.001	0.001	0.000
C.D (0.05)	0.969	0.862	0.929	0.998	0.184	0.155	0.138	0.611	0.982	0.894	0.933	0.758	0.002	0.004	0.003	0.001

whereas the genotypes T₂ (0.067 g dm⁻² day⁻¹) and T₈ (0.013 g dm⁻² day⁻¹) recorded highest during the third and fourth cutting respectively. The genotypes T₇ (0.059 g dm⁻² day⁻¹) during the first cutting and T₁ (0.012 g dm⁻² day⁻¹), T₅ (0.012 g dm⁻² day⁻¹) and T₇ (0.012 g dm⁻² day⁻¹) during the fourth cutting was on par with the genotype T₈. The lowest net assimilation rate was recorded in the genotypes T₆ (0.024 g dm⁻² day⁻¹) in the first cutting, followed by the genotypes T₄ (0.011 g dm⁻² day⁻¹), T₆ (0.011 g dm⁻² day⁻¹), T₇ (0.011 g dm⁻² day⁻¹) in the second cutting. The genotypes T₃ (0.013 g dm⁻² day⁻¹) and T₆ (0.010 g dm⁻² day⁻¹) during third and fourth cutting respectively. The genotypes T₄ (0.026 g dm⁻² day⁻¹) and T₅ (0.026 g dm⁻² day⁻¹) were on par with the genotype T₆ during the first cutting. The genotypes T₂ (0.014 g dm⁻² day⁻¹), T₃ (0.015 g dm⁻² day⁻¹) and T₅ (0.015 g dm⁻² day⁻¹) were on par with the genotype T₄ in the second cutting. The genotypes T₂ (0.011 g dm⁻² day⁻¹) and T₄ (0.011 g dm⁻² day⁻¹) were on par with the genotype T₆ during the fourth cutting.

4.3 STABILITY ANALYSIS

4.3.1 Pooled Analysis of Variance

Eight genotypes of hedge lucerne were subjected to pooled analysis of variance for different characters *viz.*, plant height, number of branches plant⁻¹, length of branches plant⁻¹, number of leaves plant⁻¹, leaf to stem ratio, number of pods cluster⁻¹, green fodder yield, dry fodder yield, dry matter production, crude protein and crude fibre content over four locations. The analysis revealed that for the genotypes, G x E interactions were significant for all the characters studied. As the G x E interactions were significant for all the characters, further analysis were done for estimating the stability parameters (Table 19).

The total sum of squares is partitioned into genotypes, Environments + (Genotype x Environment) and pooled error in the ANOVA. The mean squares due to E+ (G x E) were significant for the characters like plant height, number of branches plant⁻¹, number of leaves plant⁻¹, green fodder yield, dry fodder yield, dry matter production, crude protein and crude fibre content prioritizing the

Table 19 . Pooled Analysis of Variance (mean square) for different quantitative traits of hedge lucerne over four locations

Source of variation	df	Plant height	Number of branches plant ⁻¹	Length of branches plant ⁻¹	Number of leaves plant ⁻¹	Leaf to stem ratio	Number of pods cluster ⁻¹	Green fodder yield	Dry fodder yield	Dry matter production	Crude protein	Crude fibre
Genotypes	7	3946.99**	105.44**	4781.32**	4494.68**	0.13**	12.38**	1145.58**	195.43**	357.00**	331.14**	114.93**
Environment	3	285.86**	20.33**	14.26**	468.87**	0.03**	0.50**	249.09**	135.69**	152.39**	56.74**	51.39**
G x E interaction	21	16.23**	1.12**	8.96**	148.98**	0.004**	0.09**	54.08**	4.75**	6.43**	0.82**	0.82**
Error	96	1.31	0.46	0.40	12.99	0.0021	0.02	12.21	0.20	0.25	0.35	0.62

** Significant at 1%, * Significant at 5%

presence of G x E interaction for these traits. The mean sum of squares due to genotype was significant for the eleven characters under study (Table 20). The sum of squares due to E+ (G x E) was further partitioned into that of Environment (Linear), Genotype x Environment linear and pooled deviation (Table 20). The linear component of Environment were significant for the characters like plant height, number of branches plant⁻¹, length of branches plant⁻¹, number of leaves plant⁻¹, leaf to stem ratio, number of pods cluster⁻¹, green fodder yield, dry fodder yield, dry matter production, crude protein and crude fibre content. The variation due to G x E (linear) were significant for the characters like plant height, number of branches plant⁻¹, number of leaves plant⁻¹, leaf to stem ratio, green fodder yield and dry fodder yield. The non linear component, pooled deviation were significant for the characters like length of branches plant⁻¹, number of leaves plant⁻¹, dry fodder yield and dry matter production indicating the importance of both linear and non linear components.

4.3.2 Environmental indices

The environmental indices of eleven characters is described in the Table 21. It was found that Vellayani was favourable for all of the characters whereas Thrissur was unfavourable for all the characters. Kottarakkara was highly favourable for number of leaves and number of pods while, Ambalavayal was favourable for number of branches and dry fodder yield.

4.3.3 Stability parameters

The estimation of stability parameters i.e., mean(μ), regression coefficient (b_i) and deviation from regression (S^2_{di}) for eleven characters are furnished below (Table 22).

4.3.3.1 Plant height

The maximum plant height was observed for the genotype T₈ (116.75 cm) followed by T₁ (112.87 cm) and the lowest was observed for the genotype T₃ (71.03 cm).

Table 20. Analysis of Variance (mean square) for mean data of different quantitative traits of hedge lucerne over four locations

Source of variations	df	Plant height	Number of branches plant ⁻¹	Length of branches plant ⁻¹	Number of leaves plant ⁻¹	Leaf to stem ratio	Number of pods cluster ⁻¹	Green fodder yield	Dry fodder yield	Dry matter production	Crude protein	Crude fibre
Genotypes	7	986.74**	26.36**	1195.33**	1123.67**	0.0321**	3.09**	286.39**	48.86**	89.25**	82.78**	28.73**
E+ (G x E)	24	12.48**	0.88**	2.41	47.24*	0.0012	0.04	19.61*	5.27**	6.17**	1.95**	1.79**
Environments	3	71.46**	5.08**	3.56	117.22**	0.0017	0.13**	62.27**	33.92**	38.09**	14.18**	12.84**
Environments (Lin)	1	214.39**	15.25**	10.70*	351.65**	0.0185**	0.37**	186.82**	101.77**	114.29**	42.55**	38.54**
G x E(Lin)	7	7.88**	0.56**	2.78	63.53*	0.0015**	0.03	32.80**	2.52**	2.15	0.27	0.38
Pooled deviation	16	1.87	0.12	1.72**	21.08**	0.0007	0.02	3.39	0.46**	1.17**	0.15	0.10
Pooled error	96	1.31	0.11	0.40	3.25	0.0005	0.02	3.05	0.20	0.25	0.35	0.62
Total	31	232.48	6.63	271.78	290.31	0.0085	0.73	79.86	15.12	24.93	20.21	7.87

** Significant at 1%, * Significant at 5%

Table 21. Estimates of environmental indices (I_j) for each character of hedge lucerne under different locations

