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# Genotypic Variability in Yorkshire Fog Grass 

(Holcus lanatus L.)

A thesis<br>presented in partial fulfilment of the requirements<br>for the degree of<br>Master of Agricultural Science<br>in Agronomy<br>at<br>Massey University

Muangthong Thuantavee
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#### Abstract

Plant to plant genotypic variation in New-Zealand Yorkshire-Fog grass was examined in order to quantify the relative importance of average gene effects, dominance, epistasis and environment. The plant variability was contrasted also against topodeme variation.


Plants were grown under glasshouse conditions ( $20^{\circ}-25^{\circ} \mathrm{C}$ ), using vernalization and sixteen hour daylight to encourage growth and flowering. The confounding effect of bench position was removed by regression adjustment.

Fifty half-sib lines representing ten diverse New Zealand topodemes were examined in a one-way mating design, laid out as a randomized complete block experiment.

In general, half-sib and plant variances were much larger than the topodeme variance. This supports earlier findings that there are no major topodeme differences in New Zealand Yorkshire Fog grass germplasm.

The broad-sense heritability estimates which indicated total genotypic contribution varied from low to high. Most botanical, flowering and tillering characters had a medium to high values while the agronomic characters had medium to low estimates.

The attributes with medium to high narrow-sense heritability are several measures of leaf size, tiller development, purple colour, plant height and erectness, flavanols and panicle width. Breeding methods, such as mass selection, line selection, line breeding or simple recurrent selection should ,therefore, be appropriate for these.

The attributes with medium to high heterotic-sense heritability are leaf tensile strength, leaf hairiness, old disease, flowering period, panicle length and compactness and several aspects of tiller production. Breeding methods, such as recurrent selection with progeny testing or top cross progeny tests for high specific
combining ability should be useful, including synthetic cultivars and some kinds of recurrent bulks.

Of particular interest was the finding that there was more genetic variability for the duration of tillering and flowering periods than for tiller numbers or flower initiation. There was also evidence that the genetic activity controlling tiller number changed as the tillers aged.

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## INTRODUCTION

Yorkshire Fog grass has been judged as one of the significant grasses for farm productivity (Basnyat, 1957; Munro, 1961). It has always been valuable as a pioneer grass in drained peat swamp areas (Basnyat, 1957). It is also useful in infertile, unstable, poorly drained soil (Munro, 1961; Davies et al., 1971; Morrison and Idle, 1972; Rumball, 1983). It is capable of establishing well in humid hill county, and on unploughable steep hills (Basnyat, 1957; Hughes and Nicholson, 1961;). On such area, $H$. lanatus is one of the earliest grasses to start growth in the spring and its subsequent growth was also notable (Herriot, 1975). It has been proposed as a 'nurse' species for sown L. perenne and Trifolium rapens, for which it would consolidate the soil, protect over grazing, and speed up the fertility cycle (Thomas, 1936; Davies, 1940). Furthermore, its good persistence has been used to control erosion (Dunbar, 1974; Hornung, 1976).

Yorkshire Fog grass is more suitable for less intensive farming system, typically dairy pasture and upland sheep farms (Munro, 1961). Its growth habit and vegetativereproductive cycle make it a good candidate for a lenient system of defoliation (Levy, 1955; Beddows, 1961). Its grazing tolerance lies between perennial ryegrass and cocksfoot (Mitchell, 1956). In mixed swards and under infrequent grazing regime, $H$. lanatus dominated L.perenne (Watt, 1987) and its ground cover over 4 year in Oxford has increased from $18 \%$ to $43 \%$ (Haggars and Ellliot, 1978).

Yorkshire Fog grass is believed to have been introduced into New Zealand either as a seed impurity or a hay grass in eighteenth century (Cheeseman, 1923), and since then as a volunteer, it contributed much of New Zealand's pasture production (Munro, 1961). Massey University has been interested in Yorkshire Fog grass since 1950 (Basnyat, 1957). The first synthetic variety "Massey Basyn" was released and proved to be prominent in several areas (Robinson et al., 1980; McAdam, 1984; Watt, 1987). Evaluation on Yorkshire Fog grass germplasm of New Zealand collection was carried on by Teow (1978). In addition, factors involving sheep palatability were determined by Cameron (1979). The broad-sense heritability estimates were also initially figured out pertinent to topodeme basis.

Following previous studies, this investigation has been set up to increase the genetical knowledge of Yorkshire Fog grass. An attempt has been made to unravel the heritabilities pertinent to individual plant basis. Comparison between plant variation and topodeme variation was also carried out.

## CHAPTER 1

## LITERATURE REVIEWS

### 1.1 Yorkshire Fog Grass

### 1.1.1 Agro-botany and Agronomy

Yorkshire Fog grass or velvet grass (Holcus lanatus) is probably a native of the Iberian Peninsular (Spain and Portugal) (Vinal and Hein, 1937). It is a tufted, softly hairy perennial which can adapt to a wide range of environmental conditions, but predominates in moist and low-fertility soil (Hubbard, 1968). H. lanatus is widespread in the temperate region around the world from the limits of Northern Scandinavia and Iceland to the Caucasus mountains, North and West Africa, North America, South America, Australia, New Zealand and several sub-antarctic islands (Hulten, 1950; Bocher and Larsen, 1958; Beddows, 1961; Munro, 1961; Watton, 1975).

Although its distribution is by accident rather than design, and has caused certain weed problems (Harkess and Hope, 1974), several workers have claimed its considerable contribution to fodder production; for example, in England and Wales (Forbes et al., 1980; Watt, 1987), in Scotland (Swift, et al., 1983), in Chile, Southern Brazil, and Hawaii (Whyte, Moir and Cooper, 1959), and in Falkland Islands (Davies, et al., 1971).
H. lanatus can germinate over a wide range of soil temperature (Watt, 1976). Seedling emergence, however, is progressively delayed in accordance with decrease in mean soil temperature (Hart, 1961). It germinates well either at $22^{\circ} \mathrm{C}$. under continuous light or in diurnal fluctuating temperature ( $10^{\circ} \mathrm{C}$ and $20^{\circ} \mathrm{C}$ ) under dark condition (Thompson, Grime and Mason, 1977). It also germinates readily in the light at normal room temperature (Grime and Jarvis, 1975). Moist soil condition is indispensable for optimum germination (Watt, 1976). Most freshly collected seeds germinate rapidly in moist conditions (Watt, 1977).
H. lanatus thrives well at temperature between $12.8^{\circ} \mathrm{C}$ and $29.4^{\circ} \mathrm{C}$ (Mitchell and Lucanus, 1962). However, growth is poor at $35^{\circ} \mathrm{C}$ (Mitchell, 1956) and leafy shoot ceases development at $5^{\circ} \mathrm{C}$ (Beddows, 1961). Because it grows relatively well at low temperature, many workers regard it as a good winter grower (Munro, 1961; Hubbard, 1945; Watkin and Robinson, 1974). It is able to establish over a wide altitude range (Basnyat, 1957) and spread evenly over altitudes up to 400 m . and on all slopes up to $50^{\circ}$ (Watt, 1976).

It can inhabit on a wide light regime ranging from dense shade to open and sunny (Levy, 1970). The broader leaves are likely to intercept more light per unit area than L. perenne (Riveros, 1963) and also are more efficient than D. glomerata (Remison, 1976).

Yorkshire Fog grass can grow in most soil types, from heavy loams to sands (Hubbards, 1945). Its optimum soil pH is 5.0 to 7.5 (Davies, 1944; Watt, 1977; Kruijne and de Vries, 1963). However, it also becomes prevalent in acidic soil (Davies, 1944; Hart and McGuire, 1963). It requires a moderate to low fertility. At low nitrogen level, it has yielded equally to L. perenne under cutting regimes (Haggars, 1976; Hayes, 1976; Haggars and Standell, 1982). The application of phosphorus did not change the amount of $H$. lanatus presence in a mixed sward in Oregon (Hart and McGuire, 1963). It tends to perform best on soil low in potassium, as noted in a survey in the Netherlands (Kruije and de Vries, 1963) and in United Kingdom (Castle and Holmes, 1960). The capability to grow in such poor nutrient conditions has been ascribed to various properties. One of these is its cation exchange capacity of the root systems, which provides it with an advantage over other grasses during a resource constraint (Jackman, 1960). Also, it has been noted that the root system absorbs nutrients in the surface layers of soil (Boggie et al., 1958; Beddows, 1961). Lastly, a symbiosis of endotrophic mycorrhiza in the root has been described (Hatch, 1937; Nye, 1966).

Its growth becomes prevalent where the soil moisture content is adequate. H.lanatus seems to tolerate wet soil conditions, commonly appearing in swamp, flooded or waterlogged areas (Basnyat, 1957 ; Morrison and Idles, 1972; Watt and Haggars, 1980), but it cannot tolerate a moderately dry or dry soil (Levy, 1970). The
flooding tolerant feature is possibly attributable to the anatomy of the root, which incorporates a radial cortex and many small irregular air spaces, thereby increasing the respiratory efficiency in low aeration (Soper, 1959; Jacques and Munro, 1963). Under such conditions, the plant also tends to produce more fine roots at soil surfaces and more adventitious roots around the edge of its clump (Watt, 1977).

Growth of Yorkshire Fog grass is centered on leaf expansion on a moderate number of large tillers (Munro, 1961). According to Protich (1977), formation of tillers in Holcus lanatus can be subdivided into the following four periods: (a) "onestem plant- formation period", when a plant is in the form of a covered bud from the time of development of first green leaf to the initiation of first of the lateral buds in the tillering zone; (b) "tillering period", when tillers of the second, third and fourth order are formed; (c) the "spring development and inflorescence period", when tillering ceases and the apical buds rapidly enter into the inflorescence period and the successive ontogenetic stages (d) "spring tillering period" when formation of inflorescences on the first, second and third tillers are completed and enlargement of internodes begins; new tillers of third and fourth orders and buds of the third, fourth and fifth order are formed.

Equivalent growth is yielded from 50 tillers of Yorkshire Fog grass or cocksfoot, 80 tillers of short rotation ryegrass, 100 tillers of perennial ryegrass, or 350 tillers of browntop, at temperature $65^{\circ} \mathrm{F}$ (Munro, 1961). Tiller number and shoot dryweight in $H$. lanatus grown at $7-35^{\circ} \mathrm{C}$. followed a course similar to that in $L$. perenne and D. glomerata (Mitchell and Lucanus, 1962). However, H.lanatus can give greater yield of shoot dry weight in early spring than does L.perenne (cv. S23) (Haggar, 1976). This is possibly due to its early growth at low temperature (Watt, 1983). Comparison among weed grasses, i.e. rough stalked meadow grass, Agrotis spp. and H. lanatus with ryegrass, they were lower yielding than the best ryegrass line. However, there was one exceptional population of Yorkshire Fog (BS 3639) which showed higher mass than ryegrass (Twigg, 1978).

Yorkshire Fog grass is useful in infertile, unstable, poorly drained soil (Munro, 1961; Davies et al., 1971; Morrison and Idle, 1972; Rumball, 1983). It is capable of establishing well in humid hill county, and on unploughable steep hills
(Basnsyat, 1957; Hughes and Nicholson, 1961). Despite some of its usefulness, several drawbacks have limited its generalized utilization in pasture production. These include the low palatiblity commonly attributed to excessive flower heads, basal dead matters', rust infestation, hairiness (Munro, 1961; Rumball, 1983). However, Cameron (1979) had pointed out that hairiness was considered an unimportant factor determining sheep preference. It is very susceptible to damage by tramping and treading (Brown and Evans, 1973; Watt, 1977). H. lanatus also restricted the establishment of sown T. repens more than did L. perenne (Jacques, 1974; Smith and Allcolk, 1985), and the clover transplants grew twice as much in ryegrass swards as in Yorkshire Fog swards (Turkington et al., 1979). This is possibly due to either its greater shading (Jaques, 1974), the allelopathic effects from its root leachates towards its neighbouring plants (Newman and Rovira, 1975), or its aggressive root competition (Remison, 1976).

The onset of numerous flower heads have caused a rapid decline in acceptability (Cowlishaw \& Alder, 1960; Garner, 1963; Jacques, 1974). The density of inflorescences was one of the most important factors determining lack of sheep acceptability (Cameron, 1979).

### 1.1.2 Plant Breeding

To improve the grass, Massey Agricultural college initiated its improvement project in 1953 with collection of 151 seed samples from most districts of New Zealand (Basnyat, 1957). Spaced plants underwent evaluation for two years combined with selection to improve utilization and palatability. The criteria used were: habit of growth, the extent of leaf pubescence, the propagation of dead basal tissue, resistance to crown rust, competitivity with legumes in the sward (Jaques, 1962; Munro, 1961).

A group of promising plants were selected for progeny testing by the polycross techniques in 1959-1960 resulting in selection of 10 lines showing high general combining ability in term of maintained production, adaptability to three different soil type, limited heading and rust resistance (Basnyat, 1957; Munro, 1961). The performance of elite line was tested against ryegrass showing that its winter yield
sustained vigour throughout the year, and a high tolerance to crown rust (Munro, 1961). The cultivar was released as "Massey Basyn " in 1977 (Rumball, 1983)

Massey Basyn performance was evaluated in several temperate countries. At Glen Innes, Australia, comparison with P. aquatica cv.Sirosa, cv.Commercial and Festuca arundina cv.Demeter under mixed sward with white clover, showed that mean pasture availability was greatest initially on Massey Basyn but finally on Commercial Phalaris (Robinson, May and Scarsbrick, 1980). It established and grew well by direct drilling following burning of native grassland in the Falkland Islands (McAdam, 1984). In the uplands of Britain, Massey Basyn with 130 kg .N/ha showed similar dry matter yields to that of L. perenne (Smith and Allcock, 1985). However, L. perenne responded better than L. lanatus to high levels of nitrogen fertilizer ( $200-250 \mathrm{~kg} . \mathrm{N} / \mathrm{h}$ annually) (Watt, 1984). Similar results was affirmed at the Oxford University Field Station and additionally indicated that Massey Basyn and German Commercial had no difference in terms of yield but Massey Basyn was affected less by rust infection (Watt, 1987).

### 1.1.3 Germplasm Variability

An outcrossing species Yorkshire Fog grass may be subjected to a wide range of adaptive pressures. Its large phenotypic variability in New Zealand has been described as a secondary centre of diversity for the species (Munro, 1961; Jacques, 1962; 1974). A cluster analysis study of the phenotypic variability in several characters was conducted by Teow (1978). Based on Ward's clustering method, the 161 local populations (topodemes) were grouped into five distinct clusters.

### 1.1.4 Phenotypic and Genotypic Variability

Phenotypic variation of some characters (related to sheep acceptability) was estimated by Cameron (1979). The investigation was based on topodeme level. It is also notable that a high degree of plant variation within the topodeme prevails (the residuals of the previous two studies).

Besides the topodeme variability just discussed, several workers have made observation on specific characters in Holcus lanatus.

Phenotypic variation in leaf pubescence, in terms of hair density and hair length, is apparent. The inheritance of this character was believed to be quantitative by Beddows (1961). The genetic variation relative to phenotypic variation was low (0.2) (Cameron, 1979).

Plant form is variable in Yorkshire Fog grass. Commonly, Yorkshire Fog grass plants have an extremely prostrate growth habit (Jacques, 1974). However, it tends to grow in clumps in established swards (Beddows, 1961; Hubbard, 1968; Turkington and Harper, 1979). Its growth habit can be due to the formation of decumbent tillers in the late summer which subsequently produce roots and shoots at the nodes (Watt, 1983) Conversely, predominantly erect and semi-erect plants were available in the early selection program (Munro, 1961). Clump erectness was found to be one of most discriminating characters among groups in clustering analysis (Teow, 1978). However, the genetic variation relative to phenotypic variation was very low (0.1) (Cameron, 1979).

The major disease is crown rust (Puccinia coronata var.holci) which commonly infests old leaves during summer (Corkill, 1956; Jacques \& Munro, 1963). The phenotypic variation on disease appearance was high both among and within population (Munro, 1961). The genetic variation relative to phenotypic variation was low (0.1-0.3) (Cameron, 1979).

Panicle variation is observable. Panicle shapes are varied from lanceolate to oblong or ovate, very dense to rather loose, erect and nodding, whitish, pale green, pinkish or with a tinge of purple. The panicle size ranges from 3 to 20 cm .(Hubbard, 1968).

Yorkshire Fog grass tends to develop its maximum number of panicles during summer (October - November) in New Zealand. Flowering duration is about 3 months and varies widely over the groups of plants (Basnyat, 1957). However, time of flowering is also influenced by micrograzing pressure, soil moisture, exposure and the recurrence of annual period of moisture stress (McMillan, 1959; Cooper, 1954). The flowering date was also one of the most discriminating characters amongst groups in
the clustering study (Teow, 1978). The genotypic variation relative to phenotypic variation of flowering day was medium (0.3) (Cameron, 1979).

Yorkshire Fog grass can attain the height of $20-100 \mathrm{~cm}$.(Hubbard, 1968). The genetic variation relative to phenotypic variation in clump height was very low (0.004-0.03) (Cameron, 1979).

### 1.1.5 Heritability

Until recently, the relative contribution of genetics and environments to this variability were estimated. The heritability estimates were presented by Cameron (1979), using the split-plot-in-time model. These estimates on some of botanic and flowering characters are shown in Table 1.1. These estimates are for topodeme differences, not plant variation.

Heritability estimates based on plant to plant variation were studied recently on two adjacent populations in North Wales. Billington et al. (1988) revealed the heritability of several morphological and tillering characters (see Table 4.2). Two different quantitative genetic methods were employed in the study using maximumlikelihood technique. The populations were derived from fields with different management backgrounds. The improved field was also applied with fertilizer preceding the hay cut while the traditional field was not fertilized.

### 1.2 Quantitative Genetics

Quantitative genetics is the inheritance of those phenotypic characters between individuals that are continuously variable (quantitative) rather than due to simple segregating major gene system (qualitative) (Falconer, 1981) The same genetic principles underlie these attributes, but many genes are involved (polygenic) and the role of environment is much more pronounced. East (1910) was one of the early workers to demonstrate the relationship between classical genetics and quantitative variation. The procedures need some modified terminology and more biometrics than classical "segregating" genetic (Sprague, 1966).

Table 1.1 Broad-sense heritability estimates from split-plot-in-time model (Cameron,1979)

| Characters | Single harvest |  | Pooled harvest |  |
| :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{h}^{2}$ | se. | $\mathrm{h}^{2}$ | se. |
| Leaf tensile strength | 0.04 | (0.07) | 0.01 | (0.01) |
| Leaf pubescence | 0.20 | (0.08) | - |  |
| Leaf flavanols | 0.01 | (0.08) | - |  |
| Leaf width | 0.08 | (0.04) | - |  |
| Clump erectness | - |  | 0.10 | (0.05) |
| Clump height | - |  | 0.004 | (0.006) |
| Clump diameter | - |  | 0.06 | (0.03) |
| Clump rust | 0.10 | (0.08) | - |  |
| Green material | - |  | 0.02 | (0.02) |
| Flowering date | 0.34 | (0.09) |  |  |

Table 1.2 Heritability estimates from polycross data and the North Carolina model-2 experiment, both using REML (Billington, et al. 1988)
$\qquad$

Characters

|  | Polycross | North Carolina 2 |
| :---: | :---: | :---: |
| Impr Fld. | d. Trd Fld. | Impr Fld. Trd Fld. |


| Tiller number | 0.08 | -0.17 | 0.03 | - |
| :--- | ---: | :---: | ---: | ---: |
| Tiller dryweight (gm) | 0.19 | 0.19 | 0.01 | 0.24 |
| Stolon number | -0.29 | 0.28 | -0.10 | 0.17 |
| Stolon dryweight (gm) | -0.16 | 0.23 | -0.22 | 0.15 |
| Leaf width (mm) | -0.27 | -0.29 | 0.10 | 0.17 |
| Leaf length (mm) | 0.17 | - | - | - |
| Plant height (mm) | 0.18 | - | - | - |
| Plant diameter (mm) | -0.20 | 0.18 | - | - |
| Tiller number after cut | 0.22 | 0.19 | - | - |
| Flowering time (days) | 0.24 | 0.14 | 0.23 | 0.10 |
| Inflorescence number | 0.01 | 0.19 | 0.14 | 0.18 |
| Panicle length(mm) | 0.27 | 0.01 | - | - |
| Flag-leaf length (mm) | 0.04 | 0.11 | - | - |

Impr Fld. = Improved Field
Trd Fld. = Traditional Field

### 1.2.1 Partitioning Genetic Variance

The phenotypic value of a character for an individual can be partitioned into two main components that due to the genetic effect and that to the environmental effect (Mather and Jink, 1971; Falconer, 1981; Becker, 1984; Baker, 1986).

$$
P=G+E
$$

where: $P$ is the phenotypic value
G is the genotypic value
$E$ is the environmental effect

The genotypic value can be partitioned into three components, i.e.

$$
G=A+D+I
$$

where: A is the average allele effect ("additive")
$D$ is the heterozygote effect ("dominance")
I is the interaction between $A$ and $D$ ("epistasis")

The average effect is the sum of the "additive" (average) effects of alleles across all their backgrounds (Falconer, 1981).

The dominant effect or intra-locus effect is the sum, across loci, of heterozygote deviates within each locus (Falconer, 1981).

The epistatic effect or inter-locus effect or non-allelic effect, is the sum of main gene-effect inconsistencies among the loci (Falconer, 1981). It can be partitioned further into three parts, as follows:

$$
I=A A+A D+D D
$$

where: AA is the additive x additive interaction
AD is the additive x dominant interaction
DD is the dominant x dominant interaction

The environmental variance can also be partitioned according to the experimental model and assumptions (Cockerham, 1954). For example, in Randomized Complete Block design, the environmental variance is partitioned into the block variance and the residual (error) variance.

### 1.2.2 Genetic Experimental Designs

The experimental designs mostly employed to estimate genetical components are generations mean analysis and mating designs for variance component analysis (Spragues, 1966).

The basic generation mean model comprises $\mathrm{P}_{1}, \mathrm{P}_{2}, \mathrm{~F}_{1}, \mathrm{~F}_{2}, \mathrm{BC}$ to $\mathrm{P}_{1}\left(\mathrm{BC}_{1}\right)$, and BC to $\mathrm{P}_{2}\left(\mathrm{BC}_{2}\right)$ generation (Hayman, $1958 \mathrm{a} ; \mathrm{b}$ ). Other models have been developed to suit the nature of crop and decrease workloads. For example, model comprising $\mathrm{P}_{1}, \mathrm{P}_{2}, \mathrm{~F}_{2}, \mathrm{~F}_{3}, \mathrm{BC}_{1} \mathrm{~S}_{1}, \mathrm{BC}_{2} \mathrm{~S}_{1}$ generation is rather convenient for selfpollinated crop with a small amount of seed production (Hayman 1958b; Snape, 1987). The utilisation of generation mean analysis permits direct estimation of all epistatic parameters, but preparation of crosses usually limits the breadth of germplasm which can be studied.