Sl. No.	Character	Vellayani	Kottarakkara	Thrissur	Ambalavayal
1.	Plant height	2.28	-0.52	-4.03	2.26
2.	Number of branches	0.56	-0.20	-1.04	0.68
3.	Length of branches	0.47	0.08	-0.96	0.44
4.	Number of leaves	0.40	4.62	-4.73	0.27
5.	Leaf to stem ratio	0.85	-0.01	-0.03	0.03
6.	Number of pods	0.15	0.63	-0.12	0.04
7.	Green fodder yield	2.36	-0.66	-3.68	1.95
8.	Dry fodder yield	1.15	-0.15	-2.83	1.83
9.	Dry matter production	2.22	-0.21	-2.91	0.91
10.	Crude protein	1.34	-0.03	-1.81	0.49
11.	Crude fibre	1.26	-0.10	-1.70	0.58

Table 22. Mean performance and stability parameters for yield and its component traits in hedge lucerne over four locations

Genotype	Plant height (cm)			Number of branches			Length of branches (cm)			Number of leaves		
	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}
T ₁	112.87	1.07	0.30	5.10	1.10	-0.09	22.96	0.62	0.05	145.03	0.58	-2.56
T ₂	86.81	1.41	0.56	6.40	1.58	0.11	34.78	3.31	4.60**	119.66	3.91	70.07**
T ₃	71.03	1.46	3.11	10.31	1.30	-0.04	35.53	-1.7	5.77**	99.69	0.86	31.00
T ₄	94.53	1.75	2.11	4.29	1.05	-0.03	54.49	1.12	0.92	124.86	0.66	16.47
T ₅	101.10	0.75	-0.21	6.29	0.92	-0.10	62.49	2.09	1.48*	139.84	0.06	26.98
T ₆	110.31	0.89	-0.32	4.56	1.42	-0.02	47.66	1.05	-0.02	141.04	0.82	1.41
T ₇	86.56	0.10	5.37	3.98	0.78	0.08	33.41	0.29	0.26	112.17	0.46	-2.03
T ₈	116.75	0.56	1.44	10.34	-0.17	0.14	74.65	1.21	-0.09	143.33	0.66	1.39
Grand mean	97.49			6.41			45.75			128.20		

** Significant at 1%, * Significant at 5%

Table 22. continued

Genotype	Leaf to stem ratio			Number of pods			Green fodder yield			Dry fodder yield			Dry matter production		
	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}	Mean	b _i	S ² _{di}
T ₁	0.89	0.75	0.0005	1.51	2.12	0.011	112.35	1.12	-0.62	32.45	1.11	0.22	39.93	0.97	0.09
T ₂	0.71	2.52	-0.0004	1.58	0.24	0.001	101.28	1.34	-0.30	27.46	1.28	0.05	32.62	0.99	0.98*
T ₃	0.92	-0.17	-0.0002	1.43	1.30	0.019	90.45	2.38	-2.55	23.52	1.40	0.54	28.89	1.50	1.13*
T ₄	0.76	1.09	0.0004	2.77	1.79	0.056*	100.67	1.74	0.001	31.23	1.27	-0.03	37.59	0.94	1.04*
T ₅	0.67	0.37	-0.0004	2.14	1.23	-0.003	98.75	-1.45	9.20	27.03	0.02	0.83*	31.37	0.70	0.71*
T ₆	0.88	1.56	0.000	0.00	0.00	-0.005	110.48	1.26	-2.54	31.77	0.77	0.42	40.62	1.34	0.47
T ₇	0.84	1.18	0.0007	2.52	0.89	0.015	109.02	0.07	1.15	30.12	0.94	0.65*	40.32	0.27	3.95**
T ₈	0.83	0.68	0.0008	2.38	0.42	0.031	115.42	1.52	-1.61	34.25	1.93	0.55	40.04	1.25	0.49
Grand mean	0.81			1.79			104.93			29.74			36.42		

** Significant at 1%, * Significant at 5%

Table 22. continued

Genotype	Crude protein			Crude fibre		
	Mean	b_i	S^2_{di}	Mean	b_i	S^2_{di}
T ₁	21.38	1.42	0.02	23.35	0.97	0.098
T ₂	15.51	0.92	0.29	22.71	1.26	0.004
T ₃	14.59	1.21	-0.08	26.83	1.41	-0.139
T ₄	13.12	0.97	0.01	23.59	0.45	-0.143
T ₅	13.91	0.69	0.02	28.21	0.96	0.054
T ₆	20.26	0.89	-0.01	24.06	0.95	-0.116
T ₇	23.48	0.95	0.32	27.44	1.09	-0.027
T ₈	24.39	0.91	-0.06	29.95	0.89	-0.145
Grand mean	18.34			25.77		

** Significant at 1%, * Significant at 5%

Among the genotypes, T₁ genotype ($\mu = 112.87$, $b_i = 1.07$, $S^2_{di} = 0.30$) and the genotype T₆ ($\mu = 110.31$, $b_i = 0.89$, $S^2_{di} = -0.32$) were identified as stable ones with regression coefficient near unity and non-significant deviation from regression. The genotypes T₂, T₃ and T₄ recorded b_i value more than one (1.41, 1.46, 1.75 respectively) with non-significant minimum deviation from the regression (0.56, 3.11, 2.11 respectively). The genotypes T₅ and T₈ recorded b_i value of 0.75 and 0.58 with non-significant deviation from regression -0.21 and 1.44 respectively.

4.3.3.2 Number of branches plant⁻¹

The number of branches ranged from 4.29 (T₄) to 10.34 (T₈). The genotype T₁ ($\mu = 5.10$, $b_i = 1.10$, $S^2_{di} = -0.09$), T₄ ($\mu = 4.29$, $b_i = 1.05$, $S^2_{di} = -0.03$) and genotype T₅ ($\mu = 6.29$, $b_i = 0.92$, $S^2_{di} = -0.10$) recorded near unit regression with minimum non-significant deviation from regression. These genotypes were stable for different environments. The genotypes T₂, T₃ and T₆ showed b_i value more than one (1.58, 1.30 and 1.42 respectively) and minimum deviation from regression (0.11, -0.04 and -0.02 respectively) hence are stable and adaptable for rich environment. Regression coefficient less than one with minimum deviation from linearity was observed for the genotype T₇ ($\mu = 3.98$, $b_i = 0.78$, $S^2_{di} = 0.08$).

4.3.3.3 Length of branches

Length of the branches ranged from 74.65 cm (T₈) to 22.96 cm (T₁). T₃ ($b_i = 1.05$) was the genotype with near unit regression and non-significant deviation from regression, so T₃ was the stable genotype. T₈ was the genotype with more than unit regression and non-significant deviation from regression indicating high responsiveness towards the rich environment. The genotypes T₁ ($b_i = 0.62$) and T₇ ($b_i = 0.29$) showed less than unit regression with minimum non-significant deviation from regression. They were highly responsive in unfavourable environment.

4.3.3.4 Number of leaves plant⁻¹

The number of leaves varied from 145.03 (T₁) to T₃ (99.69). The genotype T₆ was identified as stable genotype with near unit regression ($b_i=0.82$) and non-significant deviation from regression. The genotype T₁ ($b_i=0.58$, $S^2_{di} = -2.56$) and T₇ ($b_i=0.46$, $S^2_{di} = -2.03$) were suitable for poor environment. They were performs under unfavourable environment.