The mating designs for variance component analysis are generally used much more than the former. The foundation of this procedure is due to Fisher (1918). The advancement in this area was developed by Wright (1921), Comstock and Robinson (1948) and Mather and Jink (1971), Hayman (1958a; b), Kempthorne (1957) ,Becker (1984) and Baker (1986).

Any models developed for the estimation of genetic variances involve a series of biological assumptions. The common ones are: normal diploid behaviour at meiosis; no maternal or cytoplasmic effects; no multiple alleles; linkage equilibrium; no selection; no epistasis.

Under some conditions, however, one or some of these assumptions can be exempted; but these may not be any needs to suppose relation of these assumptions, as they may be reasonable under population equilibria conditions.

The simplest mating designs are biparental mating design (BIP) and one-way mating design. The former involves crossing parents pairwise to produce full-sib family (Kearsey, 1965). And the latter involves crossing of one parent with an unknown parent to produce half-sib families (Becker, 1964). Both designs are confined to only two kinds of relationship among progenies, either sibling (full-sib / half-sib) or unrelated. However, under proper experimental design and appropriate assumptions, it can supply well-defined genetical variance components. An example of one-way mating design was showed in studying genetic components of morphological variation in Salix repens (Fowler et al., 1983).

Other designs utilize both half-sib and full-sib relationships. These are hierarchical design (North Carolina I) and factorial design (North Carolina II) (Comstock and Robinson, 1948). In the hierarchical design, each of a series of random males $(m)$ is mated to each of $f$ random females. The offsprings of the $m f$ matings comprise the relationship of half-sib ( Vm ) and full-sib - half-sib $(\mathrm{Vf}(\mathrm{m})$ ) and the unrelated $\left(\mathrm{V}_{\mathrm{e}}\right)$ (comstock and Robinson, 1948; 1952).

For the factorial design, each of a different series of males $(m)$ and females $(f)$ are mated to each other. The offsprings of $m f$ are related in the form of half-sib to males ( Vm ), half-sib to females ( Vf ), full-sib - both half-sibs, and the related ( $\mathrm{V}_{\mathrm{e}}$ ) (Comstock and Robinson 1948; 1952).

One of modifications of factorial designs which is popular and mostly applied in plant genetical analysis is diallel analysis (Cockerham, 1963; Kempthorne, 1957). The design involves the same series of males and females mating to one another. Due to its use of common parent group, the design can be modified further to several types (Griffings, 1956a; b).

1. Full diallel, offsprings derived from all full combinations of parents.
2. Partial diallels, offsprings derived from incomplete combinations which can be with or without parents and with or without reciprocal. They are used to overcome constraints from a large numbers of crosses. (Gilberts, 1958; Kempthorne and Curnow, 1961; Curnow, 1963; England, 1974).
3. Triallels (Rawlings and Cockerham, 1962a).
4. Partial triallels (Hinkelmann, 1965).
5. Tetra-allele cross designs (Rawlings and Cockerham, 1962b).

### 1.2.3 Heritability and Its Standard Error Estimates

Heritability is defined as proportion of genotypic variance to phenotypic variance (Falconer, 1981).

$$
h^{2}=v_{G} / v_{P}
$$

where: $\quad V_{G}$ is genotypic variannce
$V_{P}$ is phenotypic variance

One basic method to determine the heritability is the linear regression of genotypic values on phenotypic values (Baker, 1986). By definition;

$$
\mathrm{b}_{\mathrm{GP}}=\mathrm{v}_{\mathrm{GP} /} \mathrm{v}_{\mathrm{P}}
$$

where: $\quad \mathrm{V}_{\mathrm{GP}}$ is the covariance between genotypic and phenotypic value $\mathrm{V}_{\mathrm{P}}$ is the phenotypic variance

Since, $P=G+E$

$$
\mathrm{V}_{\mathrm{GP}}=\mathrm{V}_{(\mathrm{G})(\mathrm{G}+\mathrm{E})}=\mathrm{V}_{\mathrm{G}}+\mathrm{V}_{\mathrm{GE}}
$$

If G and E are independent, $\mathrm{V}_{\mathrm{GE}}=0, \mathrm{~V}_{\mathrm{GP}}=\mathrm{V}_{\mathrm{G}}$
Hence;

$$
\mathrm{b}_{\mathrm{GP}}=\mathrm{V}_{\mathrm{G} /} \mathrm{V}_{\mathrm{P}}
$$

Based on similar concept, parent-offspring relationship is also used to estimate the heritability. In this case, the phenotypic value of progeny $\left(\mathrm{P}_{\mathrm{i}}\right)$ is one-half maternal genetic value ( $\mathrm{G}_{\mathrm{i}}$ ), one-half paternal genetic value $\left(\mathrm{G}_{\mathrm{j}}\right)$ and an environmental deviation ( $\mathrm{E}_{\mathrm{j}}$ );

$$
\mathrm{P}_{\mathrm{i}}=0.5 \mathrm{G}_{\mathrm{i}}+0.5 \mathrm{G}_{\mathrm{j}}+\mathrm{E}_{\mathrm{i}}
$$

Under random mating situation, $\mathrm{G}_{\mathrm{i}}$ and $\mathrm{G}_{\mathrm{j}}$ will be uncorrelated. Hence;

$$
\begin{aligned}
& \mathrm{V}_{\mathrm{GP}}=\mathrm{V}_{\mathrm{Gi}(0.5 \mathrm{Gi}+0.5 \mathrm{Gj}+\mathrm{Ei})}=0.5 \mathrm{~V}_{\mathrm{G}} \\
& \mathrm{~h}^{2}=0.5 \mathrm{~V}_{\mathrm{G}} / \mathrm{V}_{\mathrm{P}}
\end{aligned}
$$

Furthermore, there is another viewpoint on heritability by considering the coefficient of determination of the regression of genotypic value on phenotypic value.

$$
\text { If } P_{i}=G_{i}+E_{i} \text { and }\left(G_{i}-\bar{G}\right)=b_{G P}\left(P_{i}-\bar{P}\right)
$$

The coefficient of determination for the regression of genotypic value is ;

$$
\mathrm{r}^{2}=\mathrm{V}_{\mathrm{GP}} / \mathrm{V}_{\mathrm{G}} \cdot \mathrm{~V}_{\mathrm{P}}=\mathrm{V}_{\mathrm{G}} / \mathrm{V}_{\mathrm{G}} \cdot \mathrm{~V}_{\mathrm{P}}=\mathrm{V}_{\mathrm{G}} / \mathrm{V}_{\mathrm{P}}=\mathrm{h}^{2}
$$

Heritability can be also estimated indirectly from differences between phenotypic and environmental variances or from the covariances between relatives. Partitioning genotypic variances into additive and non-additive portions can yield at least two common kinds of heritabilities. The broad-sense heritability considers total genetic variability in relation to the phenotypic variability ( $\mathrm{V}_{\mathrm{G}} / \mathrm{V}_{\mathrm{P}}$ ) while the narrowsense considers only the additive portion of the genetic variability in relation to phenotypic variation ( $\mathrm{V}_{\mathrm{A}} / \mathrm{V}_{\mathrm{P}}$ )(Hanson, 1963; Falconer, 1981). The proper application of these estimates in plant breeding exercise depends on mating practice. The former is appropriate for the inbred or clonal genotypes while latter is more appropriate in random mating population (Baker, 1986).

Its precision is indicated by its standard error (Falconer, 1981). A conventional way to derive the standard error of heritability is using the intra-class correlation coefficient (Robertson and Lerner, 1949). For a one-way mating design, Becker (1984) has described it as:

$$
\text { se. } h^{2}=\sqrt[4]{\frac{2(1-t)^{2}[1+(k-1) t]^{2}}{k(k-1)(s-1)}}
$$

where: $t$ is the intra-class correlation
k is the coefficient of variance component being estimated

In addition, standard error of heritability can also be derived from the variance of a ratio, using ratios of variance components (Osborne and Paterson, 1952) This procedure can be used with phenotypic and genotypic variances from any experimental models. Solutions for more complicated models were demonstrated by Gordon, et al. (1972) and Gordon (1979).

## CHAPTER 2

MATERIALS AND METHODS

### 2.1 Objectives

1. Partition genetic variance and estimate heritability.
2. Estimate the plant genetic variance and compare with topodeme variance.
3. Describe the species variation, identify those characters useful in selection and also develop guidelines for future plant breeding.
4. Elaborate tiller development and growth from the genetic point of view.

### 2.2 Source of Materials

Seeds of each line were collected from individual mother plants in an openpollination field. The offsprings of each plant therefore have one common parent (female) and many different male parents, making them half-sibs. Observations on an individual plant basis from these sibling groups make it possible to study the underlying genetic components. These lines will be called 'half-sib families' in this study.

Furthermore, the half-sib mother-plants were random individuals from several wild populations (topodemes) which previously had been grouped into clusters (Teow, 1978). This knowledge was used to define stratified samples, representing the phenotypic variation throughout New Zealand Yorkshire Fog grass. Stratified random sampling provided fifty half-sib families, five from each of ten topodemes, two of which came from each of the five clusters of Teow (see Fig.2.1). Comparison between the topodeme variation and half-sib family variation could therefore be done, in addition to the half-sib genetic analysis referred to earlier.

### 2.3 Experimental Design and Bench Layout

The experimental design was a grouped treatment Randomized Complete Block design. Nine individual plants from each half-sib family were used, arranged in three blocks, with three plants per experimental unit.


Figure 2.1 Origins of the 50 half-sib families from the 10 topodemes in 5 clusters defined by Teow (1978), the numbers refer to the seed catalogue

The experiment was set up in a glasshouse. Plants were placed in a fixed position across benches, without relocation. In this way, it was intended to used "position" as a concomitant variable in order to remove and quantify any position effect (e.g. from shading, etc.).( see Fig.2.2 and plate 2.1)

### 2.4 Experimental Crop Management

Seeds were sown in autumn (early April 1988). They were germinated in fluctuating temperature ( 8 hrs . in $10^{\circ} \mathrm{C}$ and 16 hrs .in $20^{\circ} \mathrm{C}$ ) and under continuous light conditions in a germinator. After 5 to 7 days, seedlings were removed to the glasshouse and transplanted into plastic planter bags ( 1.6 litres). The media used was sand and peat at the ratio of $3: 1$ with 250 g . of 3 -month Osmocote- $\mathrm{R}^{\mathrm{R}}$ for every 70 litres of mixed media.

At the early stages of vegetative growth, starting from the 4-5 leaf stage, plants were subjected to the ambient winter temperature of Palmerston North (heating unit was switched off) for almost 6 weeks ( 6 -th May to 17 - $^{\text {th }}$ June), in case vernalization was required. Previous studies and speculations indicated that low temperature in winter and long-day photoperiod may be a requirement for flower induction of Yorkshire Fog grass (Hill, 1988; Robertson, 1988 pers.comm.). Flowering induction and initiation were chiefly determined by a photoperiod more than 15.5 hours (Montaldo and Paredes, 1981) or between 1430 and 1845 hours (Prokudin; Kalenichenko; Mamro, 1983). Subsequently, plants were provided with artificial photosynthetic light to extend the active daylength to 16 hours a day starting from 0400 to 2000 hrs . Temperature in the glasshouse was controlled between $20-25^{\mathrm{O}} \mathrm{C}$. The aim was to provide a semblance of spring/ summer in the out-of-season glasshouse. The vernalizing treatments seemed to be effective, as the plants started their booting and heading on the first and second weeks of July.

Plants were watered by drip irrigation onto bench mats twice a day with each watering lasting about 30 minutes. Few aphids appeared, but were kept in check by pyrethroid chemical (rate $0.02 \%$ ) when necessary. Caging of individual plants with chicken-wire columns was practiced to hold up the plants because of the limited space in the glasshouse (Plate 2.3).


Door

Figure 2.2 Experimental layout in the glasshouse


Plate 2.1 Experimental layout at 4-th week (above) and at 7-th week (below).

### 2.5 Data Collection and Measurement

### 2.5.1 Leaf Blade Attributes

Seedling leaf blade width and seedling leaf blade length of the $2-$ nd and 3 -rd leaves from the ground level were measured (in millimetres) on the vegetative seedling (about one month from sowing). The leaf blade width was measured at the widest part of the leaf blade. Length was measured from the ligule to the tip. Most plants had 4 5 leaves at this stage.

Mature leaf blade width was measured (in millimetres) on the 3-th and the 4 th leaf blade from the top at two different growth stages. Firstly, at the stem elongation stage ( about 15-16 weeks after sowing), being the same time as leaf tensile strength, was measured; and secondly, at post-ripe-seed stage of the first tiller (about $30-35$ weeks and also being the end of the experiment). The latter measurement virtually coincided with the stem elongation stage of the secondary tillers. Three samples per plant were recorded in the first occasion, and only one sample per plant was recorded in the second measurement.

### 2.5.2 Tiller Numbers

Total tiller numbers of individual plant were counted every 7-10 days for two months, during vegetative stages from seedling to stem elongation (from $4^{\text {th }}$ wk. to $11^{\text {th }}$ wk. after sowing) (Plate 2.2).

At the end of experiment, tillers were classified into four groups namely: (1) dead tillers (post-flowering main tillers) (2) green tillers (secondary and tertiary tillers) (3) young tillers under 15 cm . tall and (4) aerial tiller (see Plate 2.3 and Fig 4.1 in Discussion).

After counting, each group of tillers was dried out in oven (at $75^{\circ} \mathrm{C}$ ) for 3 days and weighed separately giving tiller mass (in grams) for each group of tillers for each plant.


Plate 2.2 Stage of seedlings when the tiller counting started


Plate 2.3 Green tillers and aerial tillers

### 2.5.3 Leaf Sheath Purple Colour

Degrees of purple colour at the leaf sheath were scored three times : (1) at the vegetative stage of older plant just prior to stem elongation (about 12 wks . from sowing), (2) at stem elongation stage (about 14 wks . from sowing), and at stem elongation stage of the secondary tiller (about 33 wks . from sowing). Standard colour specimens were established, and an ordinal score from 1 to 5 was based on these scores (increasing with the increasing purple colour) (Plate 2.4). Increment of half-scores were used for border-line assessments.

### 2.5.4 Leaf Favanol and Tannin Content

The flavanoid precursors of condensed tannins in the leaf sheath were evaluated semi-quantitatively by Burn's spot test, based on the vanillin-hydrochloric acid method. The procedure, described extensively by Burn's (1963) and Jones et al. (1973) was relatively rapid and inexpensive. The test was carried out twice at the early stem elongation stage (firstly about 13 wks. and secondly about 15 wks. from sowing). An approximate 5 cm . piece of the outermost part of the leaf sheath was sampled from each plant. The sample was squeezed between two layers of Whatman ${ }^{R}$ No. 1 filter papers. The plant residual was discarded and its imprint on the paper was wetted with a few drops of test reagent. The reagent comprised two volumes of $10 \% \mathrm{w} / \mathrm{v}$ vanillin in ethanol mixed with one volume of concentrated hydrochloric acid. The reagent was normally kept on ice to keep it cool. The reaction paper was left for drying under ambient temperature ( $15^{\circ}-20^{\circ} \mathrm{C}$ ) inside a dark chamber for about 30-40 minutes. Development of a red to violet colour was scored against standards on a photograph (Plate 2.5). Ordinal scores of 1 to 5 (increasing with degree of red / violet) with half increments were based on these standards. The imprints with red and violet indicated the presence of flavan materials, while blue or green spots indicated lack of them.

### 2.5.5 Leaf Tensile Strength

Leaf tensile strength was tested during the middle-stem elongation stage (about $15-16 \mathrm{wks}$. of sowing) on the third and fourth leaf blade from the top. The

score $=1$

score $=3$


score $=2$

score $=4$
score $=5$

Plate 2.4 Leaf sheath colour score standard


Plate 2.5 Burn's spot test on flavanol standard
machine and technique were developed by Evans (1967 a; b). Three mature leaf blades were sampled from each plant during the morning. Water-soaked cotton wool was wrapped over the cut-end, and the leaves were put into a moist plastic bag until the testing period in the afternoon and evening. A 5 cm . piece was cut from about the middle of the lamina. This was inserted and held between two clamps. A motordriven spring applied load to a beam until the leaf specimens broke. A calibrated dial converted the breaking load into grams, using the regression equation of $\mathrm{Y}=-92.5+$ 5.5 $\mathrm{X}\left(\mathrm{R}^{2}=97.2 \%\right)$, where $\mathrm{Y}=$ estimate of breaking load (gms.), $\mathrm{X}=$ dial reading (Evans, 1964). The dry weight (mg.) of the tested specimens, (found after drying for 3 days at $70^{\circ} \mathrm{C}$ ) was also recorded after the break. The index of strength was estimated as:

$$
\text { Index of Strength }=\frac{\text { breaking load (gms.) }}{\text { dry weight (mg.) }}
$$

### 2.5.6 Leaf Hair

A mature leaf blade sampled at the stem elongation stage of the secondary tiller (about 30-35 wks. from sowing) was chosen randomly to examine the degree of hair intensity under a stereo-microscope. Ordinal scores 1 to 5 with a half increments were applied using the standard of Cameron(1979) (Plate 2.6).

### 2.5.7 Clump Erectness

Plant erectness scores were recorded at the older vegetative stage (about 7 wks. of sowing), this being prior to stem elongation for flowering. Ordinal scores of 1 to 5 with half increments were applied using the following definitions of angles from horizontals: (1) $0^{\circ}-15^{\circ}$; (2) $15^{\circ}-30^{\circ}$; (3) $30^{\circ}-45^{\circ}$; (4) $45^{\circ}-68^{\circ}$; (5) $68^{\circ}-90^{\circ}$. In allotting these scores, the general impression of the leaf-sheath angles of the plant were used.


Figure 2.3 Leaf hair standards for ordinal score (Cameron, 1979)

### 2.5.8 Flowering Day

Peeping day, the first anthesis day, the last flowering day were recorded (in number of days from sowing) on individual plant basis.

The peeping day was the first day when the terminal leaf-sheath showed a longitudinal split because of an enlarging inflorescence.

The first anthesis day was the first day when the first flower started to anthise.

And the last flowering day was the day when the last anthesis occurred.

The day lying half-way between the first anthesis day and the last flowering day was estimated also as the median flowering day.

### 2.5.9 Anthesis Time and Position

Anthesis time of day and anthesis position in the inflorescence on the first panicle have been recorded by ordinal scores, at the first anthesis day. For anthesis time, the scores of 1 to 4 were allocated for the time periods of $0400-0900 ; 0900-$ $1200 ; 1200-1400 ; 1400-1600$ hours, respectively. For anthesis position, the scores 1 to 3 were assigned to: top end portion, mid portion and bottom portion, respectively. Increments of half were used, also, for intermediate positions.

### 2.5.10 Panicle Size and Compactness

Panicle width and length were measured on a fully dehiscing inflorescence at the main anthesis stage (about 20 wks. from sowing). The degree of compactness was rated against ordinated standard specimens (Plate 2.7). The scores were 1 to 5 from dense to loose, with half increments.

score $=1$
score $=2$
score $=3$

score $=4$
score $=5$

Plate 2.6 Panicle compactness standard

### 2.5.11 Plant Height

Plant height was measured (in centimetres) from the soil level to top-end of panicle at late milk stage of the seed (about 25 wks . from sowing).

### 2.6 Statistical Analyses

### 2.6.1 Regression Analysis of Tiller Development

The functional relationship between tiller numbers and days was examined for each individual plant, using the "Sigmoid 2 Program" (Smith, unpubl.). The logistic function provided consistently the best fit (The other function examined was gompert). Best-fit was judged by high coefficient of determination, and by inspection of the fitted plots). Several estimates were obtained from the logistic fits namely : number of tillers at $5 \%, 50 \%, 95 \%$ of the upper asymptote, and at flowering time; also the number of days to attain $5 \%, 50 \%, 95 \%$ of upper asymptote of tiller number; the relative growth rate of tiller numbers at $5 \%, 50 \%, 95 \%$ of the upper asymptote. These calculations were assisted by an auxiliary program "Sigfits" (Smith, unpubl.). These estimates were used as data in ANOVA. These estimates provide data on firsttiller development, being estimated separately for each observational unit (plant).

### 2.6.2 Analysis of Variance

Due to some experimental units having one or two missing plants, the analysis of variance was carried out by generalized linear model procedure. The ANOVA was based on the following two models :

Model 1 (for Topodeme/Sib Families comparisons)

$$
X_{\mathrm{ijkl}}=\mu+T_{\mathrm{i}}+B_{\mathrm{j}}+T B_{\mathrm{ij}}+H_{\mathrm{k}(\mathrm{i})}+H B_{\mathrm{k}(\mathrm{i}) \mathrm{j}}+\varepsilon_{\mathrm{ijkl}}
$$

where: $X_{\mathrm{ijkl}}=$ the $i j k l-{ }^{\text {th }}$ phenotypic variate of individual plant.

$$
\begin{aligned}
& \mathrm{i}=1, \ldots \mathrm{t} \text { (no.of topodemes). } \\
& \mathrm{j}=1, \ldots \ldots \mathrm{~b} \text { (no.of blocks). } \\
& \mathrm{k}=1, \ldots \mathrm{~h} \text { (no.of half-sib families) } \\
& \mathrm{l}=1, \ldots \ldots . . \mathrm{p} \text { (no. of plants) }
\end{aligned}
$$

$\mu=$ the grand mean;
$T_{\mathrm{i}}=$ the $i$ - th topodeme effect,
$B_{\mathrm{j}}=$ the $j$ - - block effect;
$H_{\mathrm{k}(\mathrm{i})}=$ the $k$ - ${ }^{\text {th }}$ halfsib effect, nested within topodemes(error (a));
$T B_{\mathrm{ij}}=$ the interaction between topodeme and block effect;
$H B_{\mathrm{k}(\mathrm{i}) \mathrm{j}}=$ the interaction between half-sib and block
effect(error(b));
$\varepsilon_{\mathrm{ijkl}}=$ the residual variation associated with the ijkl- $^{\text {th }}$ plant.

This is a grouped treatment Randomized Complete Block design, which is analogous to a split-block design in its definition of error terms (Gomez and Gomez, 1984). Its main purpose was to compare the relative sizes of the three genotypic partitions: topodeme, half sib family and individual plant (Table 2.1).

Model 2 (for genetic analysis)

$$
X_{\mathrm{ijk}}=\mu+H_{\mathrm{i}}+B_{\mathrm{i}}+H B_{\mathrm{ij}}+\varepsilon_{\mathrm{ijk}}
$$

where: $\quad X_{\mathrm{ijk}}=$ the $i j k{ }^{-}$th phenotypic variate of individual plant; $\mathrm{i}=1, . . \mathrm{h}$ (no.of half-sib families); $j=1, \ldots . . . . . . b$ (no.of blocks); $\mathrm{k}=1$,.........p (no.of plants);
$\mu=$ the grand mean;
$H_{\mathrm{i}}=$ the $i$ - ${ }^{\text {th }}$ halfsib genotype effect;
$B_{\mathrm{j}}=$ the $j$ - th block effect;
$H B_{\mathrm{ij}}=$ the interaction between half-sib and block (experimental error);
$\varepsilon_{\mathrm{ijk}}=$ the residual variation associated with the $i j k-$ th plant.