4.3.3.5 Leaf to stem ratio

The leaf to stem ratio was highest for the genotype T₃ (0.92) and the lowest for T₅ (0.67). The genotype T₄ ($b_i=1.09$, $S^2_{di} = 0.0004$) and T₇ ($b_i=1.18$, $S^2_{di} = 0.0007$) showed near unit regression with non-significant deviation from regression, which were considered as stable genotypes across the environment. The genotype T₆ was identified as the genotype favourable for rich environment, with regression of more than one and non-significant deviation from regression. The genotype T₁ was identified as stable genotype under unfavourable environment.

4.3.3.6 Number of pods

The number of pods ranged from 0.00 (T₆) to 2.77 (T₄). The genotypes T₅ ($b_i=1.23$, $S^2_{di} = -0.003$) and T₇ ($b_i=0.89$, $S^2_{di} = 0.015$) identified as stable genotypes with near unit regression and minimum non-significant deviation from regression. The genotypes T₁ and T₃ were showed high mean, more than one regression coefficient and minimum non-significant deviation indicates high responsiveness towards environmental factors and performs well under rich environments. The genotype T₈ was stable for unfavourable environment with $b_i=0.42$ and $S^2_{di}=0.031$.

4.3.3.7 Green fodder yield

The green fodder yield varied from 90.45g (T₃) to 115.42g (T₈). The genotype T₁ had regression coefficient near unity with minimum non-significant deviation from regression, which specifies that the genotype was stable across the

environment. The genotypes T₂, T₄ and T₈ showed more than one regression coefficient with non-significant deviation from regression, which indicates the high responsiveness of these genotypes for rich environment.

4.3.3.8 Dry fodder yield

The dry fodder yield ranged from 23.52g (T₃) to 34.25g (T₈). The genotype T₁ ($\mu=32.45$, $b_i=1.11$, $S^2_{di}=0.22$) was stable genotype with near unity regression coefficient and non-significant deviation from regression. The genotypes T₂ ($\mu=27.46$, $b_i=1.28$, $S^2_{di}=0.05$) T₄ ($\mu=31.23$, $b_i=1.27$, $S^2_{di}=-0.03$) and T₈ ($\mu=34.25$, $b_i=1.93$, $S^2_{di}=0.55$) were identified as stable genotypes under favourable environment as they had more than one regression coefficient and non-significant deviation from regression. The genotype T₆ was identified as stable genotype under unfavourable environment due to the less than one regression coefficient and non-significant minimum deviation from regression.

4.3.3.9 Dry matter production

The dry matter production ranged from 40.62g (T₆) to 28.89g (T₃). The genotype T₁ ($\mu=39.93$, $b_i=0.97$, $S^2_{di}=0.09$) was identified as stable genotype with unit regression coefficient and minimum non-significant regression from deviation. The genotype T₆ ($\mu=40.62$, $b_i=1.34$, $S^2_{di}=0.47$) and T₈ ($\mu=40.04$, $b_i=1.25$, $S^2_{di}=0.49$) were stable for rich environment. They had more than one regression coefficient and non-significant deviation from regression.

4.3.3.9 Crude protein content

The crude protein content varied from 24.39% (T₈) to 13.12% (T₄) in the present study. The genotypes T₄ ($\mu=13.12$, $b_i=0.97$, $S^2_{di}=0.01$) and T₈ ($\mu=24.39$, $b_i=0.91$, $S^2_{di}=-0.06$) recoded high mean, near unit regression coefficient and non-significant deviation from regression for this trait. The genotype T₁ showed more than one regression coefficient and non-significant deviation from regression and was identified for rich environment. The genotype T₅ ($\mu=13.91$, $b_i=0.69$, $S^2_{di}=0.02$) exhibited less than unit value of regression and non-significant

Fig 9. Comparison of green fodder yield with population mean of the eight hedge lucerne genotypes

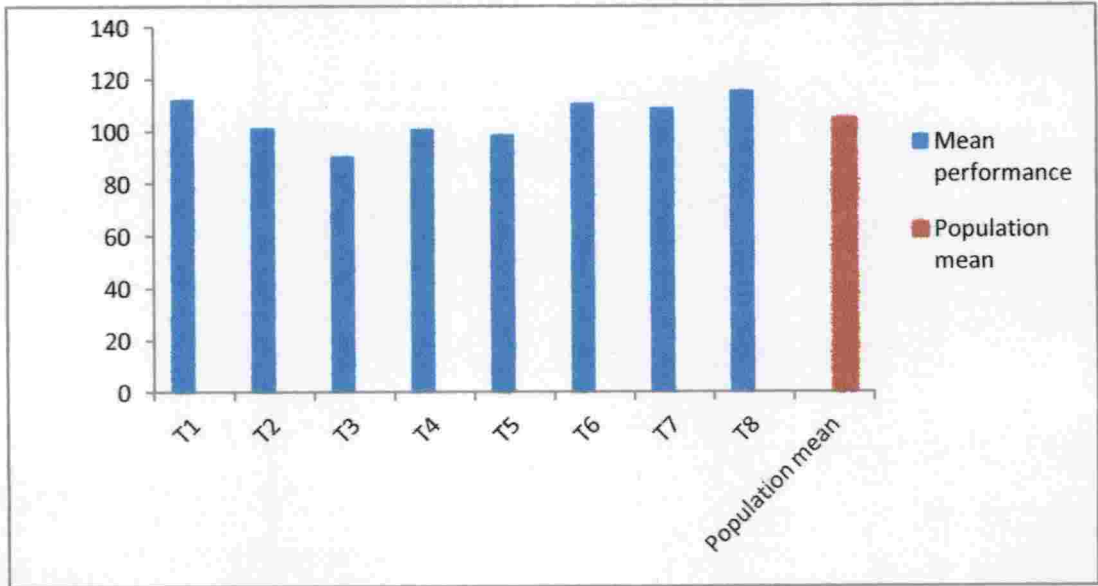


Fig 10. Comparison of dry fodder yield with population mean of the eight hedge lucerne genotypes

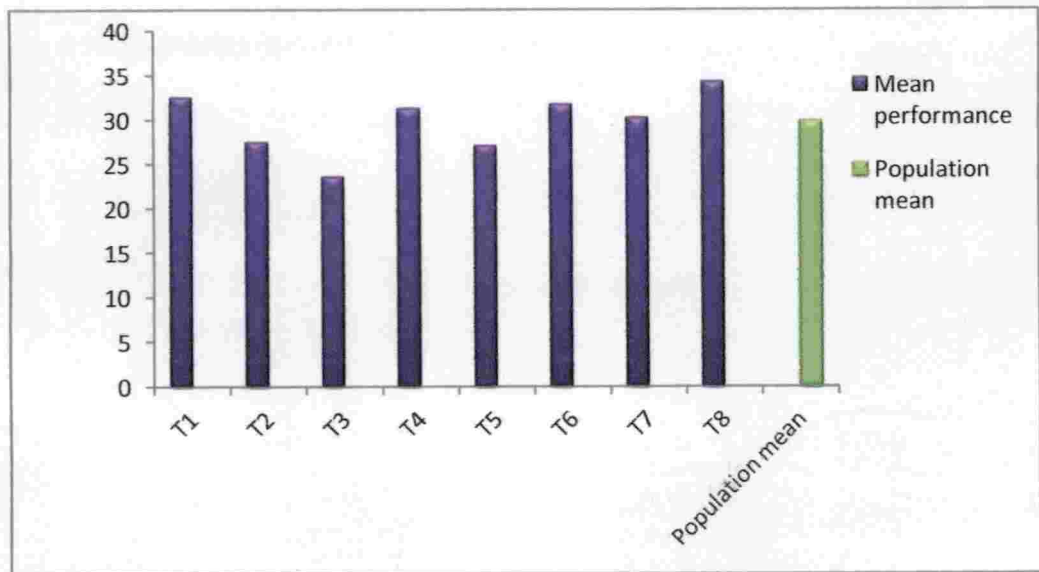


Fig 11. Comparison of dry matter production with population mean of the eight hedge lucerne genotypes

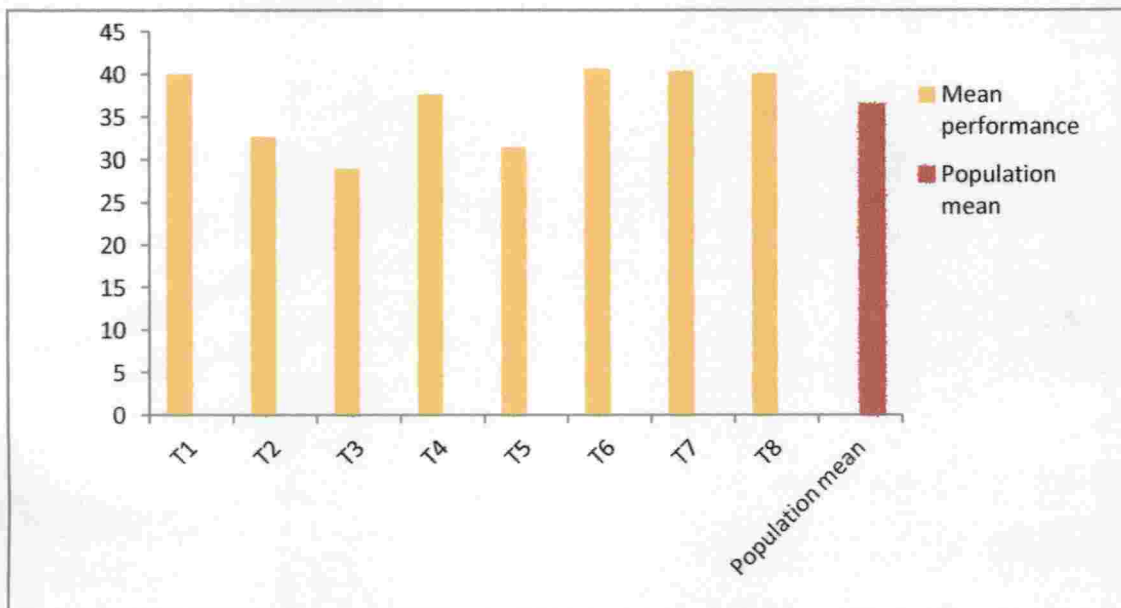


Fig12. Comparison of crude protein content with population mean of the eight hedge lucerne genotypes

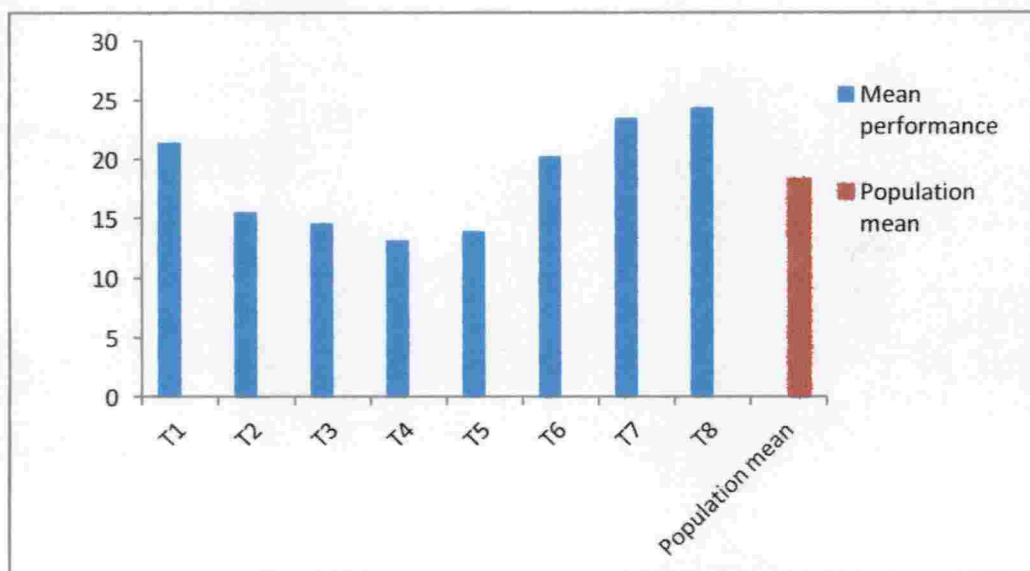
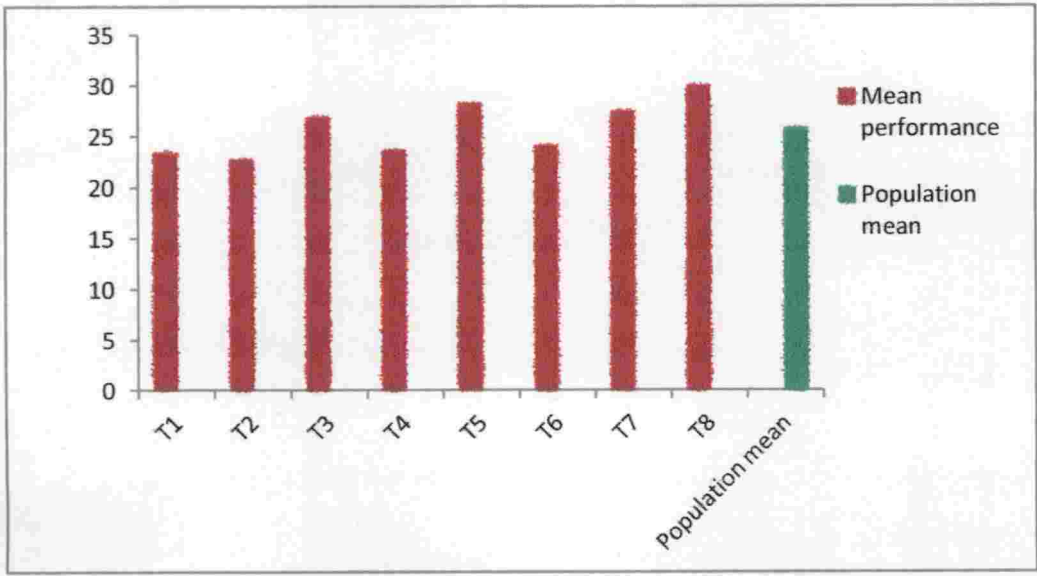


Fig 13. Comparison of crude fibre content with population mean of the eight hedge lucerne genotypes



deviation from regression and hence the genotype was suitable for poor environment.