This is an Randomized Complete Block design, with plant subsamples, intended to give a pooled genetic analysis (Table 2.2).

Both analyses of variance have been adjusted with the concomitant variable of plant bench-position to eliminate possible confounded effects due to plant position (such as shading, disease incidence, etc.). The plants of each half-sib were coded from one to three, starting from the outer edge towards the middle of the bench. These codes provided the concomitant variable.

F-tests for significance were constructed in the usual manner using randomeffect expectations of Mean Squares (Steel and Torrie, 1981; Crump, 1951; Satterthwaite, 1946).

Variance components for each effect were estimated together with their standard errors, using the program "Thwaite" (Gordon, unpubl.).

The estimator for the standard errors of the component 's estimates $\left(s^{2}\right)$ was

$$
\widehat{\operatorname{Var}}\left(\mathrm{s}^{2}\right)=2 \sum \mathrm{a}_{\mathrm{i}}^{2} \mathrm{M}_{\mathrm{i}}^{2} /\left(\mathrm{f}_{\mathrm{i}}+2\right)
$$

where: $\quad a_{i}$ 's are the linear mean-square coefficients used in computing $s^{2}$;
$M_{i}$ 's are the mean squares used in estimating $s^{2}$;
$\mathrm{f}_{\mathrm{i}}$ 's are the degrees of freedom of those mean-squares.
(Anderson and Bancroft, 1952 ; Crump, 1951).

Program "Thwaite" (Gordon, unpubl.) was used to effect these estimates.

### 2.6.3 Estimation of Genetic Variance

The biometrical variance estimates and the genetic variances were interrelated via the intra-class correlation (Falconer, 1981). The present experiment represents a one-way mating design (Falconer, 1981), and relates the model-2 experimental (biometrical) variances to the covariance between individuals within a progeny group (Baker, 1984; Falconer, 1981). As these progeny individuals were half-sibs, the

Table 2.1 Expected Mean Squares (EMS) (Model 1)

## Source

EMS

Block
Topodeme
Topodeme $\times$ Block

$$
\begin{array}{ll}
V_{w}+\tilde{p} V_{H B}+\tilde{p} h V_{T B}+\tilde{p} h T_{V B} & M S 6 \\
V_{W}+\tilde{p} V_{H B}+\tilde{p} h V_{T B}+\tilde{p} h b V_{T} & M S 5 \\
V_{W}+\tilde{p} V_{H B}+\tilde{p} V_{H B}+\tilde{p} h V_{T B} & M S 4
\end{array}
$$

Half-sib(Topodeme)
$\begin{array}{lr}V_{w}+\tilde{p} V_{H B}+\tilde{p} b V_{H(T)} & M S 3 \\ V_{W}+\tilde{p} V_{H B} & M S 2 \\ V_{W} & M S 1\end{array}$

Table 2.2 Expected Mean Squares (EMS) (Model 2)

## Source

EMS

| Block | $V_{w}+\tilde{p} V_{H B}+\tilde{p} g V_{B}$ | $M S 4$ |
| :--- | :--- | :--- |
| Half-sib | $V_{w}+\tilde{p} V_{H B}+\tilde{p} b V_{H}$ | $M S 3$ |
| Block $x$ Half-sib | $V_{w}+\tilde{p} V_{H B}$ | $M S 2$ |
| Residual (Within Plot) | $V_{w}$ | $M S 1$ |

These are the balanced expectation ( $\hat{p}=c_{1} c_{2} c_{3}$ etc.)
variance amongst progeny group is equivalent to covariance between half-sib individuals.

Therefore,the genetical model in this one-way mating design is as follows :
$\mathrm{V}_{\mathrm{H}}=\operatorname{cov} .(\mathrm{HS})=1 / 4 \mathrm{~V}_{\mathrm{A}}+1 / 16 \mathrm{~V}_{\mathrm{AA}} \ldots \ldots . . .(1)$
$\mathrm{V}_{\mathrm{W}}=\mathrm{V}_{\mathrm{Tot} .} . \operatorname{cov} .(\mathrm{HS})=3 / 4 \mathrm{~V}_{\mathrm{A}}+\mathrm{V}_{\mathrm{D}}+15 / 16 \mathrm{~V}_{\mathrm{AA}}+\mathrm{V}_{\mathrm{AD}}+\mathrm{V}_{\mathrm{DD}}+\mathrm{V}_{\mathrm{e}} . .(2)$
(Baker, 1984)
where: $\quad \mathrm{V}_{\mathrm{A}}=$ "additive" variance (average allele effect variance);
$\mathrm{V}_{\mathrm{D}}=$ "dominance" variance (heterozygote variance);
$\mathrm{V}_{\mathrm{AA}}=$ "additive x additive" variance (epistatic inconsistencies across genes when additive effects are combined);
$\mathrm{V}_{\mathrm{AD}}=$ "additive x dominant" variance (epistatic inconsistencies from additive $x$ dominance combination);
$\mathrm{V}_{\mathrm{DD}}=$ "dominant x dominant" variance (epistatic inconsistencies from dominance $x$ dominance combination);
$\mathrm{V}_{\mathrm{e}}=$ "environmental" variance.

The model 2 plot residual $\left(\mathrm{V}_{\mathrm{BH}}=\mathrm{V}_{\mathrm{EXG}}\right)$ represents an "environmental" variance for experimental-units, each consisting of (notionally) three plants (the harmonic mean of actual plants per plot, after allowing for misses, was used in some characters). Therefore, on an individual plant basis,

$$
\begin{align*}
\mathrm{V}_{\mathrm{BH}} & =\mathrm{V}_{\overline{\mathrm{x}}}(\mathrm{plt}) \\
& =\mathrm{V}_{\mathrm{e}} / \mathrm{p} \\
\text { From which } \mathrm{V}_{\mathrm{e}} & =\mathrm{p} \mathrm{~V}_{\mathrm{BH}} \ldots . . . . . . . . . . . . . . . . .(3)  \tag{3}\\
& =\text { environmental variance for plant within plots }
\end{align*}
$$

This assumption of homogeneity of environmental variances has made it possible to remove the environmental confounding within $\mathrm{V}_{\mathrm{w}}$.

The links between the biometrical variance components and the genetic variance components were as followed:
(From 1)

$$
\mathrm{V}_{\mathrm{H}} \quad=(\mathrm{MS} 3-\mathrm{MS} 2) / \mathrm{pb}
$$

$$
=1 / 4 \mathrm{~V}_{\mathrm{A}}+1 / 16 \mathrm{~V}_{\mathrm{AA}}
$$

$$
\begin{equation*}
4 \mathrm{~V}_{\mathrm{H}}=\mathrm{V}_{\mathrm{A}}+1 / 4 \mathrm{~V}_{\mathrm{AA}} \tag{4}
\end{equation*}
$$

and $\quad 3 \mathrm{~V}_{\mathrm{H}} \quad=3 / 4 \mathrm{~V}_{\mathrm{A}}+3 / 16 \mathrm{~V}_{\mathrm{AA}}$

The phenotypic variance was defined as:

$$
\begin{align*}
\mathrm{V}_{\mathrm{p}}, \quad & =\mathrm{V}_{\mathrm{H}}+\mathrm{V}_{\mathrm{w}} \ldots \ldots . . . . . . . . . . . . . . . . . . . . . . . . . . .(6) ~ \\
& =\mathrm{V}_{\mathrm{H}}+\mathrm{V}_{\mathrm{g}}+\mathrm{V}_{\mathrm{e}} \\
& =\mathrm{V}_{\mathrm{G}}+\mathrm{V}_{\mathrm{e}} \\
& =\mathrm{V}_{\mathrm{A}}+\mathrm{V}_{\mathrm{AA}}+\mathrm{V}_{\mathrm{D}}+\mathrm{V}_{A D}+\mathrm{V}_{\mathrm{DD}}+\mathrm{V}_{\mathrm{e}} . \tag{7}
\end{align*}
$$

The within- family genetic variance is, using (2) and (3),

$$
\begin{equation*}
\mathrm{V}_{\mathrm{g}} \quad=\mathrm{V}_{\mathrm{w}}-\mathrm{V}_{\mathrm{e}} \tag{8}
\end{equation*}
$$

From $\mathrm{V}_{\mathrm{W}}$ and (5),

$$
\begin{equation*}
\mathrm{V}_{\mathrm{w}}-3 \mathrm{~V}_{\mathrm{H}}=\mathrm{V}_{\mathrm{D}}+3 / 4 \mathrm{~V}_{\mathrm{AA}}+\mathrm{V}_{\mathrm{AD}}+\mathrm{V}_{\mathrm{DD}}+\mathrm{Ve}^{2} \tag{9}
\end{equation*}
$$

and (9)-(3),

$$
\begin{equation*}
\mathrm{V}_{\mathrm{h}} \quad=\mathrm{V}_{\mathrm{D}}+3 / 4 \mathrm{~V}_{\mathrm{AA}}+\mathrm{V}_{\mathrm{AD}}+\mathrm{V}_{\mathrm{DD}} \tag{10}
\end{equation*}
$$

### 2.6.4 Heritability

Various heritability estimates were made, following standard principles (Falconer, 1981; Comstock, 1952). The definitions used were as follows.

| $\mathrm{h}^{2}$ (broad sense) $\quad$ | $=$ heritability of all gene effects (genotype); |
| ---: | :--- |
|  | $=\left(V_{H}+V_{g}\right) /\left(V_{H}+V_{W}\right)$ |
|  | $=\left(V_{A}+V_{D}+V_{A A}+V_{A D}+V_{D D}\right) / V_{P}$ |

$\mathrm{h}^{2}$ (narrow sense) $\quad=$ heritability of average allele effects;

$$
=4 \mathrm{~V}_{\mathrm{H}} /\left(\mathrm{V}_{\mathrm{H}}+\mathrm{V}_{\mathrm{w}}\right)
$$

$$
=\left(\mathrm{V}_{\mathrm{A}}+1 / 4 \mathrm{~V}_{\mathrm{AA}}\right) / \mathrm{V}_{\mathrm{P}}
$$

$h_{2}$ (heterotic sense) $=$ heritability of non-average allele effects;

$$
\begin{aligned}
& =\left(\mathrm{V}_{\mathrm{w}}-3 \mathrm{~V}_{\mathrm{H}}\right) /\left(\mathrm{V}_{\mathrm{H}^{+}} \mathrm{V}_{\mathrm{W}}\right) \\
& =\left(\mathrm{V}_{\mathrm{D}}+3 / 4 \mathrm{~V}_{\mathrm{AA}}+\mathrm{V}_{\mathrm{AD}}+\mathrm{V}_{\mathrm{DD}}\right) / \mathrm{V}_{\mathrm{P}}
\end{aligned}
$$

## CHAPTER 3

## RESULTS

There were forty seven characters under investigation in this experiment. They were divided into three main categories: botanical characters, agronomic characters and tillering characters. The model 1 analysis (see methods) was used to compare the magnitude in variation between topodemes and half-sib families. The model 2 analysis was the basis for the plant genetic analyses, from which the heritability estimates were obtained.

The general value of each attribute is indicated by the grand means given in the Table 3.1. This table also summarized the overall variability in two ways: the coefficient of variation and the range (minimum and maximum). Several attributes have a high level of the coefficient of variation. These include 12 wks . and 15 wks . leaf sheath purple, anthesis time, anthesis position, panicle width and compactness, clump erectness, old disease and new disease, flavanoid at leaf sheath and almost all attributes of tillering except for the number of days to tillering. Mean differences among the fifty half-sib families accounting for each attribute are exhibited in Appendix I.

### 3.1 Topodeme, Half-sib and Plant Variance Analysis

The environmental variance, including block variance $\left(V_{B}\right)$, error (a) variance or topodeme by block interaction ( $\mathrm{V}_{\mathrm{TB}}$ ) and error (b) variance or half-sib by block interaction $\left(\mathrm{V}_{\mathrm{HB}}\right)$ is shown in Table 3.2. Most attributes was significantly influenced to some degrees by the environmental effects. The attributes which show significance on those three environmental effects simultaneously include 15 wks . leaf sheath purple, flowering peeping day, first anthesis day, median flowering day, clump erectness, flavanoid at leaf sheath, tillering number at $5 \%$ tillering, numbers of dead tillers, and numbers of days for $50 \%, 95 \%$ tillering.

Table 3.1 The grand means, their coefficients of variation and maxima and minima over all half-sib families
$\qquad$

## Botanic characters

| 1. Juvenile leaf width | mm | 25.51 | 14.12 | 21.9 | 32.1 |
| :--- | :--- | ---: | ---: | ---: | ---: |
| 2. Juvenile leaf length | mm | 72.83 | 15.94 | 59.4 | 92.9 |
| 3. Mature leaf width (15 wks) | mm | 10.31 | 11.13 | 9.1 | 11.5 |
| 4. Mature leaf width (33 wks) | mm | 9.49 | 11.75 | 8.5 | 10.7 |
| 5. Leaf sheath purple (12 wks) | score | 6.20 | 22.53 | 4.2 | 7.8 |
| 6. Leaf sheath purple (15 wks) | score | 4.32 | 35.14 | 3.0 | 6.7 |
| 7. Leaf sheath purple (33 wks) | score | 9.07 | 15.19 | 7.4 | 10.0 |
| 8. Plant height | cm | 107.82 | 9.47 | 97.6 | 121.4 |

## Flowering characters

9. Flower peeping day
10. First anthesis day
11. Median flowering day
12. Last flowering day
13. Anthesis time
14. Anthesis position
15. Panicle width
16. Panicle length
17. Panicle compactness

| days | 112.83 | 3.53 | 108.44 | 122.00 |
| :--- | ---: | ---: | ---: | ---: |
| days | 120.58 | 3.55 | 115.22 | 129.60 |
| days | 134.16 | 7.07 | 128.21 | 145.31 |
| days | 147.75 | 12.40 | 135.71 | 169.63 |
| score | 2.18 | 100.86 | 1.00 | 4.00 |
| score | 3.78 | 33.88 | 2.78 | 4.89 |
| cm. | 55.75 | 31.79 | 37.68 | 80.68 |
| cm. | 134.03 | 18.96 | 107.79 | 157.42 |
| score | 5.36 | 34.86 | 3.75 | 6.80 |

## Agronomic characters

18. Clump erectness
19. Old diseases
20. New diseases
21. Leaf hairiness
22. Leaf tensile strength
23. Flavanoid at leaf sheath

| score | 4.33 | 35.15 | 2.1 | 5.9 |
| :--- | ---: | ---: | ---: | :---: |
| score | 7.12 | 22.87 | 5.1 | 8.5 |
| score | 4.21 | 40.89 | 1.9 | 6.6 |
| score | 8.76 | 12.10 | 7.6 | 9.6 |
| mm. | 95.70 | 15.13 | 80.5 | 115.7 |
| score | 4.62 | 37.07 | 2.7 | 21.02 |

Table 3.1 (continued)
$\qquad$

## Tillering characters

| 24. Tiller No.at 5\% tillering | no. | 2.34 | 99.72 | 1.40 | 10.18 |
| :--- | :--- | ---: | ---: | ---: | ---: |
| 25. Tiller No.at 50\% tillering | no. | 21.44 | 63.58 | 13.91 | 48.81 |
| 26. Tiller No.at 95\% tillering | no. | 41.46 | 67.33 | 28.03 | 102.56 |
| 27. Tiller No.at flowering time | no. | 41.86 | 34.91 | 29.88 | 62.75 |
| 28. No.of dead tillers at end | no. | 18.08 | 46.00 | 8.67 | 30.56 |
| 29. No.of green tillers at end | no. | 28.18 | 53.83 | 18.00 | 41.25 |
| 30. No.of young tillers at end | no. | 11.37 | 86.59 | 2.44 | 26.83 |
| 31. No.of aerial tillers at end | no. | 73.77 | 47.24 | 49.89 | 108.11 |
| 32. No.of total tillers at end | no. | 132.65 | 33.62 | 101.89 | 117.14 |
| 33. No.of base tillers at end | no. | 57.67 | 40.31 | 34.67 | 91.22 |
| 34. No.of base green tiller | no. | 39.53 | 50.78 | 21.33 | 64.86 |
| 35. Dead tiller dry weight | gm. | 8.83 | 64.80 | 2.80 | 15.06 |
| 36. Green tiller dryweight | gm. | 19.43 | 66.94 | 11.66 | 35.67 |
| 37. Young tiller dryweight | gm. | 1.33 | 106.22 | 0.24 | 3.62 |
| 38. Aerial tiller dryweight | gm. | 27.92 | 53.07 | 15.84 | 45.37 |
| 39. Total tiller dryweight | gm. | 56.81 | 44.99 | 38.10 | 78.24 |
| 40. Base tiller dryweight | gm. | 55.94 | 44.31 | 17.93 | 46.51 |
| 41. Base green tiller dry weight | gm. | 20.82 | 64.76 | 12.61 | 35.91 |
| 42. No.of days for 5\% tillering | days | 28.98 | 14.49 | 24.16 | 34.88 |
| 43. No.of days for 50\% tillering | days | 63.63 | 11.78 | 54.39 | 77.56 |
| 44. No.of days for 95\% tillering | days | 98.49 | 12.74 | 83.85 | 120.63 |
| 45. RGR at 5\% tillering | - | 35.94 | 162.34 | -21.68 | 246.12 |
| 46. RGR at 50\% tillering | - | 87.34 | 74.67 | 69.13 | 284.41 |
| 47. RGR at 95\% tillering | - | 91.10 | 78.58 | 70.58 | 286.40 |

[^0]Table 3.2 Block, Error(a), Error(b) variance components and their standard error and F-significance, together with position F-significance (model 1)

| Characters | Block |  | Error(a) |  | Error(b) |  | Position |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Var. <br> (se.) | F-sig. | Var. <br> (se.) | F-sig. | Var. <br> (se.) | F-sig. | Fsig. |

## Botanic characters

| 1. Juvenile leaf width | $\begin{gathered} 0.07 \\ (0.12) \end{gathered}$ | ns | $\begin{gathered} 0.02 \\ (0.33) \end{gathered}$ | ns | $\begin{gathered} 0.26 \\ (0.81) \end{gathered}$ | ns | ns |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2. Juvenile | 0.26 | ns | 2.48 | ns | -4.76 | ns | ns |
| leaf length | (0.79) |  | (3.57) |  | (7.32) |  |  |
| 3. Mature leaf | -0.004 | ns | 0.02 | ns | -0.06 | ns | ns |
| width ( 15 wks ) | (0.004) |  | (0.03) |  | (0.08) |  |  |
| 4. Mature leaf | 0.35 | ** | -0.01 | ns | 0.16 | * | ns |
| width (33 wks) | (0.26) |  | (0.04) |  | (0.10) |  |  |
| 5. Leaf sheath | 0.05 | ** | 0.04 | * | -0.15 | ns | ns |
| purple(12 wks) | (0.05) |  | (0.05) |  | (0.10) |  |  |
| 6. Leaf sheath | 0.05 | ** | 0.08 | ** | 0.45 | ** | ** |
| purple(15 wks) | (0.05) |  | (0.11) |  | (0.20) |  |  |
| 7. Leaf sheath | 0.29 | ** | -0.04 | ns | 0.35 | ** | ns |
| purple(33 wks) | (0.22) |  | (0.06) |  | (0.17) |  |  |
| 8. Plant height | 9.35 | ** | 10.92 | ** | 11.32 | ns | ns |
|  | (7.47) |  | (6.72) |  | (8.27) |  |  |

Flower characters

| 9. Flower peeping day | $\begin{gathered} 1.26 \\ (1.02) \end{gathered}$ | ** | $\begin{gathered} 1.72 \\ (1.06) \end{gathered}$ | ** | $\begin{gathered} 1.93 \\ (1.29) \end{gathered}$ | * | ns |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| First anthesis day | 1.32 | ** | 2.40 | ** | 4.48 | ** | ns |
|  | (1.13) |  | (1.50) |  | (1.82) |  |  |
| 11. Median flowering | 3.49 | ** | 6.08 | ** | 4.04 | ** | * |
| day | (3.06) |  | (4.38) |  | (6.33) |  |  |
| 12. Last flowering day | 5.63 | * | 19.14 | * | 11.36 | ns | ** |
| 13. Anthesis time | -0.03 | ns | -0.17 | ns | 0.36 | ns | ns |
|  | (0.01) |  | (0.10) |  | (0.36) |  |  |
| 14. Anthesis position | 0.03 | * | 0.004 | ns | 0.09 | ns | ns |
|  | (0.03) |  | (0.05) |  | (0.12) |  |  |
| 15. Panicle width | 1.80 | ns | -2.43 | ns | 68.61 | ** | ** |
|  | (3.78) |  | (12.15) |  | (30.05) |  |  |
| 16. Panicle length | 83.17 | ** | 14.92 | ns | -17.47 | ns | * |
|  | (62.12) |  | (19.67) |  | (39.49) |  |  |
| 17. Panicle | -0.03 | ns | -0.06 | ns | 0.73 | ** | ns |
| compactness | (0.01) |  | (0.12) |  | (0.33) |  |  |

Table 3.2 (continued)

| Characters | Block |  | Error(a) |  | Error(b) |  | Position |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Var. <br> (se.) | F-sig. | Var. <br> (se.) | F-sig. | Var. <br> (se.) | F-sig. | Fsig. |

## Agronomic characters

| 18. | Clump erectness | $\begin{gathered} 0.19 \\ (0.15) \end{gathered}$ | ** | $\begin{gathered} 0.24 \\ (0.15) \end{gathered}$ | ** | $\begin{gathered} 0.29 \\ (0.18) \end{gathered}$ | * | ** |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 19. | Old diseases | -0.02 | ns | 0.21 | ** | 0.38 | * | ns |
|  |  | (0.01) |  | (0.16) |  | (0.22) |  |  |
|  | New diseases | 0.38 | ns | 0.33 | ** | 0.54 | ** | ns |
|  |  | (0.29) |  | (0.21) |  | (0.26) |  |  |
| 21. | Leaf hairiness | 8.11 | ** | -0.04 | ns | 0.20 | ** | ns |
|  |  | (0.08) |  | (0.03) |  | (0.10) |  |  |
| 22. | Leaf tensile strength | 34.52 | ** | -6.44 | ns | 50.20 | ** | ns |
|  |  | (26.11) |  | (7.22) |  | (21.03) |  |  |
| 23. | Flavanoid at | 0.25 | ** | 0.05 | ** | 0.72 | ** | ** |
|  | leaf sheath | (0.20) |  | (0.14) |  | (0.28) |  |  |