4.3.3.10 Crude fibre content

The highest crude fibre content was recorded for the genotype T₈ (29.95%) and the lowest for the genotype T₂ (22.71%). The genotypes T₁ ($\mu=23.35$, $b_i=0.97$, $S^2_{di} = 0.098$), T₅ ($\mu=28.21$, $b_i=0.96$, $S^2_{di} = 0.054$) and T₇ ($\mu=27.44$, $b_i=1.09$, $S^2_{di} = -0.027$) showed high mean, near unit regression and minimum non-significant deviation from regression. These genotypes hence identified as stable across environment. The genotype T₂ ($\mu=22.71$, $b_i=1.26$, $S^2_{di} = 0.004$) was identified for rich environment based on stability parameters. The genotype T₄ showed regression coefficient less than one with non-significant deviation from regression, which was stable for unfavourable environments.

4.3.4 Identification of best genotypes for each location

Based on the ranking method, the best genotypes for each location were identified. The genotypes which had a score with mean value above mean \pm SE (m) was assigned Rank 1. Those genotypes with score between mean \pm SE (m) were assigned Rank 2 and the genotypes with score less than mean \pm SE (m) were assigned Rank 3. The ranks of each genotype over four locations are described in Table 23. Based on the rank given to each genotype for the characters number of leaves plant⁻¹, green fodder yield, dry fodder yield, dry matter production, crude protein and crude fibre, the genotypes T₁, T₆ and T₈ were best suited for all locations (Table 23).

4.3.5 Identification of character imparting stability

Based on the regression coefficient, deviation from regression and environmental indices, the characters plant height, green fodder yield, dry fodder yield, dry matter production and crude protein content can be accounted as the stability imparting characters.

Table 23. Ranking method for identifying best genotypes of hedge lucerne in each location

Table 23.1. Location-I (COA, Vellayani)

Genotype	Number of leaves		Green fodder yield (g plant ⁻¹)		Dry fodder yield (g plant ⁻¹)		Dry matter production (g plant ⁻¹)		Crude protein content (%)		Crude fibre content (%)		Total of Rank
	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	
T ₁	146.23	1	116.55	1	34.31	1	41.96	1	23.61	1	24.52	3	8
T ₂	113.12	3	103.09	3	28.60	3	34.02	3	16.93	3	24.64	3	18
T ₃	104.89	3	96.04	3	25.99	3	32.65	3	16.16	3	28.49	2	17
T ₄	128.20	2	105.84	2	32.81	1	40.21	1	14.16	3	24.25	3	12
T ₅	136.53	1	96.35	3	27.18	2	33.67	3	14.99	3	29.55	1	13
T ₆	142.74	1	113.34	1	31.94	2	43.08	1	21.66	2	25.06	3	10
T ₇	111.00	3	112.21	1	30.36	2	40.48	1	24.44	1	28.64	1	9
T ₈	146.12	1	119.57	1	35.89	1	43.04	1	25.46	1	30.99	1	6

Table 23.2. Location-II (KVK, Kottarakkara)

Genotype	Number of leaves		Green fodder yield (g plant ⁻¹)		Dry fodder yield (g plant ⁻¹)		Dry matter production (g plant ⁻¹)		Crude protein content (%)		Crude fibre content (%)		Total of Rank
	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	
T ₁	147.45	1	110.69	1	31.97	1	40.19	1	21.17	1	22.71	3	8
T ₂	143.87	1	101.99	2	27.25	2	32.30	3	16.02	3	22.25	3	14
T ₃	99.37	3	88.89	3	22.84	3	27.25	3	14.54	3	26.66	1	16
T ₄	124.68	2	97.42	2	31.11	1	36.12	2	13.14	3	23.44	3	13
T ₅	144.09	1	103.91	2	28.05	2	31.17	3	14.15	3	28.52	1	12
T ₆	143.26	1	109.93	1	32.23	1	41.12	1	19.93	2	24.03	3	9
T ₇	114.81	2	108.36	1	30.16	1	42.69	1	23.06	1	27.64	1	7
T ₈	144.99	1	112.99	1	33.12	1	38.87	1	24.46	1	29.58	1	6

Table 23. 3. Location-III (Thrissur)

Genotype	Number of leaves		Green fodder yield (g plant ⁻¹)		Dry fodder yield (g plant ⁻¹)		Dry matter production (g plant ⁻¹)		Crude protein content (%)		Crude fibre content (%)		Total of Rank
	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	
T ₁	142.19	1	108.76	1	29.36	1	36.91	1	18.96	1	21.83	3	8
T ₂	106.64	3	95.51	2	23.87	2	29.46	3	13.78	3	20.76	3	16
T ₃	91.70	3	89.75	3	19.62	3	25.05	3	12.37	3	24.39	2	17
T ₄	118.68	2	95.26	2	27.60	1	35.37	1	11.26	3	22.86	3	12
T ₅	143.32	1	102.04	1	26.52	2	29.60	2	12.65	3	26.50	1	10
T ₆	135.66	1	105.76	1	29.45	1	36.30	1	18.78	2	22.36	3	9
T ₇	110.33	2	110.43	1	27.60	1	38.65	1	21.74	1	25.41	2	8
T ₈	139.21	1	110.54	1	31.23	1	36.73	1	22.67	1	28.37	1	6

Table 23. 4. Location-IV (Ambalavayal)

Genotype	Number of leaves		Green fodder yield (g plant ⁻¹)		Dry fodder yield (g plant ⁻¹)		Dry matter production (g plant ⁻¹)		Crude protein content (%)		Crude fibre content (%)		Total of Rank
	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	Mean	Rank	
T ₁	144.26	1	113.39	1	34.18	1	40.64	1	21.82	1	24.35	3	8
T ₂	115.03	3	104.52	2	30.12	2	34.69	2	15.31	3	23.23	3	15
T ₃	102.81	3	94.37	3	25.62	3	30.64	3	15.32	3	27.78	2	17
T ₄	127.89	2	104.70	2	33.41	1	38.64	1	13.95	3	23.79	3	12
T ₅	135.43	1	95.11	3	26.36	3	31.06	3	13.88	3	28.28	1	14
T ₆	142.49	1	112.17	1	33.48	1	41.99	1	20.69	2	24.79	3	9
T ₇	112.54	3	112.20	1	32.67	2	39.48	1	24.71	1	27.97	2	10
T ₈	143.00	1	118.56	1	36.75	1	41.53	1	24.98	1	31.52	1	6

Plate 3. Stable genotypes of Hedge lucerne over all locations



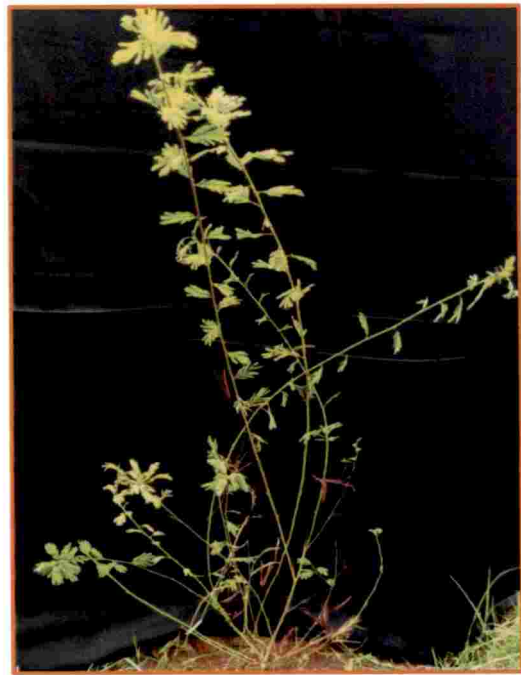
T₁ (IC 345276)



T₄ (IC 261839)



T₆ (IC 421199)



T₇ (TNDV 1)

Plate 4. Hedge lucerne suitable for favourable environments



T₂ (IC 343710)



T₈ (Thumburmuzhi local)

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Plate 5. Hedge lucerne genotypes suitable for unfavourable environment



T₅ (IC 90934)

DISCUSSION

5. DISCUSSION

Hedge lucerne is a herbaceous perennial legume belonging to the family Fabaceae. No poisonous principle is observed in foliage. Hedge lucerne is preferred by cattle for its palatable green fodder and adequate amount of crude protein. The alarming gap between demand and supply in availability of fodder in Kerala necessitates the production of high quality herbage devoid of anti-nutritional factors. However the genetic improvement achieved in hedge lucerne in terms of its productivity is very low.