Tillering characters

| 24. | Tiller No.at 5\% tillering | $\begin{aligned} & -0.03 \\ & (0.06) \end{aligned}$ | ns | $\begin{gathered} 0.15 \\ (0.04) \end{gathered}$ | ** | $\begin{gathered} 3.13 \\ (0.81) \end{gathered}$ | ** | ** |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 25. | Tiller No.at $50 \%$ tillering | $\begin{gathered} 4.0 \\ (3.88) \end{gathered}$ | * | $\begin{aligned} & -1.33 \\ & (4.90) \end{aligned}$ | ns | $\begin{gathered} 9.54 \\ (12.89) \end{gathered}$ | ns | ** |
| 26. | Tiller No.at $95 \%$ tillering | $\begin{gathered} 14.88 \\ (14.92) \end{gathered}$ | * | $\begin{aligned} & -13.28 \\ & (18.85) \end{aligned}$ | ns | $\begin{gathered} 46.00 \\ (54.90) \end{gathered}$ | ns | ** |
| 27. | Tiller No.at flowering time | $\begin{gathered} 0.81 \\ (1.92) \end{gathered}$ | ns | $\begin{gathered} 3.84 \\ (7.46) \end{gathered}$ | ns | $\begin{gathered} 15.05 \\ (15.40) \end{gathered}$ | ns | ** |
| 28. | No.of dead tillers at end | $\begin{gathered} 1.39 \\ (1.53) \end{gathered}$ | ** | $\begin{gathered} 2.95 \\ (3.36) \end{gathered}$ | ** | $\begin{gathered} 3.56 \\ (1.97) \end{gathered}$ | * | ** |
| 29. | No.of green tillers at end | $\begin{array}{r} 33.89 \\ (25.50) \end{array}$ | ** | $\begin{gathered} 31.90 \\ (15.33) \end{gathered}$ | ** | $\begin{gathered} 1.93 \\ (14.24) \end{gathered}$ | ns | ** |
| 30. | No.of young tillers at end | $\begin{aligned} & 10.48 \\ & (8.06) \end{aligned}$ | ** | $\begin{gathered} 5.27 \\ (4.44) \end{gathered}$ | ** | $\begin{aligned} & -5.19 \\ & (5.17) \end{aligned}$ | ns | ** |
| 31. | No.of aerial tillers at end | $\begin{gathered} 60.59 \\ (48.09) \end{gathered}$ | ** | $\begin{gathered} 46.21 \\ (36.90) \end{gathered}$ | ns | $\begin{aligned} & -83.72 \\ & (61.72) \end{aligned}$ | ns | ** |
| 32. | No.of total tillers at end | $\begin{gathered} 76.61 \\ (65.26) \end{gathered}$ | ** | $\begin{aligned} & 156.72 \\ & (92.31) \end{aligned}$ | ** | $\begin{gathered} -51.91 \\ (115.96) \end{gathered}$ | ns | ** |
| 33. | No.of base tillers at end | $\begin{gathered} 1.81 \\ (5.23) \end{gathered}$ | ns | $\begin{gathered} 68.25 \\ (35.73) \end{gathered}$ | ** | $\begin{gathered} 22.08 \\ (37.14) \end{gathered}$ | ns | ** |
| 34. | No.of base green tiller at end | $\begin{gathered} 7.08 \\ (7.91) \end{gathered}$ | * | $\begin{gathered} 60.83 \\ (29.15) \end{gathered}$ | ** | $\begin{gathered} 7.83 \\ (26.49) \end{gathered}$ | ns | ** |
| 35. | Dead tiller dryweight at end | $\begin{gathered} 1.48 \\ (1.28) \end{gathered}$ | ** | $\begin{gathered} 4.08 \\ (2.12) \end{gathered}$ | ** | $\begin{gathered} 7.57 \\ (2.16) \end{gathered}$ | ns | ** |
| 36. | Green tiller dryweight at end | $\begin{gathered} 38.07 \\ (27.73) \end{gathered}$ | ** | $\begin{gathered} 4.38 \\ (5.12) \end{gathered}$ | ns | $\begin{aligned} & -3.94 \\ & (9.79) \end{aligned}$ | ns | ** |

Table 3.2 (continued)


## Tillering characters

| 37 | Young tiller dry weight | $\begin{gathered} 0.17 \\ (0.13) \end{gathered}$ | ** | $\begin{gathered} 0.06 \\ (0.08) \end{gathered}$ |  | $\begin{gathered} 0.17 \\ (0.15) \end{gathered}$ | ns | ** |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 38 | Aerial tiller dry weight | $\begin{aligned} & -0.85 \\ & (0.56) \end{aligned}$ | ns | $\begin{gathered} 6.90 \\ (7.01) \end{gathered}$ | ns | $\begin{gathered} -7.47 \\ (12.83) \end{gathered}$ | ns | ** |
| 39 | Total tiller dry weight | $\begin{gathered} 18.03 \\ (16.78) \end{gathered}$ | ** | $\begin{gathered} 22.71 \\ (20.21) \end{gathered}$ | ns | $\begin{aligned} & -40.80 \\ & (35.86) \end{aligned}$ | ns | ** |
| 40 | Base tiller dry weight | $\begin{gathered} 21.87 \\ (16.78) \end{gathered}$ | ** | $\begin{aligned} & 12.49 \\ & (9.52) \end{aligned}$ |  | $\begin{gathered} 1.50 \\ (14.64) \end{gathered}$ | ns | ** |
| 41 | Base green tiller dryweight | $\begin{gathered} 35.56 \\ (26.05) \end{gathered}$ | ** | $\begin{gathered} 6.28 \\ (6.04) \end{gathered}$ | ns | $\begin{gathered} -5.06 \\ (10.80) \end{gathered}$ | ns | ** |
| 42 | No.of days for $5 \%$ tillering | $\begin{gathered} 0.14 \\ (0.19) \end{gathered}$ | ns | $\begin{gathered} 0.29 \\ (0.50) \end{gathered}$ | ns | $\begin{gathered} -0.22 \\ (1.07) \end{gathered}$ | ns | ** |
| 43 | No.of days for $50 \%$ tillering | $\begin{gathered} 4.18 \\ (3.37) \end{gathered}$ | ** | $\begin{aligned} & 1.39 \\ & (2.34) \end{aligned}$ | * | $\begin{gathered} 7.81 \\ (4.60) \end{gathered}$ | * | ** |
| 44 | No.of days for $95 \%$ tillering | $\begin{gathered} 18.52 \\ (14.18) \end{gathered}$ | ** | $\begin{gathered} 3.45 \\ (6.14) \end{gathered}$ | * | $\begin{gathered} 17.69 \\ (12.30) \end{gathered}$ | * | ** |
| 45 | RGR at $5 \%$ tillering | $\begin{aligned} & -33.59 \\ & (22.50) \end{aligned}$ | ns | $\begin{gathered} -95.43 \\ (199.12) \end{gathered}$ | ** | $\begin{aligned} & 2005.46 \\ & (509.82) \end{aligned}$ | ** | ** |
| 46 | RGR at $50 \%$ tillering | $\begin{aligned} & -43.51 \\ & (17.15) \end{aligned}$ | ns | $\begin{gathered} 103.74 \\ (245.58) \end{gathered}$ | ** | $\begin{aligned} & 1565.67 \\ & (493.79) \end{aligned}$ | ** | ns |
| 47 | RGR at $95 \%$ tillering | $\begin{aligned} & -52.11 \\ & (15.74) \end{aligned}$ | ns | $\begin{gathered} 167.65 \\ (277.71) \end{gathered}$ | ** | $\begin{aligned} & 1450.78 \\ & (529.38) \end{aligned}$ | ** | ns |

[^1]The significance of the position effect is also shown in Table 3.2. Position effects were not significant in about half of the attributes, namely juvenile leaf width and leaf length, mature leaf width, leaf tensile strength, 12 wks . and 33 wks . leaf sheath purple, plant height, flower peeping day, first anthesis day, anthesis time, anthesis position, panicle compactness, old disease, new disease and leaf hairiness. Surprisingly, plant height was not affected by position in this study. It was noteworthy that nearly all tiller attributes were affected.

The half-sib family variance had more characters with significant variance than the topodeme component (in ratio of 5 to 4 ). This indicated that more variability existed at the half-sib family level. Comparison of the topodeme and half-sib variances and also within plot variance can be made directly in Table 3.3. The halfsib variances had a higher value than the topodeme variances in almost all the characters, except in 33 wks . mature leaf width, 15 wks . leaf sheath purple, flavanoid at leaf sheath and panicle compactness. In addition, the within plot variance has the higher value than the half sib variance in every characters. This indicated that variability level of plant to plant variation within half sib lines was also predominant.

### 3.2 Genotypic Variance Analysis and Heritability Estimation

From model 2 analysis, the overall genotypic variances (half-sib families or lines) are given in Table 3.4. The block and within-plot variances are the same as in the model 1. The biometrical variance was subsequently repartitioned into genetic variances. The plot variance $\left(\mathrm{V}_{\mathrm{HB}}\right)$ and within plot variance $\left(\mathrm{V}_{\mathrm{W}}\right)$ are also presented in the same table. Most of the half-sib variance were significant ( 39 out of 47) except in median and last flowering days, anthesis time and anthesis position, numbers of green tillers and total tillers. However, the plot variance showed less numbers of significant attributes ( 27 out of 47).

The genotypic variance was repartitioned into additive variance $\left(V_{A}\right)$ and heterotic variance $\left(\mathrm{V}_{\mathrm{h}}\right)$. The phenotypic was also obtained from the overall genotypic variance and environmental variance combination. These estimates are shown in Table 3.5. About half of total characters had higher value of additive variance than heterotic variance and vice versa.

Table 3.3 Topodeme, half-sib, within-plot variance component with their standard error and the F-significance (Model 1)

| Characters | Topodeme |  | Half-sib |  | Within-plot <br> Var. <br> (se.) |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Var. <br> (se.) | F-sig. | Var. <br> (se.) | F-sig. |  |

## Botanic characters

1. Juvenile leaf width
2. Juvenile leaf length
3. Mature leaf width (15 wks)
4. Mature leaf width (33 wks)
5. Leaf sheath purple (12 wks)
6. Leaf sheath purple (15 wks)
7. Leaf sheath purple (33 wks)
8. Plant height

## Flower characters

| 9. Flower peeping day | $\begin{gathered} 0.59 \\ (0.79) \end{gathered}$ | ** | $\begin{gathered} 0.84 \\ (0.84) \end{gathered}$ | ** | $\begin{gathered} 15.90 \\ (12.98) \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 10. First anthesis day | -0.30 | ** | 1.10 | ** | 18.36 |
|  | (0.72) |  | (1.20) |  | (14.99) |
| 11. Median flowering day | -2.52 | ns | -0.43 | ns | 90.00 |
|  | (1.63) |  | (3.18) |  | (73.48) |
| 12. Last flowering day | -7.00 | ns | -0.34 | ns | 335.78 |
|  | (5.94) |  | (11.76) |  | (28.74) |
| 13. Anthesis time | 0.03 | ns | -0.18 | ns | 4.83 |
|  | (0.05) |  | (0.16) |  | (3.94) |
| 14. Anthesis position | 0.01 | ns | -0.03 | ns | 1.64 |
|  | (0.03) |  | (0.06) |  | (1.34) |
| 15. Panicle width | 10.59 | ns | 19.05 | ** | 314.03 |
|  | (10.01) |  | (19.81) |  | (256.40) |
| 16. Panicle length | -9.92 | ns | 76.87 | ** | 645.54 |
|  | (7.38) |  | (34.49) |  | (527.08) |
| 17. Panicle compactness | 0.08 | ** | 0.06 | ** | 3.49 |
|  | (0.09) |  | (0.19) |  | (2.85) |

Table 3.3 (continued)

| Characters | Topodeme |  | Half-sib |  | Within-plot <br> Var. <br> (se.) |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Var. <br> (se.) | F-sig. | Var. <br> (se.) | F-sig. |  |

## Agronomic characters

| 18. Clump erectness | $\begin{gathered} 0.08 \\ (0.11) \end{gathered}$ | ** | $\begin{gathered} 0.35 \\ (0.16) \end{gathered}$ | ** | $\begin{array}{r} 2.13 \\ (0.19) \end{array}$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 19. Old diseases | 0.03 | ** | 0.05 | * | 2.65 |
|  | (0.09) |  | (0.13) |  | (0.22) |
| 20. New diseases | 0.30 | ** | -0.06 | ns | 2.96 |
|  | (0.23) |  | (0.13) |  | (2.42) |
| 21. Leaf hairiness | 0.02 | ns | 0.005 | * | 1.12 |
|  | (0.02) |  | (0.05) |  | (0.91) |
| 22. Leaf tensile strength | -3.33 | ns | 15.55 | ** | 209.46 |
|  | (2.39) |  | (14.30) |  | (18.34) |
| 23. Flavanoid at | 0.31 | ** | 0.09 | ** | 2.94 |
| leaf sheath | (0.19) |  | (0.17) |  | (2.40) |

Tillering characters
24. Tiller No.at $5 \%$
tillering
25. Tiller No.at $50 \%$ tillering
26. Tiller No.at $95 \%$ tillering
27. Tiller No.at flowering time
28. No.of dead tillers at end
29. No.of green tillers at end
30. No.of young tillers at end
31. No.of aerial tillers at end
32. No.of total tillers at end
33. No.of base tillers at end
34. No.of base green tillers at end

| -0.11 | $*$ | 0.12 | $* *$ | 5.46 |
| :---: | :--- | :---: | :--- | ---: |
| $(0.17)$ |  | $(0.47)$ |  | $(0.46)$ |
| 2.21 | ns | 14.08 | $* *$ | 185.81 |
| $(2.05)$ |  | $(9.36)$ |  | $(15.59)$ |
| 15.18 | ns | 42.26 | $* *$ | 779.09 |
| $(14.67)$ |  | $(36.46)$ |  | $(636.12)$ |
| 4.16 | $*$ | 39.73 | $* *$ | 213.65 |
| $(5.39)$ |  | $(15.92)$ |  | $(17.93)$ |
| 2.74 | $* *$ | 13.74 | $* *$ | 69.98 |
| $(2.80)$ |  | $(5.87)$ |  | $(5.91)$ |
| -8.20 | ns | -1.49 | ns | 230.13 |
| $(6.03)$ |  | $(6.86)$ |  | $(19.55)$ |
| 1.00 | $*$ | 11.90 | $* *$ | 96.91 |
| $(2.73)$ |  | $(5.96)$ |  | $(79.13)$ |
| 31.32 | $* *$ | 55.73 | ns | 1214.34 |
| $(31.58)$ |  | $(39.98)$ |  | $(100.33)$ |
| -19.62 | ns | 54.05 | ns | 1989.22 |
| $(43.99)$ |  | $(67.30)$ |  | $(1677.82)$ |
| -6.21 | $*$ | 33.99 | $* *$ | 50.38 |
| $(17.58)$ |  | $(25.51)$ |  | $(45.92)$ |
| -13.13 | ns | 17.31 | $*$ | 402.96 |
| $(12.18)$ |  | $(16.70)$ |  | $(329.02)$ |
| -0.58 | $*$ | 4.68 | $* *$ | 30.57 |
| $(0.98)$ |  | $(2.04)$ | $*$ | $(24.96)$ |
| 1.46 | ns | 11.64 | $*$ | 169.17 |
| $(3.22)$ |  | $(7.09)$ |  | $(138.13)$ |

Table 3.3 (continued)

| Characters | Topodeme |  | Half-sib |  | Within-plot <br> Var. <br> (se.) |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Var. <br> (se.) | F-sig. | Var. (se.) | F-sig. |  |

Tillering characters

| 37. | Young tiller dryweight at end | $\begin{aligned} & -0.02 \\ & (0.04) \end{aligned}$ | ns | $\begin{gathered} 0.28 \\ (0.13) \end{gathered}$ | ** | $\begin{gathered} 2.00 \\ (0.17) \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 38. | Aerial tiller dryweight at end | $\begin{aligned} & -0.08 \\ & (3.69) \end{aligned}$ | ns | $\begin{aligned} & 10.19 \\ & (8.23) \end{aligned}$ | ns | $\begin{gathered} 219.53 \\ (14.09) \end{gathered}$ |
| 39. | Total tiller dryweight at end | $\begin{aligned} & -9.17 \\ & (8.01) \end{aligned}$ | ns | $\begin{gathered} 15.17 \\ (19.93) \end{gathered}$ | ns | $\begin{aligned} & 653.09 \\ & (55.49) \end{aligned}$ |
| 40. | Base tiller dryweight at end | $\begin{aligned} & -5.52 \\ & (7.91) \end{aligned}$ | ns | $\begin{gathered} 17.24 \\ (10.66) \end{gathered}$ | ** | $\begin{aligned} & 228.33 \\ & (19.65) \end{aligned}$ |
| 41. | Base green tiller dryweight at end | $\begin{aligned} & 1.04 \\ & (3.59) \end{aligned}$ | ns | $\begin{aligned} & 11.92 \\ & (7.58) \end{aligned}$ | * | $\begin{aligned} & 181.84 \\ & (15.56) \end{aligned}$ |
| 42. | No.of days for $5 \%$ tillering | $\begin{gathered} 0.70 \\ (0.53) \end{gathered}$ | ** | $\begin{gathered} 2.46 \\ (1.02) \end{gathered}$ | ** | $\begin{gathered} 17.64 \\ (1.49) \end{gathered}$ |
| 43. | No.of days for $50 \%$ tillering | $\begin{gathered} 2.21 \\ (2.05) \end{gathered}$ | ** | $\begin{gathered} 4.22 \\ (3.26) \end{gathered}$ | ** | $\begin{gathered} 56.16 \\ (45.85) \end{gathered}$ |
| 44. | No.of days for $95 \%$ tillering | $\begin{gathered} 9.20 \\ (6.76) \end{gathered}$ | ** | $\begin{aligned} & 10.53 \\ & (8.50) \end{aligned}$ | ** | $\begin{gathered} 157.50 \\ (128.60) \end{gathered}$ |
| 45. | RGR at 5\% tillering | $\begin{aligned} & -18.38 \\ & (90.19) \end{aligned}$ | * | $\begin{array}{r} 210.62 \\ (324.72) \end{array}$ | ** | $\begin{aligned} & 3404.53 \\ & (287.22) \end{aligned}$ |
| 46. | RGR at $50 \%$ tillering | $\begin{gathered} -1.23 \\ (126.18) \end{gathered}$ | ** | $\begin{array}{r} 56.49 \\ (283.68) \end{array}$ | ** | $\begin{aligned} & 4254.19 \\ & (357.00) \end{aligned}$ |
| 47. | RGR at 95\% tillering | $\begin{gathered} -5.03 \\ (142.94) \end{gathered}$ | ** | $\begin{array}{r} 54.08 \\ (300.30) \end{array}$ | ** | $\begin{aligned} & 5124.98 \\ & (430.08) \end{aligned}$ |

* Significant at 5\% probability level
** Significant at $1 \%$ probability level

Table 3.4 Genotypic variance from half-sib $\left(\mathrm{V}_{\mathrm{H}}\right)$ and Plot variance $\left(\mathrm{V}_{\mathrm{HB}}\right)$ with their standard errors (Model 2)

| Characters | Half-sib |  | Plot |  |
| :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} V_{H} \\ \text { (se.) } \end{gathered}$ | F-sig. | $\begin{gathered} \mathrm{V}_{\mathrm{HB}} \\ (\mathrm{se} .) \end{gathered}$ | F-sig. |

## Botanic characters

| 1. Juvenile leaf width | 3.63 | ** | 0.28 | ns |
| :---: | :---: | :---: | :---: | :---: |
|  | (1.05) |  | (0.75) |  |
| 2. Juvenile leaf length | 30.95 | ** | -2.62 | ns |
|  | (9.17) |  | (7.07) |  |
| 3. Mature leaf width ( 15 wks ) | 0.20 | ** | -0.06 | ns |
|  | (0.07) |  | (0.07) |  |
| 4. Mature leaf width ( 33 wks ) | 0.12 | ** | 0.15 | * |
|  | (0.07) |  | (0.09) |  |
| 5. Leaf sheath purple | 0.40 | ** | -0.11 | ns |
| (12 wks) | (0.12) |  | (0.09) |  |
| 6. Leaf sheath purple | 0.46 | ** | 0.53 | ** |
| (15 wks) | (0.19) |  | (0.19) |  |
| 7. Leaf sheath purple | 0.09 | ** | 0.31 | ** |
| (33 wks) | (0.09) |  | (0.15) |  |
| 8. Plant height | 10.58 | ** | 22.86 | ** |
|  | (6.71) |  | (9.12) |  |

## Flower characters

| 9. Flower peeping day | $\begin{gathered} 0.99 \\ (0.92) \end{gathered}$ | ** | $\begin{gathered} 3.59 \\ (1.40) \end{gathered}$ | ** |
| :---: | :---: | :---: | :---: | :---: |
| First anthesis day | 0.42 | ** | 6.74 | ** |
|  | (1.15) |  | (1.97) |  |
| 11. Median flowering day | -2.60 | ns | 9.39 | ns |
|  | (2.98) |  | (6.52) |  |
| 12. Last flowering day | -6.50 | ns | 27.87 | ns |
|  | (11.02) |  | (23.43) |  |
| 13. Anthesis time | $-0.13$ $(0.38)$ | ns | $0.21$ | ns |
| 14. Anthesis position | -0.02 | ns | 0.10 | ns |
|  | (0.05) |  | (0.11) |  |
| 15. Panicle width | 23.17 | ** | 61.81 | ** |
|  | (18.12) |  | (26.63) |  |
| 16. Panicle length | 49.54 | ** | 0.79 | ns |
|  | (27.66) |  | (38.96) |  |
| 17. Panicle compactness | 0.13 | ** | 0.67 | ** |
|  | (0.18) |  | (0.29) |  |

Table 3.4 (continued)

| Characters | Half-sib |  | Plot |  |
| :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} v_{H} \\ \text { (se.) } \end{gathered}$ | F-sig. | $\begin{aligned} & \mathrm{v}_{\mathrm{HB}} \\ & (\mathrm{se} .) \end{aligned}$ | F-sig. |

## Agronomic characters

18. Clump erectness
19. Old diseases
20. New diseases
21. Leaf hairiness
22. Leaf tensile strength
23. Flavanoid at leaf sheath

## Tillering characters

| 24. Tiller No.at $5 \%$ tillering | $\begin{aligned} & -0.01 \\ & (0.42) \end{aligned}$ | ** | $\begin{gathered} 3.33 \\ (0.76) \end{gathered}$ |  |
| :---: | :---: | :---: | :---: | :---: |
| Tiller No.at $50 \%$ tillering | 14.21 | ** | 8.74 | ns |
|  | (8.45) |  | (11.80) |  |
| 26. Tiller No.at $95 \%$ tillering | $49.69$ | ** | $35.72$ | ns |
| 27. Tiller No.at flowering time | 36.07 | ** | 18.77 | ns |
|  | (14.04) |  | (14.67) |  |
| 28. No.of dead tillers at end | 13.35 | ** | 13.43 | ** |
|  | (5.46) |  | (5.79) |  |
| 29. No.of green tillers at end | -8.75 | ns | 26.59 |  |
|  | (7.45) |  | (16.80) |  |
| 30. No.of young tillers at end | 10.59 | ** | 12.14 | , |
|  | (5.52) |  | (6.98) |  |
| 31. No.of aerial tillers | 76.45 | * | -43.04 | ns |
| at end | (43.27) |  | (62.47) |  |
| 32. No.of total tillers at end | 32.07 | ns | 87.67 | ns |
|  | (68.67) |  | (124.95) |  |
| 33. No.of base tillers at end | 22.16 | ** | 82.91 | ** |
|  | (25.99) |  | (42.10) |  |
| 34. No.of base green tillers | 2.53 | * | 60.96 | ** |
| at end | (16.97) |  | (31.28) |  |
| 35. Dead tiller dryweight | 3.30 | ** | 5.17 | ** |
| at end | (1.87) |  | (2.46) |  |
| 36. Green tiller dryweight | 11.25 | ** | -0.98 | ns |
| at end | (6.59) |  | (9.46) |  |