The identification of superior genotypes remarks the success of crop improvement activities. Phenotypic stability and potential for high yield under favourable environment considered most in selecting superior genotypes. To analyze G x E interaction a number of parametric statistical procedures have been developed in which Eberhart and Russell model (1966) is the most widely used method. The nature of adaptation of genotypes can be studied from two main factors namely, regression coefficient (linear sensitivity) and the deviation from mean squares due to regression (non-linear sensitivity).

Based on the stability parameters the extent of deviation of yield and yield related characters over environments is studied to identify the best genotypes which are widely adapted. Multilocation testing of genotypes provides an opportunity for plant breeders to study the adaptability of genotypes to a particular environment and also to understand the stability of the genotypes over different environments. Thus information on Genotype x Environment interaction is of major importance to the plant breeders in identifying an improved stable genotype.

Realizing the importance of the stability analysis in performance of genotypes, in the present study eight genotypes of hedge lucerne were evaluated across four environments. The salient findings from this present study entitled "Genotype x Environment interaction in Hedge lucerne (*Desmanthus virgatus* (L.)

Willd.) for yield and quality” have been critically analyzed and discussed in light of available literature under the following subheads.

1. Pooled analysis of variance
2. Stability analysis for yield and its attributing traits

5.1. POOLED ANALYSIS OF VARIANCE

From the pooled analysis of variance, the genotypes showed significant differences for all the characters studied, which revealed the presence of genetic variability among the genotypes. Significant differences among the genotypes gave greater opportunity for selecting suitable genotypes with high mean value for all the characters of interest. Environments were highly significant for the characters viz., plant height (cm), number of branches plant⁻¹, length of branches plant⁻¹(cm), number of leaves plant⁻¹, leaf to stem ratio, number of pods cluster⁻¹, green fodder yield (g), dry fodder yield (g), dry matter production (g), crude protein content (%) and crude fibre content (%/) suggesting the divergence among growing environments. The effect of genotype x environment were significant for all the characters studied, which indicated the differential response of genotypes to varying environment. Therefore, these genotypes must be evaluated over a wide range of environments where they are ultimately cultivated for commercial purposes. Similar findings were also reported by Palathingal (2003) in rice, Bikash *et al.* (2013) in pearl millet, Preeti *et al.* (2016) in wheat, Mehraj *et al.* (2017) in oats and Patil *et al.* (2017) in okra.

The joint regression analysis by stability analysis revealed that G x E interaction (linear) was highly significant for all the characters studied viz., plant height (cm), number of branches plant⁻¹, length of branches plant⁻¹, number of leaves plant⁻¹, leaf to stem ratio, number of pods cluster⁻¹, green fodder yield, dry fodder yield, dry matter production, crude protein content and crude fibre content indicating that the genotypes had divergent linear response to the environmental changes for these characters. Comparable findings were reported by Saranya (2016) for the characters plant height, number of branches, number of leaves

plant⁻¹ and total fresh weight in Neelamari, Preeti *et al.* (2016) for the characters plant height and dry matter yield in wheat, Mehraj (2017) for the traits plant height, fodder yield ha⁻¹, green fodder yield and leaf to stem ratio in oats.

5.2. STABILITY ANALYSIS FOR YIELD AND ITS ATTRIBUTING TRAITS

Several workers proposed different models for identifying stable genotypes which exhibit least interaction with environments. Finlay and Wilkinson (1963) developed a dynamic approach for interpretation of varying environments. They considered mean value of the genotype and their regression coefficient. But Eberhart and Russell improved this model by adding another stability parameter, *i.e.*, the deviation from regression and provided fresh approach to G x E interaction analysis. Eberhart and Russell model (1966) considered three stability parameters like (i) mean performance (μ), (ii) regression coefficient (b_i) and (iii) deviation from regression (S^2_{di}). Linear component of G x E interaction is measured by using b_i value and also gives an idea about response of genotype. G x E interaction of unpredictable type (*i.e.*, predictable or unpredictable type) is measured from S^2_{di} value.

The result interpretation of present study was done by using the parameters like regression coefficient, mean value and deviation from regression for stability. Once the genotypes were found to be stable based on non-significant deviation from regression ($S^2_{di}=0$), then the type of stability was based on regression coefficient and mean value. If b_i is equal to unity, a genotype is considered as stable or has the same performance in all the environment, if b_i is more than unity, it is considered to have less than average stability or good performance in favourable environments and if b_i is less than unity, it is suggested to have above average stability or good performance under poor environments (Eberhart and Rusell, 1966).

5.2.1. Plant height (cm)

Two genotypes, T₁ and T₆ have recorded the highest mean value than population mean for the character plant height with The regression coefficient

near unity with non-significant deviation from regression, suggested that these two genotypes were stable for this character. The genotype T₂ had more than unity regression (1.41) and non-significant deviation from regression (0.56) and was found to be stable for favourable environments. The genotype T₅ was found to be stable for unfavourable environments with less than unity regression and non-significant deviation from regression. Similar results were observed by Sunil (2004) in turmeric, Panwar *et al.* (2011) in ocimum, Javia (2014) in okra, Saranya (2016) in neelamari, Preeti *et al.* (2016) in wheat, Patil *et al.* (2017) in oats also observed stability for the character plant height.

5.2.2. Number of branches plant⁻¹

Among the genotypes evaluated T₁ (5.10), T₄ (4.29) and T₅ (6.29) with regression near to unity and minimum deviation from regression was widely adapted with average stability. The genotypes T₃ (10.31) and T₆ (4.56) had regression coefficient greater than unity and non-significant deviation from regression, indicated that these genotypes were stable under favourable environments with predictable performance. Less than unity regression and non-significant deviation from regression was recorded for the genotype T₇ (3.98), which indicated the adaptability of the genotype under unfavourable environment. Similar findings were recorded by Ottai *et al.* (2006) in roselle, Abou *et al.* (2006) in white mustard, Saranya (2016) in neelamari and Patil *et al.* (2017) in okra.

5.2.3. Length of branches plant⁻¹

The genotype T₆ was considered as stable for the character length of branches plant⁻¹ because of a regression coefficient near unity. The genotypes T₈ (b_i=1.21) recorded more than unit regression and non-significant deviation from regression, indicated that the genotype was suitable for favourable environment. The genotypes T₁ and T₇ reported less than unity regression for the character length of branches plant⁻¹ with minimum non-significant deviation from regression, which suggested that these genotypes were suitable for unfavourable environments. Similar results for variation in stability parameters for the character

length of branches plant⁻¹ were observed by Singh and Arya (2014) in *Vigna radiata* and Ramesh *et al.* (2017) in pigeon pea.