Table 3.4 (continued)

| Characters | Half-sib |  | Plot |  |
| :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} \mathrm{v}_{\mathrm{H}} \\ \text { (se.) } \end{gathered}$ | F-sig. | $\begin{gathered} \mathrm{v}_{\mathrm{HB}} \\ (\mathrm{se} .) \end{gathered}$ | F-sig. |

## Tillering characters

| 37. Young tiller dryweight at end | $\begin{gathered} 0.21 \\ (0.11) \end{gathered}$ | ** | $\begin{gathered} 0.22 \\ (0.14) \end{gathered}$ |  |
| :---: | :---: | :---: | :---: | :---: |
| 38. Aerial tiller dryweight at end | $\begin{gathered} 8.12 \\ (7.78) \end{gathered}$ | ns | $\begin{gathered} 0.90 \\ (13.00) \end{gathered}$ | ns |
| 39. Total tiller dryweight at end | $\begin{array}{r} 3.52 \\ (18.08) \end{array}$ | ns | $\begin{aligned} & -16.54 \\ & (36.36) \end{aligned}$ | ns |
| 40. Base tiller dryweight at end | $\begin{gathered} 14.59 \\ (10.03) \end{gathered}$ | ** | $\begin{gathered} 11.08 \\ (14.85) \end{gathered}$ | ns |
| 41. Base green tiller dryweight | $\begin{aligned} & 11.21 \\ & (7.14) \end{aligned}$ | * | $\begin{gathered} -0.25 \\ (10.63) \end{gathered}$ | ns |
| 42. No.of days for $5 \%$ tillering | $\begin{gathered} 2.61 \\ (0.98) \end{gathered}$ | * | $\begin{gathered} 0.42 \\ (1.01) \end{gathered}$ | ns |
| 43. No.of days for $50 \%$ tillering | $\begin{gathered} 5.89 \\ (3.37) \end{gathered}$ | ** | $\begin{gathered} 9.30 \\ (4.42) \end{gathered}$ | ** |
| 44. No.of days for $95 \%$ tillering | $\begin{aligned} & 17.95 \\ & (9.33) \end{aligned}$ | ** | $\begin{gathered} 21.27 \\ (11.78) \end{gathered}$ | * |
| 45. Relative growth rate at $5 \%$ tillering | $\begin{gathered} 155.42 \\ (280.41) \end{gathered}$ | ** | $\begin{aligned} & 1942.37 \\ & (455.59) \end{aligned}$ | ** |
| 46. Relative growth rate at $50 \%$ tillering | $\begin{gathered} 42.63 \\ (266.84) \end{gathered}$ | ** | $\begin{aligned} & 1711.72 \\ & (470.17) \end{aligned}$ | ** |
| 47. Relative growth rate at $95 \%$ tillering | $\begin{array}{r} 33.04 \\ (287.16) \end{array}$ | ** | $\begin{aligned} & 1676.03 \\ & (513.99) \end{aligned}$ | ** |

* Significant at $5 \%$ probability level
** Significant at $1 \%$ probability level

Table 3.5 Genetic Variance components repartitioned into additive variance ( $\mathrm{V}_{\mathrm{A}}$ ) and heterotic variance $\left(V_{h}\right)$, together with phenotypic-variance $\left(V_{P}\right)$
Characters

## Botanic characters

| 1. Juvenile leaf width | 14.5 | 21.23 | 16.59 |
| :--- | :--- | ---: | ---: |
| 2. Juvenile leaf length | 123.80 | 49.72 | 165.71 |
| 3. Mature leaf width (15 wks) | 0.80 | 0.88 | 1.52 |
| 4. Mature leaf width (33 wks) | 0.44 | -1.91 | -1.04 |
| 5. Leaf sheath purple (12 wks) | 1.60 | 1.08 | 2.35 |
| 6. Leaf sheath purple (15 wks) | 1.84 | -0.65 | 2.76 |
| 7. Leaf sheath purple (33 wks) | 0.36 | 0.74 | 1.99 |
| 8. Plant height | 42.32 | 8.98 | 114.85 |

## Flowering characters

9. Flower peeping day

| 3.96 | 2.97 | 16.89 |
| ---: | ---: | ---: |
| 1.68 | -0.95 | 18.78 |
| -10.40 | 71.71 | 87.40 |
| 26.00 | 277.80 | 329.28 |
| 0.52 | 4.64 | 4.70 |
| -0.08 | 1.42 | 1.62 |
| 92.68 | 73.29 | 337.20 |
| 198.16 | 494.73 | 695.08 |
| 0.52 | 1.24 | 3.62 |

## Agronomic characters

18. Clump erectness
1.44
$-0.26$
2.67
19. Old diseases
20. New diseases
21. Leaf Hair
0.92
0.87
2.72
0.04
0.61
3.19
22. Leaf tensile strength
41.84
63.12
1.13
23. Flavanoid at leaf sheath
1.44
-0.42
219.92
$\begin{array}{llll}\text { 23. Flavanoid at leaf sheath } & 1.44 & -0.42 & 3.30\end{array}$

Table 3.5 (continued)


Tillering characters

| 24. Tiller No.at 5\% tillering | -0.04 | -3.37 | 5.45 |
| :--- | ---: | ---: | ---: |
| 25. Tiller No.at 50\% tillering | 56.84 | 118.27 | 200.02 |
| 26. Tiller No.at 95\% tillering | 198.76 | 528.22 | 828.78 |
| 27. Tiller No.at flowering time | 144.28 | 51.95 | 249.72 |
| 28. No.of dead tillers at end | 53.40 | -8.07 | 83.33 |
| 29. No.of green tillers at end | -35.00 | 181.40 | 221.38 |
| 30. No.of young tillers at end | 42.36 | 29.70 | 107.50 |
| 31. No.of aerial tillers at end | 305.80 | 1111.52 | 1290.79 |
| 32. No.of total tillers at end | 128.28 | 1641.41 | 2021.29 |
| 33. No.of base tillers | 88.64 | 240.09 | 562.54 |
| 34. No.of base green tillers | 10.12 | 223.46 | 405.49 |
| 35. Dead tiller dry weight at end | 13.20 | 6.19 | 33.87 |
| 36. Green tiller dryweight at end | 45.00 | 138.25 | 180.42 |
| 37. Young tiller dryweight at end | 0.84 | 0.76 | 2.21 |
| 38. Aerial tiller dryweight at end | 32.48 | 192.63 | 227.67 |
| 39. Total tiller dryweight at end | 14.08 | 689.02 | 656.61 |
| 40. Base tiller dryweight | 58.36 | 151.28 | 242.92 |
| 41. Base green tiller dry weight | 44.84 | 148.91 | 193.05 |
| 42. No.of days for 5\% tillering | 10.40 | 8.65 | 20.24 |
| 43. No.of days for 50\% tillering | 23.56 | 11.98 | 62.05 |
| 44. No.of days for 95\% tillering | 71.80 | 43.02 | 175.45 |
| 45. RGR at $5 \%$ tillering | 621.68 | -2578.06 | 3559.95 |
| 46. RGR at $50 \%$ tillering | 170.52 | -752.10 | 4296.82 |
| 47. RGR at 95\% tillering | 132.16 | 249.17 | 5158.02 |

[^2]The relative contribution of genetic variance to the phenotypic variance was viewed in the forms of narrow-sense heritability (average allele), heterotic-sense heritability (non-additive) and broad-sense heritability, (general genotypic) respectively. The comparison among these three estimates can be done in Table 3.6. The characters which have high narrow-sense heritability include juvenile leaf width and length, 12 wks . and 15 wks . leaf sheath purple, tiller numbers at flowering and numbers of dead tillers. The high heterotic heritability estimates include 15 wks . and 33 wks. mature leaf width, median and last flowering day, anthesis time and position, panicle length, tiller numbers at $95 \%$, numbers of green tiller, numbers of aerial tiller, and numbers of total tiller, green tiller dry-weight, aerial tiller dry-weight, base tiller dry-weight and total tiller dry-weight. Finally, the broad-sense heritability estimates are high in most characters especially in flowering and tillering characters.

Table 3.6 Heritability estimates for narrow sense $\left(h^{2} N\right)$, heterotic sense $\left(h^{2} h\right)$ and broad sense $\left(h_{B}{ }_{B}\right)$
$\qquad$

## Botanic characters

| 1. Juvenile leaf width | 0.88 | 0.07 | 0.95 |
| :--- | ---: | ---: | :--- |
| 2. Juvenile leaf length | 0.75 | 0.30 | 1.05 |
| 3. Mature leaf width $(15 \mathrm{wks})$ | 0.53 | 0.58 | 1.11 |
| 4. Mature leaf width $(33 \mathrm{wks})$ | -0.42 | 1.84 | 1.41 |
| 5. Leaf sheath purple $(12 \mathrm{wks})$ | 0.68 | 0.46 | 1.14 |
| 6. Leaf sheath purple $(15 \mathrm{wks})$ | 0.66 | -0.24 | 0.43 |
| 7. Leaf sheath purple $(33 \mathrm{wks})$ | 0.18 | 0.37 | 0.55 |
| 8. Plant height | 0.37 | 0.08 | 0.45 |

## Flowering characters

9. Flower peeping day

| 0.23 | 0.18 | 0.41 |
| ---: | ---: | ---: |
| 0.09 | -0.05 | 0.04 |
| -0.12 | 0.82 | 0.70 |
| -0.08 | 0.84 | 0.76 |
| 0.11 | 0.98 | 0.88 |
| -0.05 | 0.88 | 0.83 |
| 0.27 | 0.22 | 0.49 |
| 0.29 | 0.71 | 1.00 |
| 0.14 | 0.34 | 0.49 |

## Agronomic characters

18. Clump erectness
0.54
19. Old diseases (Rust)
20. New diseases (other)
21. Leaf hairiness
22. Leaf tensile strength
23. Flavanoid at leaf sheath
0.10
0.29
0.04
0.19
0.44

| -0.10 | 0.44 |
| ---: | ---: |
| 0.32 | 0.42 |
| -0.02 | 0.27 |
| 0.54 | 0.58 |
| 0.29 | 0.48 |
| -0.13 | 0.31 |

Table 3.6 (continued)


## Tillering characters

| 24. Tiller No.at $5 \%$ tillering | -0.007 | -0.62 | -0.63 |
| :---: | :---: | :---: | :---: |
| 25. Tiller No.at $50 \%$ tillering | 0.28 | 0.59 | 0.88 |
| 26. Tiller No.at $95 \%$ tillering | 0.24 | 0.64 | 0.88 |
| 27. Tiller No.at flowering time | 0.58 | 0.21 | 0.79 |
| 28. No.of dead tillers at end | 0.64 | -0.10 | 0.54 |
| 29. No.of green tillers at end | -0.16 | 0.82 | 0.66 |
| 30. No.of young tillers at end | 0.39 | 0.28 | 0.67 |
| 31. No.of aerial tillers at end | 0.24 | 0.86 | 1.10 |
| 32. No.of total tillers at end | 0.06 | 0.81 | 0.88 |
| 33. No.of base tillers at end | 0.16 | 0.43 | 0.58 |
| 34. No.of base green tiller at end | 0.03 | 0.55 | 0.58 |
| 35. Dead tiller dry weight at end | 0.39 | 0.18 | 0.57 |
| 36. Green tiller dryweight at end | 0.25 | 0.77 | 1.02 |
| 37. Young tiller dryweight at end | 0.38 | 0.34 | 0.72 |
| 38. Aerial tiller dryweight at end | 0.14 | 0.85 | 0.99 |
| 39. Total tiller dryweight at end | 0.13 | 1.07 | 1.21 |
| 40. Base tiller dryweight at end | 0.24 | 0.62 | 0.86 |
| 41. Base green tiller dry weight | 0.23 | 0.77 | 1.00 |
| 42. No.of days for $5 \%$ tillering | 0.51 | 0.43 | 0.94 |
| 43. No.of days for $50 \%$ tillering | 0.38 | 0.19 | 0.57 |
| 44. No.of days for $95 \%$ tillering | 0.41 | 0.25 | 0.65 |
| 45. RGR at $5 \%$ tillering | 0.17 | -0.72 | -0.55 |
| 46. RGR at $50 \%$ tillering | 0.04 | -0.18 | -0.14 |
| 47. RGR at $95 \%$ tillering | 0.03 | 0.05 | 0.07 |

* Significant at $5 \%$ probability level
** Significant at $1 \%$ probability level


## CHAPTER 4

## DISCUSSION

### 4.1 Comparison Among Topodeme, Half-sib and Plant Variations

The topodeme variation is derived from the differentiation among means of local populations, the open-pollinated seeds were collected from several locations throughout New Zealand. Whereas the half sib variation is confined to among plants within each topodeme. In the other word, the half sib variation is the allele effects amongst single plants within topodemes originally used as self mother plants. The within plot variation is the plant to plant variation within half sib families or lines. Hence, the total plant to plant variation within topodemes is the half sib variance and within variance combined.

In this study, the half-sib family variance has a higher value than topodeme variance in most characters ( 39 out of 47 characters) The exception were: 33 wks . mature leaf blade width, 15 wks . leaf sheath purple ,flavanoid, anthesis time, anthesis position, panicle compactness, new disease and leaf hairiness. And plant-to-plant within half sib variance has a higher value than topodeme variance in every character. This has affirmed the speculation from the previous work conducted by Cameron (1979)

It is of some interest to compare this result with those from other species. The within-population of Trifolium repens from a uniform pasture found a great deal of variation in several characters; and even as great as that between populations from different environments in some cases (Burdon and Harper, 1980). The breeding system of a species could affect on the amount of genetic variation within and between populations (Levin, 1978). The population of cross-fertilizng species was less differentiated inter se than the population of self-fertilizing species. In Trifolium spp., outbreeders had more within population heterogeneity for quantitative characters and less between-population heterogeneity than inbreeder (Katznelson, 1969 cited from

Levin,1978). However, other workers found a great deal variation in predominantly selfing-species and concluded that patterns of variation was not confined to one group of species or the others (Allard 1975; Jain, 1976).

In practice, selection could be more effective on the half-sib family level than the topodeme level. The germplasm collection and maintenance would be more benefitial to pay attention on subsamples within topodemes or half-sib families than among samples of topodemes.

The ecotypes of Holcus spp. in New Zealand was proposed by earlier workers (Munro, 1961). The high level of half-sib and plant variations (and much higher than the topodeme in some traits) in the present finding may suggest that there are no ecotype nor major topodeme differences in New Zealand. The situation was quite similar to Phalaris tuberosa in Australia where Trumble and Cashmore (1934) found no evidence of ecotypic differentiation among samples from various parts of Australia, despite the fact that the species had at that time been established in relatively small but widespread areas for long time.

### 4.2 Genetic Variance and Heritability

Significant genetic variation is detected among half sib progenies for numerous Yorkshire Fog grass characteristics. These results concur with earlier reports in Yorkshire Fog grass for several characters (Cameron,1979; Billington et al.,1988).

In the analysis of quantitative variability and heritability in predominantly cross-fertilized forage species, it is convenient to make use of family groups produced by natural crossing. The offspring is often derived from the ovules of a maternal plant which has been pollinated without control of male parentage (pollen), and these form half-sib progenies or lines. For the analysis, it is assumed that the offspring were produced under random mating (no inbreeding). However, some traits may be also under the influence of maternal effect and phenotypic assortive mating. The maternal effect might cause bias estimates of heritability if they were ignored. From such fact, the use of field collected maternal sibships needs to be cautious. Paternal analysis
indicated that these progeny were not likely to be half-sibs (Ellstrand,1984), which could cause overestimates of heritability.

Some estimates of phenotypic and genotypic variances are negative. And they, in turn, have caused the inflated or negative heritability estimates in some attributes. This is possible because of the sampling distribution of trivial parameters or non-random sampling of genotypes from the natural population (Falconer,1981) It is also possible that estimates of narrow-sense heritabilities may be biased by the confounding of nonadditive genetic variance (Mitchell-Olds and Rutledge, 1986).

These heritability estimates are on an individual plant basis, and vary from low to high. The broad-sense heritability estimates are low (0.04) for the first day of anthesis to very high (1.0) for juvenile leaf-width, mature leaf width at 15 and 30 weeks, the panicle length and purple leaf sheath at 12 weeks. The narrow-sense heritability estimates are relatively high to medium in most of the botanical and tillering attributes. But most of agronomic traits showed medium to low narrow-sense heritability. Although, Moll and Stuber (1974) concluded that the genetic variability of many important agronomic traits of forage crops had been found to be predominantly additive.

### 4.2.1 Botanical Characters

The heritability estimates for the most botanical characters are similar to those of other workers with other grasses. The broad-sense and narrow-sense heritability estimates for juvenile leaf width and leaf length of Yorkshire Fog grass are relatively high while the heritability estimates of Italian ryegrass seedlings for leaf width and leaf length were medium, ( 0.38 and 0.42 , respectively)( Cooper and Edwards, 1961).

The broad-sense heritability estimates of mature leaf blade width both at 15 and 30 weeks are high and the narrow-sense one is medium at 15 weeks and high at 30 weeks. Similar result was shown in Bermudagrass (Cynodon dactylon) whose broadsense and narrow-sense heritability estimates of leaf blade width were high and medium ( 0.83 and 0.62 , respectively). However, Cameron (1979) and Billington (1988) had found that the broad-sense and narrow-sense heritability for this attributes was relatively low ( 0.08 and 0.17 ,respectively). Furthermore, the leaf width mean
tended to change with time. This study has unraveled some genetic variance pattern on it. At 15 weeks, there are almost half additive variance and heterotic variance, but at 30 weeks, it alters to become all heterotic. The leaf-width grand mean towards the narrowness indicates that the leaf narrowness is under the heterotic heritability.

There are some contrasting patterns in broad-sense and narrow-sense heritabilities for leaf sheath purple of different time periods. The difference possibly indicates that there has been a trigger, or change of genetic control. A possible external trigger may have been the caging, which occurred between the two measuring periods. Further research should resolve this issue.

The broad-sense and narrow-sense heritability estimates of plant height in Yorkshire Fog grass are medium and low in value, respectively. The pattern was very similar to other grasses. In the following examples, the broad-sense heritability estimates were ranged from 0.4-0.6 namely: for Nebraska populations Indiangrass (Sorghastrum nutans) was 0.4 (Vogel, et al., 1980), for reed canarygrass (Phalaris arundizacea) in Eastern Canada population was 0.54 (Sachs and Coulman, 1983), for sand bluestem (Andropogon halhi) was 0.62 (Riley,1982), for Rhodes grass (Chloris gayana) was 0.66 (Quesenberry et al.,1978). The narrow-sense heritability was also very similar to guineagrass (Panicum maximum) which was rather low (0.2) (Usberti and Jain, 1978).

### 4.2.2 Flowering Characters

Most of these flowering characters have medium to high broad-sense heritability with heterotic variances prevailing. There are some variations amongst different flowering measurements. The first day and the median day of flowering may be under different sets of gene control. The first anthesis day has very small additive genetic variance, only 4 percent and very large environmental variance, about 96 percent. This suggests an invariant mechanism for flowering initiation. In contrast, later flowering controls have stronger genetic variability. Both the median flowering day and last flowering have a very high genetic variance and all of which is heterotic. (as shown by $h_{h}^{2}$ ). But they have fewer environmental variance, only about 30 percent.

It shows the same trend in this study. Billington (1988) found that the narrowsense heritability of flowering time in Yorkshire Fog grass was relatively low in traditional field and medium in improved field population, respectively. In general, the heritability estimates for median flowering or heading day are quite similar to other grasses. In Indiangrass (Sorghastrum nutans) of Nebraska populations was 0.5-0.7 (Vogel, et al.,1980), in (Lolium perenne) was 0.94, in canarygrass (Phalaris arundizacea) was 0.94 (Sachs and Coulman, 1983) and in sand bluestem was 0.73 (Riley,1982).

The panicle length had nearly the same amount of narrow-sense heritability as Billington (1988) had found, but the broad-sense heritability was considerably larger.

### 4.2.2 Agronomic Characters

For agronomic characters, most have a medium broad-sense heritability and low to medium for broad-sense heritability.

Clump erectness at vegetative stage, just prior to stem elongation, had a relatively medium (0.44) estimate for narrow-sense heritability. This was different to a previous study by Cameron (1979) which reported a low estimate (0.10). The differences in the two results arise from this: Cameron's material was a different sample from the same germplasm but it could also be due to scoring at different stages of growth.

Sheep performance has been associated with leaf cellulose content which may be positively correlated with leaf tensile strength in ryegrass (Lolium spp.). Weight gains have been reported highest on the grasses with the lowest strengths (Wilson,1965; Evan,1967b). The present study reveals significant genetic differences in leaf tensile strength. This has been reported also amongst lines of weeping lovegrass (Eragrostis curvala) and amongst clones of Bermudagrass (Cynodon dactylon), sideoats grama (Bouteloua curtipendula) and sand bluestem (Andropogon hallii) (Kneebone, 1960). In this study, a medium level of broad-sense heritabilty was found which was different from Cameron's result which showed a very low value (0.01-0.04). It was also different from other grasses. The broad-sense heritability estimates for leaf tensile strength in tall fescue (Festuca arundinacea) were relatively
high, ranking from 0.83 in June to 0.93 in August to 0.85 in October. The narrowsense heritability estimates were also high (0.7-0.8) (Nguyen,et al.,1982). The genetic control mechanism might change according to the seasonal cycle or growth stages. Further investigation is needed to resolve the issue.

The flavanol level had a relatively medium narrow-sense heritability. The pattern was rather similar to leaf sheath purple at 12 weeks. It has been summarized that the purple colour is flavanoid in nature and the similarity of the two heritabilities may support this possibility.

Leaf hair has both high broad-sense and narrow-sense heritability. It contrasted to Cameron's (1979) result which indicated a low broad-sense heritability (0.2). The result was similar to that for Medicago where the narrow-sense heritability of hair density was medium (0.55)(Kitch, et al.,1985).

Leaf diseases are categorized into old disease,i.e. mostly rust, and new diseases, i.e. leaf spot (symptom similar to Helminthosporium leaf spot). Both have relatively low to medium broad-sense heritabilities and low narrow-sense heritabilities. These results are similar to those of other grasses. The realized heritabilities for rust resistance on eight cultivars of tall fescue (Festuca arundinacea) ranked from 0.07, $0.08,0.16,0.18,0.36,0.45,0.49$ and 0.52 , respectively. It was concluded that there might. be different gene system for rust resistance in different population. Also, the low heritability one might be the result of some non-additive gene action for rust resistance (Wofford and Watson, 1982). In this study, plants have a low narrow-sense heritability on rust resistance, while the heterotic variance is three time higher than the additive variance. The high non-additive variance indicated it might not be easy to select for in traditional selection nursery methods. This contrasted to Munro's (1961) recommendation for rust resistance relating to easily selected major genes. In case of leaf spot, in meadow fescue (Festuca pratensis), the narrow-sense heritability for Helminthosporium was medim (0.49) (Frandsen et al.,1981). This indicated that it might be easier to select for leaf spot disease resistance in the traditional selection nursery methods.

### 4.3 Genetic Variance on Tiller Development

Tiller development starting from sowing till flowering observed by tiller numbers has expressed virtually in a logistic function (Fig.4.1). Growth analysis of a permanent pasture in Normandy in spring revealed that Holcus lanatus growth followed a sigmoid curve (Lemaire, et al.,1982). In Lolium, however, the tiller number in the early stage were increasing exponentially (Cooper and Edwards, 1961). It is interesting to note that the lower half of a logistic is exponential (Causton, 1977)

Grasses are likely to developed the tillers successively and continuously without any distinct termination of the whole tillering process. This complies very well to Protich's (1977) descriptive work. Although flowering tillers died soon after seed maturity, the new young tillers emerged from the ground thereafter. During the heading and seed development periods, grasses had possessed a great number of elongated green tillers and aerial tillers directly from their green tillers.

For tiller number, the broad-sense heritability estimates across time are from zero to very high (0.88). The narrow-sense estimates are from zero to medium (0.28) and then low (0.07)(Fig.4.1 and Fig.4.2). Billigton et al. (1988) unraveled the same pattern of medium broad-sense heritability and low narrow-sense heritability for ten week growth of Yorkshire Fog grass. Similar trend also occurs in the other grasses. The broad-sense heritability estimates of tillers on two month-old Lolium from sowing were medium to high (0.4-0.8) (Cooper and Edwards, 1961). In reed canary grass, both broad-sense and narrow-sense heritability were high for tiller number (Casler, 1984). In guineagrass, heritability estimates based on parent-offspring regression for total tiller number were relatively low to medium (0.3) (Usberti and Jain, 1978). In maize, however, the genetic component of variation for tillering was believed due to general combining ability (Rood and Major, 1981)

Both broad-sense and narrow-sense heritabilities for flowering tillers are high and medium, respectively. This was somewhat comparable to what Billington et al. (1988) finding which revealed a medium to low heritabilities for both improved-field and traditional field population. The young tillers which have emerged after flower tiller died, show the same pattern of genetic and environment variation. This might indicate the recycle of genetic control in Yorkshire Fog grass.


Figure 4.1 Genotypic variance of tiller number development from sowing to flowering stage

No. of
Tillers


Figure 4.2 Genotypic variance of tiller number after main tiller flowering stage (33 weeks)

For dry matter, a high broad-sense and medium to low narrow-sense heritabilities are obtained for every type of tiller at old plant stage ( 33 wks .)(Fig.4.3). That basal tiller mass (which included dead (post-flowering) tillers, green tillers and young tillers) represents the mass in pasture, and is of particular interest. It has a medium broad-sense heritability ( 0.43 ) and low narrow-sense heritability ( 0.16 ), which are somewhat comparable to other grass species. The broad-sense heritability of mass (yield) in several grasses was medium to low (Clements,1969; Marum et al,1979; Oram, et al.,1974, Shenk and Westerhaus,1982). Dry matter/plant in Lolium multiforum had medium broad-sense heritability (0.48)(Bugge,1984). Also, in reed canary grass, for the tiller dry weight per plant, broad-sense heritabilities were relatively medium (0.3-0.6) (Casler, 1981); as it was in Lolium perenne (0.53) (Utz and Oettler, 1978).

Similar results of heritability estimates for dry matter with respect to variability within established genotypes of crossed fertilized species have been drawn by Cooper(1959) on Lolium, Gardner (1963) on yield of maize and by Kehr and Gardner (1960) on forage yield in lucern

The relative growth rate at $5 \%, 50 \%$ and $95 \%$ asymptote show quite a similar patterns in their variance components (Fig.4.1). At very young stages, the plant has only environmental variances in action. The $95 \%$ stage has 92 percent of environmental variance with only 3 percent additive and 5 percent heterotic variance. The timing to reach $5 \%, 50 \%$, and $95 \%$ of growth have results very different to those of relative growth rate and tiller numbers. These generally are high and medium broad-sense heritabilities and relatively medium narrow-sense heritabilities. This showed, clearly, the different genetic perspectives represented by growth rate and timings. As for the flowering attributes, the duration of events was shown to have greater genetic variabilities than either their initiation (for flowering) or the rates of change (for tillering).

### 4.4 Implication for plant breeding

The detailed genetic analysis of a locally populations is of practical interest in setting up a effective plant breeding programme. The initial step in any of them is the

Tiller
Dry weighl
(gmi.)


Figure 4.3 Genotypic variance of tiller dry matler after main tiller flowering stage
choice of a suitable base population. The alternatives, in the case of cross-fertilized species, will often include:

1. the improvement of an established populations by intra-population selection, or
2. the formation of a more widely based genetic population by the incorporation of introduced materials (wide crosses).

The useful genetic variation presented in the local New Zealand populations of Holcus has been found to be quite appreciable for most of the characters studied. The genetic advance under selection for these characters depends on the amount of genetic variation available and on its heritability.

These results indicate that genetic advance for the characters: juvenile leaf blade width and length, 15 weeks mature leaf blade width, purple leaf sheath, plant height, clump erectness, flavanol, panicle width, number of dead tillers and young tillers, dead tiller mass, young tiller mass, number of days to reach $5 \%, 50 \%$ and $95 \%$ of growth stage should be possible using breeding methods which utilize additive genetic variation. The traditional breeding methods such as mass selection, line selection, line breeding or simple recurrent selection should be efficient methods for the improvement of these attributes.

Many characters exhibit low narrow-sense heritability but high heterotic heritability, are included : 30 wks . mature leaf blade width, leaf tensile strength, leaf hair, old disease, median and last flowering day, panicle length and compactness, total tiller number, green tiller number, number of tiller at $50 \%$ and $95 \%$ of growth stage, basal tiller number, total tiller mass, basal tiller mass and green tiller mass. These require some combination of progeny testing and recurrent selection or top cross progeny tests for high specific combining ability for development of synthetic cultivars or special forms of recurrent selection bulks.

Some further research would be desirable. For example, estimates of correlation was needed because it would assist in estimating the relative efficiency of direct and indirect selection for characters which were easier to evaluate than others.

For instance, a high total genetic contribution in juvenile leaf size criteria might be used as indirect selection for some other high genetic correlated responses.

For those characters which had different genotypic variances across time (eg. purple leaf sheath, flowering day and tiller number development), it would be good practice to select at the period with a higher level of genotypic variance. For instance, amongst the flowering characters, selection would be more effective on the median flowering day, than on the first day of flowering. Also, the number of tillers would best be selected in the later stages of development ( $50 \%$ and $95 \%$ of growth stages).

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## APPENDIX 1

Duncan's multiple range test for juvenile leaf blade width

| No. | Clu | us 7 | Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 |  | 69 | 147 | 24.9 | defghijk |
| 2 |  |  |  | 148 | 24.3 | defghijk |
| 3 |  |  |  | 150 | 22.5 | ijk |
| 4 |  |  |  | 151 | 24.9 | defghijk |
| 5 |  |  |  | 154 | 21.9 | k |
| 6 |  |  | 45 | 429 | 26.2 | cdefghijk |
| 7 |  |  |  | 430 | 25.2 | defghijk |
| 8 |  |  |  | 431 | 32.1 | a |
| 9 |  |  |  | 432 | 26.8 | cdefghi |
| 10 |  |  |  | 435 | 28.2 | bcde |
| 11 | 2 |  | 97 | 260 | 25.9 | defghijk |
| 12 |  |  |  | 261 | 26.1 | cdefghijk |
| 13 |  |  |  | 262 | 26.7 | cdefghij |
| 14 |  |  |  | 264 | 27.9 | bcdef |
| 15 |  |  |  | 266 | 26.9 | cdefghi |
| 16 |  |  | 31 | 369 | 21.9 | k |
| 17 |  |  |  | 371 | 27.2 | bcdefgh |
| 18 |  |  |  | 372 | 27.3 | bcdefg |
| 19 |  |  |  | 376 | 22.9 | hijk |
| 20 |  |  |  | 377 | 25.4 | defghi |
| 21 | 3 |  | 15 | 36 | 27.6 | bcdef |
| 22 |  |  |  | 37 | 27.4 | bcdefg |
| 23 |  |  |  | 38 | 23.9 | efghijk |
| 24 |  |  |  | 40 | 25.7 | defghijk |
| 25 |  |  |  | 42 | 24.4 | defghijk |
| 26 |  | 27 | 7 | 64 | 26.9 | cdefghi |
| 27 |  |  |  | 66 | 22.3 | jk |
| 28 |  |  |  | 67 | 25.2 | defghijk |
| 29 |  |  |  | 69 | 23.2 | ghijk |
| 30 |  |  |  | 71 | 23.6 | fghijk |
| 31 | 4 |  |  | 233 | 24.0 | defghijk |
| 32 |  |  |  | 234 | 31.3 | ab |
| 33 |  |  |  | 235 | 22.8 | ijk |
| * 34 |  |  |  | 236 | 23.9 | defghijk |
| 35 |  |  |  | 237 | 25.4 | defghijk |
| 36 |  | 154 |  | 460 | 23.6 | fghijk |
| 37 |  |  |  | 461 | 30.3 | abc |
| 38 |  |  |  | 462 | 25.1 | defghijk. |
| 39 |  |  |  | 463 | 23.6 | fghijk |
| 40 |  |  |  | 465 | 25.6 | defghijk |
| 41 | 5 |  |  | 25 | 26.3 | cdefghijk |
| 42 |  |  |  | 26 | 22.1 | k |
| 43 |  |  |  | 27 | 26.8 | cdefghi |
| 44 |  |  |  | 31 | 25.9 | defghijk |
| 45 |  |  |  | 32 | 24.8 | defghijk |
| 46 |  | 142 |  | 414 | 26.1 | cdefghijk |
| 47 |  |  |  | 415 | 23.0 | ghijk |
| 48 |  |  |  | 416 | 24.0 | defghijk |
| 49 |  |  |  | 418 | 28.3 | abcd |
| 50 |  |  |  | 420 | 26.8 | cdefghi |

Means with the same letter are not significantly different at $5 \%$


Means with the same letter are not significantly different at $5 \%$

```
Duncan's multiple range test for mature leaf blade width at }15\mathrm{ weeks
\begin{tabular}{|c|c|c|c|c|c|}
\hline No. & Clus & s 'ropo & Half-sib & Means & \\
\hline 1 & 1 & 69 & 147 & 9.6 & efgh \\
\hline 2 & & & 148 & 10.8 & abcdef \\
\hline 3 & & & 150 & 10.7 & abcdef \\
\hline 4 & & & 151 & 10.4 & abcdefgh \\
\hline 5 & & & 154 & 10.0 & defgh \\
\hline 6 & & 145 & 429 & 10.0 & defgh \\
\hline 7 & & & 430 & 11.5 & ab \\
\hline 8 & & & 431 & 10.9 & abcde \\
\hline 9 & & & 432 & 10.4 & abcdefgh \\
\hline 1.0 & & & 435 & 10.9 & abcde \\
\hline 11. & 2 & 97 & 260 & 9.6 & efgh \\
\hline 12 & & & 261 & 9.8 & defgh \\
\hline 13 & & & 262 & 10.7 & abcdef \\
\hline 14 & & & 264 & 10.2 & abcdefgh \\
\hline 15 & & & 266 & 10.1 & bcdefgh \\
\hline 16 & & 131 & 369 & 9.1 & h \\
\hline 17 & & & 371 & 10.4 & abcdefgh \\
\hline 18 & & & 372 & 10.6 & abcdef \\
\hline 19 & & & 376 & 9.9 & defgh \\
\hline 20 & & & 377 & 10.5 & abcdefg \\
\hline 21 & 3 & 15 & 36 & 10.7 & abcdef \\
\hline 22 & & & 37 & 11.1 & abcd \\
\hline 23 & & & 38 & 11.3 & \(a b c\) \\
\hline 24 & & & 40 & 10.7 & abcdef \\
\hline 25 & & & 42 & 9.8 & defgh \\
\hline 26 & & 27 & 64 & 11.3 & \(a b c\) \\
\hline 27 & & & 66 & 10.5 & abodef \\
\hline 28 & & & 67 & 10.4 & abodefigh \\
\hline 29 & & & 69 & 9.7 & efgh \\
\hline 30 & & & 71 & 10.5 & abcdef \\
\hline 31 & 4 & 91 & 233 & 9.5 & fgh \\
\hline 32 & & & 234 & 10.7 & abcdef \\
\hline 33 & & & 235 & 10.2 & bcdefgh \\
\hline 34 & & & 236 & 9.5 & fgh \\
\hline 35 & & & 237 & 10.4 & abcdefgh \\
\hline 36 & & 154 & 460 & 10.5 & abcdef \\
\hline 37 & & & 461 & 10.0 & defgh \\
\hline 38 & & & 462 & 10.4 & abcdefg \\
\hline 39 & & & 463 & 9.9 & defgh \\
\hline 40 & & & 465 & 10.0 & cdefgh \\
\hline 41 & 5 & 13 & 25 & 10.6 & abcdef \\
\hline 42 & & & 26 & 10.6 & abcdef \\
\hline 43 & & & 27 & 9.9 & defgh \\
\hline 44 & & & 31 & 10.0 & defgh \\
\hline 45 & & & 32 & 9.9 & defgh \\
\hline 46 & & 142 & 414 & 9.4 & fgh \\
\hline 47 & & & 415 & 10.2 & abcdefgh \\
\hline 48 & & & 416 & 9.2 & gh \\
\hline 49 & & & 418 & 11.5 & a \\
\hline 50 & & & 420 & 10.9 & abcde \\
\hline
\end{tabular}
Means with the same letter are not significantly different at 5%
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Duncan's multiple range test for mature leaf blade width at 33 weeks
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Means with the same letter are not significantly different at 5%
```