5.2.4. Number of leaves plant⁻¹

The genotype T₆ which had the highest mean value (141.04), near unity regression (0.82) and minimum non-significant deviation from regression (1.41), was considered as a stable genotype. The genotypes T₁ and T₇ were stable under unfavourable environments. Similar results for variability in stability parameters for the trait number of leaves plant⁻¹ was reported by Saranya (2016) in neelamari, Nanavati (2016) in maize and Mehraj *et al.* (2017) in oats.

5.2.5. Leaf to stem ratio

Among the genotypes T₄ ($\mu=0.76$, $b_i=1.09$, $S^2_{di}=0.0004$) and T₇ ($\mu=0.84$, $b_i=1.18$, $S^2_{di}=0.0007$) were considered as the stable genotypes because of the desirable mean, near unity regression and mean deviation from linearity, which can be suggested for wider environments. Whereas, the genotype T₆ ($\mu=0.88$, $b_i=1.56$, $S^2_{di}=0.000$) was suitable for favourable environments due to greater than unit regression and non-significant deviation from linearity, while T₁ ($\mu=0.89$, $b_i=0.75$, $S^2_{di}=0.0005$) genotype was suitable for unfavourable environments. These results were in accordance with the results of Nanavati (2016) in maize, and Mehraj *et al.* (2017) in oats.

5.2.6. Number of pods cluster⁻¹

The genotypes T₅ and T₇ recorded near unit regression (1.23 and 0.89) with non-significant S^2_{di} value (-0.003 and 0.015), indicated that the genotypes were stable across the environment. The genotypes T₁ and T₃ had higher than unity and minimum non-significant deviation from regression, which indicated that these genotypes were adaptable for favourable environments. The genotype T₈ was suitable for unfavourable environment and had regression coefficient of less than unity and non-significant deviation from regression. These findings are agreement with Ramesh *et al.* (2017) in pigeon pea.

5.2.7. Green fodder yield

The genotype had the high mean yield (112.35 g) than the population yield (104.93 g), near unit regression (1.12) and non-significant minimum deviation from regression (-0.62) was considered as stable genotype for the character green fodder yield. The genotype T₄ and T₈ genotypes performed well under favourable environment with regression coefficient of 1.74 and 1.52 respectively. The variable stability parameters were reported in green fodder yield by Nanavati (2016) in forage maize and Mehraj *et al.* (2017) in oats.

5.2.8. Dry fodder yield

The genotype T₁ ($\mu=32.45$, $b_i=1.11$, $S^2_{di}=0.22$) possessed higher dry fodder yield than the population mean and were considered as highly adaptable genotypes having average stability and was expected to perform well in all the environment. The genotypes T₂, T₄ and T₈ had higher mean than population mean, more than unity regression with non-significant deviation from linearity, which suggested that these genotypes were stable under favourable environment and perform better under rich environment. The genotype T₆ ($\mu=31.77$, $b_i=0.77$, $S^2_{di}=0.42$) were stable for unfavourable environment with high mean, less than unity regression and minimum deviation from regression. Similar findings for the variable stability parameters in dry fodder yield was noticed by Bikash *et al.* (2013) in pearl millet and Mehraj *et al.* (2017) in oats.

5.2.9. Dry matter production

The genotypes T₁, T₆ and T₈ had higher mean performance for dry matter production than the population mean. The genotype T₁ was stable with regression coefficient near unity ($b_i=0.97$) and non-significant deviation from regression ($S^2_{di}=0.09$). The genotypes T₆ ($b_i=1.34$) and T₈ ($b_i=1.25$) had regression coefficient greater than unity and minimum deviation from regression, which indicated that these genotypes were stable for favourable environment. The same findings for variation in stability parameters was observed by Preeti *et al.* (2016) in wheat.

5.2.10. Crude protein content

Two genotypes, T₄ and T₈ were stable for the character crude protein content in all the environment with regression coefficient of 0.97 and 0.91, with minimum deviation from regression of 0.01 and -0.06 respectively. The genotype T₁ ($b_i=1.42$) was stable under favourable environment. The genotype T₅ ($b_i=0.69$) had less than unity regression and non-significant deviation from regression (0.02) and was stable under unfavourable environment and poor environment. Comparable findings for the variable stability parameters for protein content was reported by Saeed *et al.* (1985) in sorghum, Peterson *et al.* (1992) in wheat, Shi *et al.* (1999) in rice, Gurmu *et al.* (2009) in soybean.

5.2.11. Crude fibre content

Among the genotypes, T₁, T₅ and T₇ had near unity regression coefficient (0.97, 0.96 and 1.09) and minimum deviation from regression (0.098, 0.054 and -0.027) and their performances for the character crude fibre content can be predicted. They were well adapted for all the environments. The genotype T₂ ($b_i=1.26$, $S^2_{di}=0.004$) was found to be suitable for unfavourable environment. The genotype T₄ ($b_i=0.45$, $S^2_{di}= -0.143$) was suitable for poor environment for the trait crude fibre content.

Table 24. Comparison of hedge lucerne genotypes on the basis of mean performance and stability parameters

Sl. No.	Character	Stable	Favourable environment	Unfavourable environment	Population mean
1.	Plant height (cm)	T ₁ , T ₆	T ₂	T ₅	97.49
2.	Number of branches	T ₁ , T ₄ , T ₅	T ₃ , T ₆	T ₇	6.41
3.	Length of branches (cm)	T ₆	T ₈	T ₁ , T ₈	45.75
4.	Number of leaves	T ₆	-	T ₁ , T ₇	128.20
5.	Leaf to stem ratio	T ₄ , T ₇	T ₆	T ₁	0.81
6.	Number of pods	T ₅ , T ₇	T ₁ , T ₃	T ₈	1.79
7.	Green fodder yield (g plant ⁻¹)	T ₁	T ₂ , T ₄ , T ₈	-	104.93
8.	Dry fodder yield (g plant ⁻¹)	T ₁ , T ₈	T ₂ , T ₄ , T ₈	T ₆	29.74
9.	Dry matter production (g plant ⁻¹)	T ₁	T ₆ , T ₈	-	36.42
10.	Crude protein	T ₄ , T ₈	T ₁	T ₆	18.34
11.	Crude fibre	T ₁ , T ₇	T ₂	T ₅	25.77

SUMMARY

6. SUMMARY

The present study on “Genotype×Environment interaction in hedge lucerne (*Desmanthus virgatus* (L.) Willd.) for yield and quality” was carried out to identify stable genotypes of eight *Desmanthus virgatus* in varied environments with respect to yield and quality in four locations of Kerala viz., College of Agriculture, Vellayani, College of Horticulture, Thrissur, Krishi Vigyan Kendra, Kottarakkara, Kollam and Regional Agricultural Research Station, Ambalavayal, Wayanad.