Dumoan's multiple range test for leaf sheath purple at 15 weeks


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Duncan's multiple range test for leaf sheath purple at 33 weeks
\begin{tabular}{|c|c|c|c|c|c|}
\hline No. & Clus & S Topo & Half-sib & Mean & \\
\hline 1 & 1. & 69 & 147 & 8.3 & \(a b c\) \\
\hline 2 & & & 148 & 9.0 & \(a b c\) \\
\hline 3 & & & 150 & 9.2 & \(a b c\) \\
\hline 4 & & & 151 & 8.9 & \(a b c\) \\
\hline 5 & & & 1.54 & 9.9 & \(a b\) \\
\hline 6 & & 145 & 429 & 9.4 & abo \\
\hline 7 & & & 430 & 8.9 & abc \\
\hline 8 & & & 431 & 9.4 & \(a b c\) \\
\hline 9 & & & 432 & 9.6 & ab \\
\hline 10 & & & 435 & 9.0 & \(a b c\) \\
\hline 11 & 2 & 97 & 260 & 9.4 & \(a b c\) \\
\hline 12 & & & 261 & 9.0 & \(a b c\) \\
\hline 13 & & & 262 & 9.8 & \(a b\) \\
\hline 14 & & & 264 & 9.0 & ab \\
\hline 15 & & & 266 & 10.0 & a \\
\hline 16 & & 131 & 369 & 7.4 & bc \\
\hline 17 & & & 371 & 8.0 & \(a b c\) \\
\hline 18 & & & 372 & 9.6 & \(a b\) \\
\hline 19 & & & 376 & 9.1 & \(a b c\) \\
\hline 20 & & & 377 & 9.4 & \(a b c\) \\
\hline 21 & 3 & 15 & 36 & 9.7 & \(a b\) \\
\hline 22 & & & 37 & 9.9 & \(a b\) \\
\hline 23 & & & 38 & 8.2 & \(a b c\) \\
\hline 24 & & & 40 & 9.0 & \(a b c\) \\
\hline 25 & & & 42 & 9.6 & \(a b\) \\
\hline 26 & & 27 & 64 & 9.3 & \(a b c\) \\
\hline 27 & & & 66 & 8.1 & abc \\
\hline 28 & & & 67 & 8.5 & \(a b c\) \\
\hline 29 & & & 69 & 8.9 & \(a b c\) \\
\hline 30 & & & 71 & 9.8 & \(a b\) \\
\hline 31 & 4 & 91 & 233 & 8.4 & \(a b c\) \\
\hline 32 & & & 234 & 8.9 & \(a b c\) \\
\hline 33 & & & 235 & 8.9 & \(a b c\) \\
\hline 34 & & & 236 & 7.9 & \(b \mathrm{c}\) \\
\hline 35 & & & 237 & 9.0 & \(a b c\) \\
\hline 36 & & 154 & 460 & 7.9 & bc \\
\hline 37 & & & 461 & 9.7 & ab \\
\hline 38 & & & 462 & 9.6 & ab \\
\hline 39 & & & 463 & 9.6 & \(a b\) \\
\hline 40 & & & 465 & 10.0 & a \\
\hline 41 & 5 & 13 & 25 & 9.3 & \(a b c\) \\
\hline 42 & & & 26 & 8.4 & \(a b c\) \\
\hline 43 & & & 27 & 8.4 & \(a b c\) \\
\hline 44 & & & 31 & 9.6 & \(a b c\) \\
\hline 45 & & & 32 & 9.2 & \(a b c\) \\
\hline 46 & & 142 & 414 & 7.9 & bc \\
\hline 47 & & & 415 & 9.6 & \(a b\) \\
\hline 48 & & & 416 & 8.9 & abc \\
\hline 49 & & & 418 & 9.7 & ab \\
\hline 50 & & & 420 & 8.9 & abc \\
\hline
\end{tabular}
Means with the same letter are not significantly different at \(5 \%\)
```

| No.Clus |  | s Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 105.6 | abcdefg |
| 2 |  |  | 148 | 109.2 | abcdefg |
| 3 |  |  | 150 | 110.7 | abcdefg |
| 4 |  |  | 151 | 109.6 | abcdefg |
| 5 |  |  | 154 | 113.9 | abcdef |
| 6 |  | 145 | 429 | 98.9 | efg |
| 7 |  |  | 430 | 107.9 | abcdefg |
| 8 |  |  | 431 | 112.3 | abcdefg |
| 9 |  |  | 432 | 114.6 | abcde |
| 10 |  |  | 435 | 105.8 | abcdefg |
| 11 | 2 | 97 | 260 | 104.2 | bcdefg |
| 12 |  |  | 261 | 107.7 | abcdefg |
| 13 |  |  | 262 | 121.4 | a |
| 14 |  |  | 264 | 114.6 | abcde |
| 15 |  |  | 266 | 111.9 | abcdefg |
| 16 |  | 131 | 369 | 98.5 | fg |
| 17 |  |  | 371 | 110.6 | abcdefg |
| 18 |  |  | 372 | 114.8 | abcde |
| 19 |  |  | 376 | 106.9 | abcdefg |
| 20 |  |  | 377 | 108.3 | abcdefg |
| 21 | 3 | 15 | 36 | 117.5 | $a b$ |
| 22 |  |  | 37 | 111.5 | abcdefg |
| 23 |  |  | 38 | 104.7 | bcdefg |
| 24 |  |  | 40 | 101.4 | cdefg |
| 25 |  |  | 42 | 106.7 | abcdefg |
| 26 |  | 27 | 64 | 108.6 | abcdefg |
| 27 |  |  | 66 | 102.4 | bcdef |
| 28 |  |  | 67 | 97.6 | $g$ |
| 29 |  |  | 69 | 104.4 | bcdefg |
| 30 |  |  | 71 | 107.1 | abcdefg |
| 31 | 4 | 91 | 233 | 116.0 | abcd |
| 32 |  |  | 234 | 116.9 | $a b c$ |
| 33 |  |  | 235 | 104.6 | bcdefg |
| 34 |  |  | 236 | 106.3 | abcdefg |
| 35 |  |  | 237 | 114.1 | abcdef |
| 36 |  | 154 | 460 | 100.2 | defg |
| 37 |  |  | 461 | 100.8 | defg |
| 38 |  |  | 462 | 108.0 | abcdefg |
| 39 |  |  | 463 | 101.4 | cdefg |
| 40 |  |  | 465 | 113.5 | abcdefg |
| 41 | 5 | 13 | 25 | 106.8 | abcdefg |
| 42 |  |  | 26 | 105.8 | abcdefg |
| 43 |  |  | 27 | 112.5 | abcdefg |
| 44 |  |  | 31 | 109.4 | abcdefg |
| 45 |  |  | 32 | 100.0 | defg |
| 46 |  | 142 | 414 | 106.8 | abcdefg |
| 47 |  |  | 415 | 101.6 | cdefg |
| 48 |  |  | 416 | 100.6 | defg |
| 49 |  |  | 418 | 107.4 | abcdefg |
| 50 |  |  | 420 | 108.5 | abcdefg |



| No.Clus |  | s Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 139.2 | $a b$ |
| 2 |  |  | 148 | 130.4 | b |
| 3 |  |  | 150 | 138.1 | $a b$ |
| 4 |  |  | 151 | 136.1 | $a b$ |
| 5 |  |  | 154 | 137.1 | $a b$ |
| 6 |  | 145 | 429 | 131.2 | b |
| 7 |  |  | 430 | 128.4 | b |
| 8 |  |  | 431 | 137.6 | $a b$ |
| 9 |  |  | 432 | 129.9 | b |
| 10 |  |  | 435 | 129.9 | b |
| 11 | 2 | 97 | 260 | 132.1 | b |
| 12 |  |  | 261 | 145.3 | a |
| 13 |  |  | 262 | 136.9 | $a b$ |
| 14 |  |  | 264 | 131.6 | b |
| 15 |  |  | 266 | 135.4 | $a b$ |
| 16 |  | 131 | 369 | 130.8 | b |
| 17 |  |  | 371 | 135.8 | $a b$ |
| 18 |  |  | 372 | 136.6 | $a b$ |
| 19 |  |  | 376 | 133.1 | $a b$ |
| 20 |  |  | 377 | 136.4 | $a b$ |
| 21 | 3 | 15 | 36 | 139.2 | $a b$ |
| 22 |  |  | 37 | 133.4 | $a b$ |
| 23 |  |  | 38 | 134.3 | $a b$ |
| 24 |  |  | 40 | 128.2 | b |
| 25 |  |  | 42 | 136.5 | $a b$ |
| 26 |  | 27 | 64 | 131.3 | b |
| 27 |  |  | 66 | 129.2 | b |
| 28 |  |  | 67 | 134.8 | $a b$ |
| 29 |  |  | 69 | 132.9 | $a b$ |
| 30 |  |  | 71 | 135.1 | $a b$ |
| 31 | 4 | 91 | 233 | 137.6 | $a b$ |
| 32 |  |  | 234 | 130.4 | b |
| 33 |  |  | 235 | 130.9 | b |
| 34 |  |  | 236 | 135.0 | $a b$ |
| 35 |  |  | 237 | 133.8 | $a b$ |
| 36 |  | 154 | 460 | 130.8 | b |
| 37 |  |  | 461 | 131.8 | b |
| 38 |  |  | 462 | 138.0 | $a b$ |
| 39 |  |  | 463 | 137.1 | $a b$ |
| 40 |  |  | 465 | 134.8 | $a b$ |
| 41 | 5 | 13 | 25 | 132.3 | $a b$ |
| 42 |  |  | 26 | 136.3 | $a b$ |
| 43 |  |  | 27 | 128.6 | b |
| 44 |  |  | 31 | 139.2 | $a b$ |
| 45 |  |  | 32 | 135.3 | $a b$ |
| 46 |  | 142 | 414 | 134.0 | $a b$ |
| 47 |  |  | 415 | 137.4 | $a b$ |
| 48 |  |  | 416 | 134.3 | $a b$ |
| 49 |  |  | 418 | 131.8 | b |
| 50 |  |  | 420 | 132.2 | ab |

Duncan's multiple range test for the median flowering day


Means with the same letter are not significantly different at $5 \%$


| No.Clus |  | Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 2.4 | $a b$ |
| 2 |  |  | 148 | 1.3 | $a b$ |
| 3 |  |  | 150 | 1.0 | b |
| 4 |  |  | 151 | 2.9 | ab |
| 5 |  |  | 154 | 1.4 | ab |
| 6 |  | 145 | 429 | 1.4 | $a b$ |
| 7 |  |  | 430 | 1.0 | b |
| 8 |  |  | 431. | 1.4 | ab |
| 9 |  |  | 432 | 2.3 | $a b$ |
| 10 |  |  | 435 | 3.1 | ab |
| 11. | 2 | 97 | 260 | 1.4 | $a b$ |
| 12 |  |  | 261 | 3.1 | $a b$ |
| 13 |  |  | 262 | 2.0 | $a b$ |
| 14 |  |  | 264 | 1.7 | $a b$ |
| 15 |  |  | 266 | 2.2 | $a b$ |
| 16 |  | 131 | 369 | 3.6 | ab |
| 17 |  |  | 371 | 2.0 | ab |
| 18 |  |  | 372 | 2.1 | ab |
| 19 |  |  | 376 | 2.2 | ab |
| 20 |  |  | 377 | 1.7 | $a b$ |
| 21 | 3 | 15 | 36 | 1.8 | ab |
| 22 |  |  | 37 | 2.0 | $a b$ |
| 23 |  |  | 38 | 1.1 | b |
| 24 |  |  | 40 | 1.6 | $a b$ |
| 25 |  |  | 42 | 3.2 | $a b$ |
| 26 |  | 27 | 64 | 2.8 | $a b$ |
| 27 |  |  | 66 | 1.0 | b |
| 28 |  |  | 67 | 2.2 | $a b$ |
| 29 |  |  | 69 | 3.0 | $a b$ |
| 30 |  |  | 71 | 1.8 | $a b$ |
| 31 | 4 | 91 | 233 | 2.9 | ab |
| 32 |  |  | 234 | 2.3 | $a b$ |
| 33 |  |  | 235 | 2.9 | $a b$ |
| 34 |  |  | 236 | 3.2 | $a b$ |
| 35 |  |  | 237 | 3.4 | $a b$ |
| 36 |  | 154 | 460 | 2.0 | $a b$ |
| 37 |  |  | 461 | 2.7 | $a b$ |
| 38 |  |  | 462 | 1.7 | $a b$ |
| 39 |  |  | 463 | 1.9 | $a b$ |
| 40 |  |  | 465 | 2.8 | $a b$ |
| 41 | 5 | 13 | 25 | 2.0 | $a b$ |
| 42 |  |  | 26 | 2.3 | $a b$ |
| 43 |  |  | 27 | 2.1 | $a b$ |
| 44 |  |  | 31 | 2.2 | $a b$ |
| 45 |  |  | 32 | 4.0 | a |
| 46 |  | 142 | 414 | 2.0 | $a b$ |
| 47 |  |  | 415 | 1.7 | $a b$ |
| 48 |  |  | 416 | 1.6 | $a b$ |
| 49 |  |  | 418 | 2.2 | $a b$ |
| 50 |  |  | 420 | 1.9 | ab |

Means with the same letter are not significantly different at $5 \%$

Duncan's multiple range test for anthesis position


Means with the same letter are not significantly different at $5 \%$


[^3]

Duncan's multiple range test for panicle compactness

| No.Clus Topo Half-sib |  |  |  | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 6.4 | $a b c$ |
| 2 |  |  | 148 | 5.9 | $a b c$ |
| 3 |  |  | 150 | 6.8 | a |
| 4 |  |  | 151 | 6.3 | abc |
| 5 |  |  | 154 | 3.9 | bc |
| 6 |  | 145 | 429 | 5.1 | $a b c$ |
| 7 |  |  | 430 | 4.1 | $a b c$ |
| 8 |  |  | 431 | 6.4 | abc |
| 9 |  |  | 432 | 6.0 | $a b c$ |
| 10 |  |  | 435 | 6.0 | $a b c$ |
| 11 | 2 | 97 | 260 | 4.7 | abc |
| 12 |  |  | 261 | 5.6 | $a b c$ |
| 13 |  |  | 262 | 5.4 | $a b c$ |
| 14 |  |  | 264 | 4.9 | abc |
| 15 |  |  | 266 | 3.9 | bc |
| 16 |  | 131 | 369 | 6.3 | $a b c$ |
| 17 |  |  | 371 | 4.8 | abc |
| 18 |  |  | 372 | 3.8 | c |
| 19 |  |  | 376 | 4.8 | abc |
| 20 |  |  | 377 | 4.0 | abc |
| 21 | 3 | 15 | 36 | 4.8 | $a b c$ |
| 22 |  |  | 37 | 6.1 | $a b c$ |
| 23 |  |  | 38 | 6.7 | $a b$ |
| 24 |  |  | 40 | 5.3 | $a b c$ |
| 25 |  |  | 42 | 6.1 | $a b c$ |
| 26 |  | 27 | 64 | 6.0 | abc |
| 27 |  |  | 66 | 4.6 | $a b c$ |
| 28 |  |  | 67 | 5.9 | abc |
| 29 |  |  | 69 | 3.9 | bc |
| 30 |  |  | 71 | 5.5 | abc |
| 31 | 4 | 91 | 233 | 4.4 | $a b c$ |
| 32 |  |  | 234 | 6.0 | abc |
| 33 |  |  | 235 | 3.8 | c |
| 34 |  |  | 236 | 6.5 | abc |
| 35 |  |  | 237 | 4.6 | $a b c$ |
| 36 |  | 154 | 460 | 6.2 | abc |
| 37 |  |  | 461 | 4.8 | abc |
| 38 |  |  | 462 | 5.3 | $a b c$ |
| 39 |  |  | 463 | 5.3 | abc |
| 40 |  |  | 465 | 4.3 | $a b c$ |
| 41 | 5 | 13 | 25 | 6.2 | abc |
| 42 |  |  | 26 | 6.2 | abc |
| 43 |  |  | 27 | 6.3 | abc |
| 44 |  |  | 31 | 6.4 | abc |
| 45 |  |  | 32 | 5.0 | $a b c$ |
| 46 |  | 142 | 414 | 6.3 | $a b c$ |
| 47 |  |  | 415 | 4.9 | $a b c$ |
| 48 |  |  | 416 | 5.4 | $a b c$ |
| 49 |  |  | 418 | 6.2 | $a b c$ |
| 50 |  |  | 420 | 5.3 | abc |

Means with the same letter are not significantly different at $5 \%$


Duncan's multiple range test for old disease


Duncan's multiple range test for new diseases


Means with the same letter are not significantly different at $5 \%$

| No.Clus |  | Topo69 | $\begin{aligned} & \text { Half-sib } \\ & 147 \end{aligned}$ | Means |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 |  |  | 9.1 | $a b c$ |  |
| 2 |  |  | 148 | 8.8 | abcd |  |
| 3 |  |  | 150 | 8.2 | abcd |  |
| 4 |  |  | 151 | 8.9 | abcd |  |
| 5 |  |  | 154 | 8.3 | abcd |  |
| 6 |  | 145 | 429 | 9.1 | abcd |  |
| 7 |  |  | 430 | 8.8 | abcd |  |
| 8 |  |  | 431 | 9.2 | abc |  |
| 9 |  |  | 432 | 8.6 | abcd |  |
| 10 |  |  | 435 | 8.3 | abcd |  |
| 11 | 2 | 97 | 260 | 9.1 | abcd |  |
| 12 |  |  | 261 | 8.9 | abcd |  |
| 13 |  |  | 262 | 8.9 | abcd |  |
| 14 |  |  | 264 | 9.6 | a |  |
| 15 |  |  | 266 | 9.6 | $a b$ |  |
| 16 |  | 131 | 369 | 8.8 | abcd |  |
| 17 |  |  | 371 | 8.8 | abcd |  |
| 18 |  |  | 372 | 8.7 | abcd |  |
| 19 |  |  | 376 | 9.0 | abcd |  |
| 20 |  |  | 377 | 9.4 | abc |  |
| 21 | 3 | 15 | 36 | 8.6 | abcd |  |
| 22 |  |  | 37 | 7.9 | cd |  |
| 23 |  |  | 38 | 8.7 | abcd |  |
| 24 |  |  | 40 | 9.3 | abc |  |
| 25 |  |  | 42 | 8.4 | abcd |  |
| 26 |  | 27 | 64 | 8.8 | abod |  |
| 27 |  |  | 66 | 9.1 | abcd |  |
| 28 |  |  | 67 | 8.9 | abcd |  |
| 29 |  |  | 69 | 9.3 | abc |  |
| 30 |  |  | 71 | 7.6 | d |  |
| 31 | 4 | 91 | 233 | 8.3 | abcd |  |
| 32 |  |  | 234 | 8.3 | abcd |  |
| 33 |  |  | 235 | 8.8 | abcd |  |
| 34 |  |  | 236 | 9.0 | abcd |  |
| 35 |  |  | 237 | 9.4 | $a b c$ |  |
| 36 |  | 154 | 460 | 8.4 | abcd |  |
| 37 |  |  | 461 | 8.7 | abcd |  |
| 38 |  |  | 462 | 9.1 | abcd |  |
| 39 |  |  | 463 | 8.0 | bcd |  |
| 40 |  |  | 465 | 8.6 | abcd |  |
| 41 | 5 | 13 | 25 | 9.1 | abcd |  |
| 42 |  |  | 26 | 8.7 | abcd |  |
| 43 |  |  | 27 | 8.4 | abcd |  |
| 44 |  |  | 31 | 8.2 | abcd |  |
| 45 |  |  | 32 | 8.3 | abcd |  |
| 46 |  | 142 | 414 | 8.4 | abcd |  |
| 47 |  |  | 415 | 8.8 | abcd |  |
| 48 |  |  | 416 | 8.3 | abcd |  |
| 49 |  |  | 418 | 8.9 | abcd |  |
| 50 |  |  | 420 | 8.4 | abcd |  |



Duncan's multiple range test for flavanoid leaf sheath



Duncan's multiple range test for tiller number at $50 \%$ upper asymptote


Means with the same letter are not significantly different at $5 \%$


Means with the same letter are not significantly different at 5\%

```
Duncan's multiple range test for tiller number at flowering
\begin{tabular}{|c|c|c|c|c|c|}
\hline No. & Clus & us Topo & Half-sib & Means & \\
\hline 1 & 1 & 69 & 147 & 34.7 & defgh \\
\hline 2 & & & 148 & 32.9 & efgh \\
\hline 3 & & & 150 & 40.3 & bcdefgh \\
\hline 4 & & & 151 & 62.8 & a \\
\hline 5 & & & 154 & 36.2 & cdefgh \\
\hline 6 & & 145 & 429 & 43.8 & abcdefgh \\
\hline 7 & & & 430 & 32.1 & gh \\
\hline 8 & & & 431 & 47.2 & abcdefgh \\
\hline 9 & & & 432 & 52.8 & abcde \\
\hline 10 & & & 435 & 39.9 & bcdefgh \\
\hline 11 & 2 & 97 & 260 & 39.8 & bcdefgh \\
\hline 12 & & & 261 & 44.0 & abcdefgh \\
\hline 13 & & & 262 & 38.0 & cdefgh \\
\hline 14 & & & 264 & 51.7 & abcdefgh \\
\hline 15 & & & 266 & 50.4 & abcdefgh \\
\hline 16 & & 131 & 369 & 33.6 & defgh \\
\hline 17 & & & 371 & 32.0 & fgh \\
\hline 18 & & & 372 & 37.1 & cdefgh \\
\hline 19 & & & 376 & 33.3 & defgh \\
\hline 20 & & & 377 & 43.0 & bcdefgh \\
\hline 21 & 3 & 15 & 36 & 44.7 & abcdefgh \\
\hline 22 & & & 37 & 35.7 & cdefgh \\
\hline 23 & & & 38 & 35.7 & cdefgh \\
\hline 24 & & & 40 & 30.4 & h \\
\hline 25 & & & 42 & 49.4 & abcdefgh \\
\hline 26 & & 27 & 64 & 32.4 & fgh \\
\hline 27 & & & 66 & 30.5 & h \\
\hline 28 & & & 67 & 55.8 & \(a b c\) \\
\hline 29 & & & 69 & 51.0 & abcdefg \\
\hline 30 & & & 71 & 46.4 & abcdefgh \\
\hline 31 & 4 & & 233 & 46.1 & abcdefgh \\
\hline 32 & & & 234 & 52.3 & abcdef \\
\hline 33 & & & 235 & 32.4 & fgh \\
\hline 34 & & & 236 & 34.7 & defgh \\
\hline 35 & & & 237 & 35.6 & defgh \\
\hline 36 & & 154 & 460 & 36.0 & cdefgh \\
\hline 37 & & & 461 & 43.6 & abcdefgh \\
\hline 38 & & & 462 & 46.4 & abcdefgh \\
\hline 39 & & & 463 & 37.0 & cdefgh \\
\hline 40 & & & 465 & 45.9 & abcdefgh \\
\hline 41 & 5 & 13 & 25 & 43.0 & bcdefgh \\
\hline 42 & & & 26 & 29.9 & h \\
\hline 43 & & & 27 & 35.2 & defgh \\
\hline 44 & & & 31 & 41.7 & bcdefgh \\
\hline 45 & & & 32 & 36.7 & cdefgh \\
\hline 46 & & 142 & 414 & 44.3 & abcdefgh \\
\hline 47 & & & 415 & 35.0 & defgh \\
\hline 48 & & & 416 & 49.1 & abcdefgh \\
\hline 49 & & & 418 & 53.1 & abcd \\
\hline 50 & & & 420 & 52.6 & abcde \\
\hline
\end{tabular}
```

[^4]

| Duncan' <br> No. Clus |  | S Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 69 |  | 30.3 | ab |
| 2 |  |  | 148 | 22.8 | $a b$ |
| 3 |  |  | 150 | 32.7 | $a b$ |
| 4 |  |  | 151 | 27.4 | $a b$ |
| 5 |  |  | 154 | 25.3 | $a b$ |
| 6 |  | 145 | 429 | 20.6 | $a b$ |
| 7 |  |  | 430 | 29.7 | $a b$ |
| 8 |  |  | 431 | 24.1 | $a b$ |
| 9 |  |  | 432 | 33.1 | $a b$ |
| 10 |  |  | 435 | 19.5 | b |
| 11 | 2 | 97 | 260 | 27.3 | ab |
| 12 |  |  | 261 | 26.9 | $a b$ |
| 13 |  |  | 262 | 22.2 | $a b$ |
| 14 |  |  | 264 | 31.0 | $a b$ |
| 15 |  |  | 266 | 26.7 | $a b$ |
| 16 |  | 131 | 369 | 21.2 | $a b$ |
| 17 |  |  | 371 | 30.1 | $a b$ |
| 18 |  |  | 372 | 30.8 | $a b$ |
| 19 |  |  | 376 | 26.4 | $a b$ |
| 20 |  |  | 377 | 26.7 | $a b$ |
| 21 | 3 | 15 | 36 | 34.4 | $a b$ |
| 22 |  |  | 37 | 36.3 | $a b$ |
| 23 |  |  | 38 | 33.0 | $a b$ |
| 24 |  |  | 40 | 22.2 | $a b$ |
| 25 |  |  | 42 | 36.3 | $a b$ |
| 26 |  | 27 | 64 | 18.0 | b |
| 27 |  |  | 66 | 26.7 | $a b$ |
| 28 |  |  | 67 | 35.8 | $a b$ |
| 29 |  |  | 69 | 27.1 | $a b$ |
| 30 |  |  | 71 | 25.1 | $a b$ |
| 31 | 4 | 91 | 233 | 28.2 | $a b$ |
| 32 |  |  | 234 | 33.2 | $a b$ |
| 33 |  |  | 235 | 41.3 | $a$ |
| 34 |  |  | 236 | 28.8 | $a b$ |
| 35 |  |  | 237 | 25.4 | $a b$ |
| 36 |  | 154 | 460 | 28.3 | $a b$ |
| 37 |  |  | 461 | 32.0 | $a b$ |
| 38 |  |  | 462 | 21.0 | $a b$ |
| 39 |  |  | 463 | 27.4 | $a b$ |
| 40 |  |  | 465 | 35.1 | $a b$ |
| 41 | 5 | 13 | 25 | 20.2 | $a b$ |
| 42 |  |  | 26 | 31.8 | $a b$ |
| 43 |  |  | 27 | 26.4 | $a b$ |
| 44 |  |  | 31 | 20.8 | $a b$ |
| 45 |  |  | 32 | 28.9 | $a b$ |
| 46 |  | 142 | 414 | 26.6 | $a b$ |
| 47 |  |  | 415 | 36.9 | $a b$ |
| 48 |  |  | 416 | 36.3 | $a b$ |
| 49 |  |  | 418 | 22.9 | $a b$ |
| 50 |  |  | 420 | 32.1 | $a b$ |

Means with the same letter are not significantly different at 5\%
$\left.\begin{array}{ccccc}\text { Duncan's multiple range test } & \text { for } & \text { young } & \text { tiller number } \\ \text { No. Clus Topo Half-sib } & \\ 1 & 1 & 69 & 147 & \text { Means }\end{array}\right]$


| No.Clus |  | S Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 53.4 | cde |
| 2 |  |  | 148 | 47.9 | cde |
| 3 |  |  | 150 | 68.5 | abcde |
| 4 |  |  | 151 | 45.8 | de |
| 5 |  |  | 154 | 68.0 | abcde |
| 6 |  | 145 | 429 | 44.5 | de |
| 7 |  |  | 430 | 51.9 | cde |
| 8 |  |  | 431 | 51.8 | cde |
| 9 |  |  | 432 | 60.4 | abcde |
| 10 |  |  | 435 | 45.8 | de |
| 11 | 2 | 97 | 260 | 54.9 | cde |
| 12 |  |  | 261 | 59.6 | abcde |
| 13 |  |  | 262 | 54.0 | cde |
| 14 |  |  | 264 | 60.0 | abcde |
| 15 |  |  | 266 | 56.3 | cde |
| 16 |  | 131 | 369 | 47.1 | de |
| 17 |  |  | 371 | 64.2 | abcde |
| 18 |  |  | 372 | 65.2 | abcde |
| 19 |  |  | 376 | 49.8 | cde |
| 20 |  |  | 377 | 57.7 | bcde |
| 21 | 3 | 15 | 36 | 81.9 | abc |
| 22 |  |  | 37 | 52.7 | cde |
| 23 |  |  | 38 | 63.8 | abcde |
| 24 |  |  | 40 | 47.9 | cde |
| 25 |  |  | 42 | 91.2 | a |
| 26 |  | 27 | 64 | 34.7 | e |
| 27 |  |  | 66 | 45.3 | de |
| 28 |  |  | 67 | 74.1 | abcde |
| 29 |  |  | 69 | 63.9 | abcde |
| 30 |  |  | 71 | 64.9 | abcde |
| 31 | 4 | 91 | 233 | 57.8 | bcde |
| 32 |  |  | 234 | 64.3 | abcde |
| 33 |  |  | 235 | 76.4 | abcd |
| 34 |  |  | 236 | 57.4 | bode |
| 35 |  |  | 237 | 54.8 | ccie |
| 36 |  | 1.54 | 460 | 55.1 | cde |
| 37 |  |  | 461 | 63.1 | abcde |
| 38 |  |  | 462 | 49.4 | cde |
| 39 |  |  | 463 | 62.0 | abcde |
| 40 |  |  | 465 | 89.0 | $a b$ |
| 41 | 5 | 13 | 25 | 43.6 | de |
| 42 |  |  | 26 | 51.4 | cde |
| 43 |  |  | 27 | 48.8 | cde |
| 44 |  |  | 31 | 47.6 | de |
| 45 |  |  | 32 | 55.0 | cde |
| 46 |  | 142 | 414 | 53.8 | cde |
| 47 |  |  | 415 | 67.8 | abcde |
| 48 |  |  | 416 | 61.4 | abcde |
| 49 |  |  | 418 | 50.7 | cde |
| 50 |  |  | 420 | 59.2 | abcde |

Means with the same letter are not significantly different at $5 \%$

Duncan's multiple range test for base green tiller number

| No. | Clu | as Topo | Half-sib | Means |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 43.3 | abcdef |  |
| 2 |  |  | 148 | 32.6 | bcdef |  |
| 3 |  |  | 150 | 59.5 | $a b c$ |  |
| 4 |  |  | 151 | 31.1 | cdef |  |
| 5 |  |  | 154 | 42.9 | abcdef |  |
| 6 |  | 145 | 429 | 28.6 | def |  |
| 7 |  |  | 430 | 38.3 | abcdef |  |
| 8 |  |  | 431 | 31.6 | bcdef |  |
| 9 |  |  | 432 | 42.6 | abcdef |  |
| 10 |  |  | 435 | 29.5 | def |  |
| 11 | 2 | 97 | 260 | 38.1 | abcdef |  |
| 12 |  |  | 261 | 35.4 | bcdef |  |
| 13 |  |  | 262 | 30.2 | cdef |  |
| 14 |  |  | 264 | 37.1 | abcdef |  |
| 15 |  |  | 266 | 32.9 | bcdef |  |
| 16 |  | 131 | 369 | 32.3 | bodef |  |
| 17 |  |  | 371 | 48.1 | abcdef |  |
| 18 |  |  | 372 | 41.7 | abcdef |  |
| 19 |  |  | 376 | 35.9 | abcdef |  |
| 20 |  |  | 377 | 34.4 | bcdef |  |
| 21 | 3 | 15 | 36 | 52.2 | abcdef |  |
| 22 |  |  | 37 | 38.8 | abcdef |  |
| 23 |  |  | 38 | 47.0 | abcdef |  |
| 24 |  |  | 40 | 35.7 | abcdef |  |
| 25 |  |  | 42 | 60.7 | $a b$ |  |
| 26 |  | 27 | 64 | 21.3 | f |  |
| 27 |  |  | 66 | 36.0 | abcdef |  |
| 28 |  |  | 67 | 55.4 | abcdef |  |
| 29 |  |  | 69 | 41.3 | abcdef |  |
| 30 |  |  | 71 | 41.4 | abcdef |  |
| 31 | 4 | 91 | 233 | 37.8 | abcdef |  |
| 32 |  |  | 234 | 44.7 | abcdef |  |
| 33 |  |  | 235 | 57.9 | abcd |  |
| 34 |  |  | 236 | 41.2 | abcdef |  |
| 35 |  |  | 237 | 34.3 | bcdef |  |
| 36 |  | 154 | 460 | 40.1 | abcdef |  |
| 37 |  |  | 461 | 41.4 | abcdef |  |
| 38 |  |  | 462 | 31.6 | bcdef | . |
| 39 |  |  | 463 | 44.1 | abcdef |  |
| 40 |  |  | 465 | 64.9 | a |  |
| 41 | 5 |  | 25 | 28.6 | def |  |
| 42 |  |  | 26 | 42.8 | abcdef |  |
| 43 |  |  | 27 | 33.2 | bedef |  |
| 44 |  |  | 31 | 27.6 | ef |  |
| 45 |  |  | 32 | 43.4 | abcdef |  |
| 46 |  | 142 | 414 | 39.4 | abcdef |  |
| 47 |  |  | 415 | 53.4 | abcdef |  |
| 48 |  |  | 416 | 44.1 | abcdef |  |
| 49 |  |  | 418 | 26.8 | ef |  |
| 50 |  |  | 420 | 38.9 | abcdef |  |

Means with the same letter are not significantly different at 5\%


Duncan's multiple range test for dead tiller dry weight


[^5]