The eight genotypes of hedge lucerne viz., T₁ (IC 345276), T₂ (IC 343710), T₃ (IC 89910), T₄ (IC 261839), T₅ (IC 90934), T₆ (IC 421199), T₇ (TNDV 1) and T₈ (Thumburmuzhi local) were evaluated in a Randomized Block Design (RBD) with four replications over four locations during 2017-2018. Eberhart and Russell model (1966) was used to analyze the stability and adaptability of yield and yield related characters of these genotypes. The present study showed significant mean squares due to genotypes for yield, quality and other component characters and this revealed the existence of high variability among the genotype studied. The mean squares due to Genotype x Environment interaction were significant for all the characters indicating the varying response of genotypes towards changing environment. Hence further analysis was done to assess the stability of genotypes.

In the stability analysis, the mean squares due to Environments + (Genotype x Environment) were significant for the characters viz., plant height, number of branches plant⁻¹, number of leaves plant⁻¹, green fodder yield, dry fodder yield, dry matter production, dry matter production, crude protein content and crude fibre content strengthening the presence of G x E interaction for these characters. The sum of squares due to E + (G x E) was further partitioned into Environment (linear), Genotype x Environment (linear) and pooled deviation. Environment (linear) was significant for the characters such as, plant height, number of branches plant⁻¹, length of branches plant⁻¹, number of leaves plant⁻¹, leaf to stem ratio, number of pods plant⁻¹, green fodder yield, dry fodder yield,

dry matter production, crude protein content and crude fibre content. The mean squares due to Genotype x Environment (linear) was significant for the traits like, plant height, number of branches plant⁻¹, number of leaves plant⁻¹, leaf to stem ratio, green fodder yield and dry fodder yield. This indicated that the major component for differences in stability was due to both linear and non linear components.

The estimation of environmental indices for all the characters in all the four locations (Vellayani, Kottarakkara, Thrissur and Ambalavayal) revealed that Vellayani and Ambalavayal were most favourable or suitable environment for cultivation of hedge lucerne and Thrissur was the unfavourable environment for hedge lucerne cultivation.

The stability of the genotype was measured from the mean performance of a genotype along with two stability parameters viz., regression coefficient (b_i) and deviation from regression coefficient (S^2_{di}).

The genotypes T₁ (IC 345276), T₄ (IC 261839), T₆ (IC 421199) and T₇ (TNDV 1) were identified as stable genotypes having regression coefficient near unity and non-significant deviation from regression with wider adaptability over environment for most of the characters. The genotype T₁ (IC 345276) was stable over all locations for different characters such as plant height, number of branches, green fodder yield, dry fodder yield, dry matter production and crude fibre. The genotype T₄ (IC 261839) was stable for the characters number of branches, leaf to stem ratio and crude protein across the locations. The genotype T₆ (IC 421199) was stable over locations for the characters viz., plant height, length of branches and number of leaves, while the genotype T₇ (TNDV 1) was stable for leaf to stem ratio, number of pods and crude fibre.

The genotypes T₂ (IC 343710) and T₈ (Thumburmuzhi local) were stable genotypes for favourable environment. The genotype T₂ (IC 343710) showed stable performance for the characters such as leaf to stem ratio, green fodder yield, dry fodder yield and crude fibre. The genotype T₈ (Thumburmuzhi local) was

stable in favourable environment for length of branches, green fodder yield, dry fodder yield and dry matter production. The genotype T₅ (IC 90934) was found to be stable for the characters plant height and crude fibre in unfavourable environments.

The present study revealed that the genotypes T₁ (IC 345276), T₄ (IC 261839), T₆ (IC 421199) and T₇ (TNDV 1) were stable over the four different locations *viz.*, College of Agriculture, Vellayani, College of Horticulture, Thrissur, Krishi Vigyan Kendra, Kottarakkara and Regional Agricultural Research Station, Ambalavayal, Wayanad. The genotypes T₂ (IC 343710) and T₈ (Thumburmuzhi local) showed stable performance under favourable environments *viz.*, College of Agriculture, Vellayani and Regional Agricultural Research Station, Ambalavayal, Wayanad, while the genotype T₅ (IC 90934) was suitable for unfavourable environment *i.e.*, College of Horticulture, Thrissur. The superior genotypes identified in the present study can be further promoted to farm trials before releasing them as a variety.

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**GENOTYPE×ENVIRONMENT INTERACTION IN
HEDGE LUCERNE (*Desmanthus virgatus* (L.) Willd.) FOR
YIELD AND QUALITY**

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Abstract of the thesis

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ABSTRACT

The present work on “Genotype×Environment interaction in hedge lucerne (*Desmanthus virgatus* (L.) Willd.) for yield and quality” was carried out across four locations in Kerala viz., College of Agriculture, Vellayani, College of Horticulture, Thrissur, Krishi Vigyan Kendra, Kottarakkara and Regional Agricultural Research Station, Ambalavayal, Wayanad, during 2016-2018 with an objective to identify stable genotypes of *Desmanthus virgatus* in varied environments with respect to yield and quality.

The eight genotypes of hedge lucerne viz., T₁ (IC 345276), T₂ (IC 343710), T₃ (IC 89910), T₄ (IC 261839), T₅ (IC 90934), T₆ (IC 421199), T₇ (TNDV 1) and T₈ (Thumburmuzhi local) were evaluated in a Randomized Block Design (RBD) with four replications over four locations during 2017-2018. Eberhart and Russell model (1966) was used to analyze the stability and adaptability of yield and yield related characters of these genotypes. Based on the environmental indices, COA, Vellayani and RARS, Ambalavayal was found most favourable for all of the characters, while COH, Thrissur was unfavourable for all the characters studied.

In the pooled analysis of variance for evaluation of hedge lucerne over locations, significant differences among the genotypes and environments were noticed for all the characters studied, suggesting that genotypes interacted significantly with environments.

Stability analysis revealed that the genotype T₁ (IC 345276) was stable over all locations for different characters such as plant height, number of branches, green fodder yield, dry fodder yield, dry matter production and crude fibre. The genotype T₄ (IC 261839) was stable for the characters number of branches, leaf to stem ratio and crude protein across the locations. The genotype T₆ (IC 421199) was stable over locations for the characters viz., plant height, length of branches and number of leaves,

while the genotype T₇ (TNDV 1) was stable for leaf to stem ratio, number of pods and crude fibre.

The genotypes T₂ (IC 343710) and T₈ (Thumburmuzhi local) were identified as stable genotypes for favourable environments. The genotype T₂ (IC 343710) showed stable performance for the characters such as leaf to stem ratio, green fodder yield, dry fodder yield and crude fibre. The genotype T₈ (Thumburmuzhi local) was stable in favourable environment for length of branches, green fodder yield, dry fodder yield and dry matter production. The genotype T₅ (IC 90934) was found to be stable for the characters plant height and crude fibre in unfavourable environments.

The present study revealed that the genotypes T₁ (IC 345276), T₄ (IC 261839), T₆ (IC 421199) and T₇ (TNDV 1) were stable over the four different locations viz., College of Agriculture, Vellayani, College of Horticulture, Thrissur, Krishi Vigyan Kendra, Kottarakkara and Regional Agricultural Research Station, Ambalavayal, Wayanad. The genotypes T₂ (IC 343710) and T₈ (Thumburmuzhi local) showed stable performance under favourable environments viz., College of Agriculture, Vellayani and Regional Agricultural Research Station, Ambalavayal, Wayanad, while the genotype T₅ (IC 90934) was suitable for unfavourable environment i.e., College of Horticulture, Thrissur.

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