```
Duncan's multiple range test for young tiller dry weight
\begin{tabular}{|c|c|c|c|c|c|}
\hline No. & Clus & S Topres & Half-sib & Mea & \\
\hline 1 & 1 & 69 & 147 & 0.9 & cdefg \\
\hline 2 & & & 148 & 1.4 & bcdefg \\
\hline 3 & & & 150 & 2.2 & abcdefg \\
\hline 4 & & & 151 & 0.3 & g \\
\hline 5 & & & 154 & 2.2 & abcdefg \\
\hline 6 & & 145 & 429 & 0.9 & cdef \\
\hline 7 & & & 430 & 0.8 & defg \\
\hline 8 & & & 431 & 1.0 & bcdefg \\
\hline 9 & & & 432 & 1.4 & bcdefg \\
\hline 10 & & & 435 & 1.4 & bcdefg \\
\hline 11 & 2 & 97 & 260 & 0.9 & cdefg \\
\hline 12 & & & 261 & 0.9 & cdefg \\
\hline 13 & & & 262 & 1.0 & bcdefg \\
\hline 14 & & & 264 & 0.7 & defg \\
\hline 15 & & & 266 & 0.9 & cdefg \\
\hline 16 & & 131 & 369 & 0.9 & cdefg \\
\hline 17 & & & 371 & 2.3 & abcde \\
\hline 18 & & & 372 & 1.6 & bcdefg \\
\hline 19 & & & 376 & 0.7 & defg \\
\hline 20 & & & 377 & 0.7 & efg \\
\hline 21 & 3 & 15 & 36 & 2.0 & abcdefg \\
\hline 22 & & & 37 & 0.2 & \(g\) \\
\hline 23 & & & 38 & 1.2 & bcdefg \\
\hline 24 & & & 40 & 1.6 & bcdefg \\
\hline 25 & & & 42 & 3.6 & a \\
\hline 26 & & 27 & 64 & 0.3 & fg \\
\hline 27 & & & 66 & 0.9 & cdefg \\
\hline 28 & & & 67 & 2.9 & \(a b\) \\
\hline 29 & & & 69 & 1.9 & abcdefg \\
\hline 30 & & & 71 & 2.0 & abcdefg \\
\hline 31 & 4 & 91 & 233 & 1.7 & bcdefg \\
\hline 32 & & & 234 & 1.4 & bcdefg \\
\hline 33 & & & 235 & 2.7 & abcde \\
\hline 34 & & & 236 & 1.0 & bcdefg \\
\hline 35 & & & 237 & 0.9 & cdefg \\
\hline 36 & & 154 & 460 & 1.2 & bcdefg \\
\hline 37 & & & 461 & 0.9 & cdefg \\
\hline 38 & & & 462 & 1.1 & bcdefg \\
\hline 39 & & & 463 & 2.8 & abc \\
\hline 40 & & & 465 & 1.5 & bcdefg \\
\hline 41 & 5 & 13 & 25 & 0.9 & cdef \\
\hline 42 & & & 26 & 1.3 & bcdef \\
\hline 43 & & & 27 & 0.7 & def \\
\hline 44 & & & 31 & 0.9 & cdef \\
\hline 45 & & & 32 & 2.3 & abcdef \\
\hline 46 & & 142 & 414 & 1.2 & bcdef \\
\hline 47 & & & 415 & 2.7 & abcd \\
\hline 48 & & & 416 & 0.6 & efg \\
\hline 49 & & & 418 & 0.5 & efg \\
\hline 50 & & & 420 & 0.8 & cdefg \\
\hline
\end{tabular}
```

[^6]
## Duncan's multiple range test for base green tiller dryweight

| No. | clus | Is Topo | Half-sib | Means |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 21.5 | abcdef |  |
| 2 |  |  | 148 | 19.0 | bcdef |  |
| 3 |  |  | 150 | 25.6 | abcdef |  |
| 4 |  |  | 151 | 12.8 | f |  |
| 5 |  |  | 154 | 20.3 | abcdef |  |
| 6 |  | 145 | 429 | 14.3 | ef |  |
| 7 |  |  | 430 | 18.2 | cdef |  |
| 8 |  |  | 431 | 16.7 | def |  |
| 9 |  |  | 432 | 22.2 | abcdef |  |
| 10 |  |  | 435 | 16.1 | def |  |
| 11 | 2 | 97 | 260 | 14.9 | ef |  |
| 12 |  |  | 261 | 22.2 | abcdef |  |
| 13 |  |  | 262 | 14.3 | ef |  |
| 14 |  |  | 264 | 19.5 | bcdef |  |
| 15 |  |  | 266 | 17.0 | def |  |
| 16 |  | 131 | 369 | 12.6 | f |  |
| 17 |  |  | 371 | 23.0 | abcdef |  |
| 18 |  |  | 372 | 21.6 | abcdef |  |
| 19 |  |  | 376 | 18.9 | bcdef |  |
| 20 |  |  | 377 | 25.5 | abcdef |  |
| 21 | 3 | 15 | 36 | 32.1 | abcd |  |
| 22 |  |  | 37 | 35.9 | a |  |
| 23 |  |  | 38 | 21.2 | abcdef |  |
| 24 |  |  | 40 | 17.4 | cdef |  |
| 25 |  |  | 42 | 24.7 | abcdef |  |
| 26 |  | 27 | 64 | 16.0 | def |  |
| 27 |  |  | 66 | 23.2 | abcdef |  |
| 28 |  |  | 67 | 21.6 | abcdef |  |
| 29 |  |  | 69 | 22.3 | abcdef |  |
| 30 |  |  | 71 | 18.8 | bcdef |  |
| 31 | 4 | 91 | 233 | 18.3 | cdef |  |
| 32 |  |  | 234 | 22.4 | abcdef |  |
| 33 |  |  | 235 | 33.7 | abc |  |
| 34 |  |  | 236 | 22.8 | abcdef |  |
| 35 |  |  | 237 | 13.5 | f |  |
| 36 |  | 154 | 460 | 15.3 | ef |  |
| 37 |  |  | 461 | 20.8 | abcdef |  |
| 38 |  |  | 462 | 14.0 | f |  |
| 39 |  |  | 463 | 35.0 | $a b$ |  |
| 40 |  |  | 465 | 30.8 | abcde |  |
| 41 | 5 | 13 | 25 | 14.8 | ef |  |
| 42 |  |  | 26 | 28.1 | abcdef |  |
| 43 |  |  | 27 | 19.7 | abcdef |  |
| 44 |  |  | 31 | 17.4 | cdef |  |
| 45 |  |  | 32 | 22.9 | abcdef |  |
| 46 |  | 142 | 414 | 13.3 | f |  |
| 47 |  |  | 415 | 32.0 | abcdef |  |
| 48 |  |  | 416 | 20.2 | abcdef |  |
| 49 |  |  | 418 | 16.6 | def |  |
| 50 |  |  | 420 | 21.7 | abcdef |  |


| No. | Clus | us Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 21.3 | bode |
| 2 |  |  | 148 | 21.6 | cde |
| 3 |  |  | 150 | 33.3 | abcde |
| 4 |  |  | 151 | 26.2 | bcde |
| 5 |  |  | 154 | 26.0 | bcde |
| 6 |  | 145 | 429 | 29.1 | abcde |
| 7 |  |  | 430 | 26.7 | bcde |
| 8 |  |  | 431 | 24.6 | bode |
| 9 |  |  | 432 | 33.7 | abcde |
| 10 |  |  | 435 | 27.5 | abcde |
| 11 | 2 | 97 | 260 | 18.0 | de |
| 12 |  |  | 261 | 22.4 | bcde |
| 13 |  |  | 262 | 28.3 | abcde |
| 14 |  |  | 264 | 29.7 | abcde |
| 15 |  |  | 266 | 27.0 | bode |
| 16 |  | 131 | 369 | 28.1 | abcde |
| 17 |  |  | 371 | 26.2 | bcde |
| 18 |  |  | 372 | 26.9 | bcde |
| 1.9 |  |  | 376 | 30.4 | abcde |
| 20 |  |  | 377 | 33.6 | abcde |
| 21 | 3 | 15 | 36 | 23.7 | bcde |
| 22 |  |  | 37 | 23.7 | bcde |
| 23 |  |  | 38 | 28.6 | abcde |
| 24 |  |  | 40 | 21.7 | bcde |
| 25 |  |  | 42 | 29.4 | abcde |
| 26 |  | 27 | 64 | 36.9 | $a b c$ |
| 27 |  |  | 66 | 32.5 | abcde |
| 28 |  |  | 67 | 26.4 | bcde |
| 29 |  |  | 69 | 26.3 | bode |
| 30 |  |  | 71 | 36.3 | abcd |
| 31 | 4 |  | 233 | 29.7 | abcde |
| 32 |  |  | 234 | 38.1 | $a b$ |
| 33 |  |  | 235 | 20.3 | bcde |
| 34 |  |  | 236 | 15.8 | e |
| 35 |  |  | 237 | 32.8 | abcde |
| 36 |  | 154 | 460 | 23.5 | bcde |
| 37 |  |  | 461 | 23.1 | bcde |
| 38 |  |  | 462 | 26.8 | bcde |
| 39 |  |  | 463 | 26.4 | bcde |
| 40 |  |  | 465 | 26.0 | bcde |
| 41 | 5 | 13 | 25 | 33.7 | abcde |
| 42 |  |  | 26 | 18.8 | cde |
| 43 |  |  | 27 | 29.0 | abcde |
| 44 |  |  | 31 | 35.0 | abcd |
| 45 |  |  | 32 | 35.8 | abcd |
| 46 |  | 142 | 414 | 21.5 | bcde |
| 47 |  |  | 415 | 31.0 | abcde |
| 48 |  |  | 416 | 21.0 | bcde |
| 49 |  |  | 418 | 45.4 | a |
| 50 |  |  | 420 | 30.8 | abcde |

Means with the same letter are not significantly different at $5 \%$

| No.Clus |  | Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 24.3 | cdef |
| 2 |  |  | 148 | 22.2 | def |
| 3 |  |  | 150 | 30.7 | abcdef |
| 4 |  |  | 151 | 19.9 | def |
| 5 |  |  | 154 | 30.8 | abcdef |
| 6 |  | 145 | 429 | 21.1 | def |
| 7 |  |  | 430 | 24.1 | cdef |
| 8 |  |  | 431 | 25.7 | cdef |
| 9 |  |  | 432 | 30.3 | abcdef |
| 10 |  |  | 435 | 22.8 | def |
| 11 | 2 | 97 | 260 | 20.1 | def |
| 12 |  |  | 261 | 46.4 | abcdef |
| 13 |  |  | 262 | 26.0 | cdef |
| 14 |  |  | 264 | 29.2 | abcdef |
| 15 |  |  | 266 | 26.6 | bcdef |
| 16 |  | 131 | 369 | 17.9 | def |
| 17 |  |  | 371 | 31.1 | abcdef |
| 18 |  |  | 372 | 34.2 | abcdef |
| 19 |  |  | 376 | 25.3 | cdef |
| 20 |  |  | 377 | 37.3 | abcdef |
| 21 | 3 | 15 | 36 | 46.5 | a |
| 22 |  |  | 37 | 43.1 | $a b c$ |
| 23 |  |  | 38 | 28.5 | abcdef |
| 24 |  |  | 40 | 22.6 | def |
| 25 |  |  | 42 | 36.7 | abcdef |
| 26 |  | 27 | 64 | 23.3 | cdef |
| 27 |  |  | 66 | 28.2 | abcdef |
| 28 |  |  | 67 | 30.6 | abcdef |
| 29 |  |  | 69 | 35.5 | abcdef |
| 30 |  |  | 71 | 31.7 | abcdef |
| 31 | 4 | 91 | 233 | 26.9 | abcdef |
| 32 |  |  | 234 | 31.1 | abcdef |
| 33 |  |  | 235 | 41.6 | abcd |
| 34 |  |  | 236 | 29.3 | abcdef |
| 35 |  |  | 237 | 20.9 | def |
| 36 |  | 154 | 460 | 21.6 | def |
| 37 |  |  | 461 | 35.8 | abcdef |
| 38 |  |  | 462 | 22.3 | def |
| 39 |  |  | 463 | 42.9 | $a b c$ |
| 40 |  |  | 465 | 46.2 | $a b$ |
| 41 | 5 | 13 | 25 | 21.9 | def |
| 42 |  |  | 26 | 32.1 | abcdef |
| 43 |  |  | 27 | 27.4 | abcdef |
| 44 |  |  | 31 | 27.3 | abcdef |
| 45 |  |  | 32 | 28.6 | abcdef |
| 46 |  | 142 | 414 | 18.3 | def |
| 47 |  |  | 415 | 40.5 | abcde |
| 48 |  |  | 416 | 29.9 | abcdef |
| 49 |  |  | 418 | 27.3 | abcdef |
| 50 |  |  | 420 | 31.6 | abcdef |

Means with the same letter are not significantly different at $5 \%$

```
Duncan's multiple range test for total tiller dry weight
\begin{tabular}{|c|c|c|c|c|c|}
\hline No. & Clus & S Topo & Half-sib & Means & \\
\hline 1 & 1 & 69 & 147 & 45.6 & bcd \\
\hline 2 & & & 148 & 47.0 & bod \\
\hline 3 & & & 150 & 64.0 & abcd \\
\hline 4 & & & 151 & 46.0 & bcd \\
\hline 5 & & & 154 & - 56.8 & abcd \\
\hline 6 & & 145 & 429 & 50.2 & abcd \\
\hline 7 & & & 430 & 50.8 & abod \\
\hline 8 & & & 431 & 50.2 & abcd \\
\hline 9 & & & 432 & 64.0 & abcd \\
\hline 10 & & & 435 & 50.4 & abcd \\
\hline 11 & 2 & 97 & 260 & 38.1 & d \\
\hline 12 & & & 261 & 58.8 & abcd \\
\hline 13 & & & 262 & 54.2 & abcd \\
\hline 14 & & & 264 & 58.9 & abcd \\
\hline 15 & & & 266 & 53.6 & abcd \\
\hline 16 & & 131 & 369 & 46.1 & bed \\
\hline 17 & & & 371 & 57.3 & abcd \\
\hline 18 & & & 372 & 61.1 & \(a b c d\) \\
\hline 19 & & & 376 & 55.7 & abcd \\
\hline 20 & & & 377 & 70.9 & \(a b\) \\
\hline 21 & 3 & 15 & 36 & 70.2 & \(a b\) \\
\hline 22 & & & 37 & 66.9 & abcd \\
\hline 23 & & & 38 & 57.1 & abcd \\
\hline 24 & & & 40 & 44.3 & bcd \\
\hline 25 & & & 42 & 66.1 & abcd \\
\hline 26 & & 27 & 64 & 60.3 & abcd \\
\hline 27 & & & 66 & 51.2 & abcd \\
\hline 28 & & & 67 & 52.2 & abcd \\
\hline 29 & & & 69 & 54.9 & abcd \\
\hline 30 & & & 71 & 60.9 & abcd \\
\hline 31 & 4 & 91 & 233 & 56.6 & abcd \\
\hline 32 & & & 234 & 69.2 & \(a b c\) \\
\hline 33 & & & 235 & 62.0 & abcd \\
\hline 34 & & & 236 & 42.7 & bcd \\
\hline 35 & & & 237 & 55.8 & abcd \\
\hline 36 & & 154 & 460 & 45.1 & bcd \\
\hline 37 & & & 461 & 58.9 & abcd \\
\hline 38 & & & 462 & 49.2 & abcd \\
\hline 39 & & & 463 & 69.2 & \(a b c\) \\
\hline 40 & & & 465 & 78.2 & a \\
\hline 41 & 5 & 13 & 25 & 55.6 & abcd \\
\hline 42 & & & 26 & 50.9 & abcd \\
\hline 43 & & & 27 & 56.4 & abcd \\
\hline 44 & & & 31 & 62.3 & abcd \\
\hline 45 & & & 32 & 64.4 & abcd \\
\hline 46 & & 142 & 414 & 38.8 & cd \\
\hline 47 & & & 415 & 71.4 & \(a b\) \\
\hline 48 & & & 416 & 50.8 & abcd \\
\hline 49 & & & 418 & 72.7 & \(a b\) \\
\hline 50 & & & 420 & 62.5 & abcd \\
\hline
\end{tabular}
```

[^7]Duncan's multiple range test for number of days to reach $5 \%$ asymptote


Duncan's multiple range test for number of days to reach $50 \%$ upper asymptote

| No. | Clus | ( Topo | Half-sib | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 61.6 | bcde |
| 2 |  |  | 148 | 54.4 | e |
| 3 |  |  | 150 | 67.1 | bc |
| 4 |  |  | 151 | 65.4 | bcde |
| 5 |  |  | 154 | 62.2 | bcde |
| 6 |  | 145 | 429 | 62.8 | bcde |
| 7 |  |  | 430 | 59.3 | cde |
| 8 |  |  | 431 | 66.2 | bod |
| 9 |  |  | 432 | 65.6 | bod |
| 10 |  |  | 435 | 60.0 | cde |
| 11 | 2 | 97 | 260 | 63.1 | bcde |
| 12 |  |  | 261 | 69.4 | $a b c$ |
| 13 |  |  | 262 | 64.5 | bcde |
| 14 |  |  | 264 | 66.6 | bc |
| 15 |  |  | 266 | 64.5 | bcde |
| 16 |  | 131 | 369 | 61.8 | bode |
| 17 |  |  | 371 | 63.5 | bcde |
| 18 |  |  | 372 | 64.1 | bcde |
| 19 |  |  | 376 | 60.3 | bcde |
| 20 |  |  | 377 | 62.6 | bcde |
| 21 | 3 | 15 | 36 | 77.8 | a |
| 22 |  |  | 37 | 69.0 | abc |
| 23 |  |  | 38 | 61.5 | bcde |
| 24 |  |  | 40 | 60.9 | bcde |
| 25 |  |  | 42 | 67.7 | bc |
| 26 |  | 27 | 64 | 59.6 | cde |
| 27 |  |  | 66 | 60.9 | bcde |
| 28 |  |  | 67 | 68.7 | abc |
| 29 |  |  | 69 | 67.3 | $b c$ |
| 30 |  |  | 71 | 67.7 | bc |
| 31 | 4 | 91 | 233 | 60.4 | bcde |
| 32 |  |  | 234 | 63.6 | bcde |
| 33 |  |  | 235 | 61.4 | bcde |
| 34 |  |  | 236 | 61.3 | bcde |
| 35 |  |  | 237 | 61.0 | bode |
| 36 |  | 154 | 460 | 64.8 | bcde |
| 37 |  |  | 461 | 71.3 | $a b$ |
| 38 |  |  | 462 | 64.1 | bcde |
| 39 |  |  | 463 | 65.2 | bcde |
| 40 |  |  | 465 | 65.1 | bcde |
| 41 | 5 | 13 | 25 | 62.2 | bcde |
| 42 |  |  | 26 | 62.2 | bcde |
| 43 |  |  | 27 | 61.4 | bcde |
| 44 |  |  | 31 | 60.2 | cde |
| 45 |  |  | 32 | 61.1 | bcde |
| 46 |  | 142 | 414 | 55.5 | de |
| 47 |  |  | 415 | 58.5 | cde |
| 48 |  |  | 416 | 65.8 | bcd |
| 49 |  |  | 418 | 65.1 | bode |
| 50 |  |  | 420 | 66.2 | bcd |

Duncan's multiple range test for number of days to reach $95 \%$ upper asymptote

| No.Clus Topo Half-sib |  |  |  | Means |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 69 | 147 | 97.5 | bcdef |
| 2 |  |  | 148 | 83.9 | f |
| 3 |  |  | 150. | 99.3 | bcdef |
| 4 |  |  | 151 | 102.5 | bcde |
| 5 |  |  | 154 | 92.2 | cdef |
| 6 |  | 145 | 429 | 97.8 | bcdef |
| 7 |  |  | 430 | 92.3 | cdef |
| 8 |  |  | 431 | 104.5 | abcde |
| 9 |  |  | 432 | 103.8 | abcde |
| 10 |  |  | 435 | 96.0 | bcdef |
| 11 | 2 | 97 | 260 | 98.4 | bcdef |
| 12 |  |  | 261 | 109.7 | $a b c$ |
| 13 |  |  | 262 | 98.6 | bcdef |
| 14 |  |  | 264 | 105.1 | abcde |
| 15 |  |  | 266 | 101.6 | bcdef |
| 16 |  | 131 | 369 | 93.7 | cdef |
| 17 |  |  | 371 | 94.9 | cdef |
| 18 |  |  | 372 | 97.0 | bodef |
| 19 |  |  | 376 | 91.3 | def |
| 20 |  |  | 377 | 96.8 | bcdef |
| 21 | 3 | 15 | 36 | 120.6 | a |
| 22 |  |  | 37 | 106.3 | abcd |
| 23 |  |  | 38 | 95.5 | bcdef |
| 24 |  |  | 40 | 96.4 | bcdef |
| 25 |  |  | 42 | 106.6 | abcd |
| 26 |  | 27 | 64 | 91.9 | cdef |
| 27 |  |  | 66 | 92.1 | cdef |
| 28 |  |  | 67 | 105.0 | abcde |
| 29 |  |  | 69 | 105.5 | abcde |
| 30 |  |  | 71 | 104.3 | abcde |
| 31 | 4 | 91 | 233 | 91.7 | cdef |
| 32 |  |  | 234 | 97.4 | bcdef |
| 33 |  |  | 235 | 94.2 | cdef |
| 34 |  |  | 236 | 94.3 | cdef |
| 35 |  |  | 237 | 92.8 | cdef |
| 36 |  | 154 | 460 | 97.6 | bcdef |
| 37 |  |  | 461 | 112.9 | $a b$ |
| 38 |  |  | 462 | 98.7 | bcdef |
| 39 |  |  | 463 | 102.5 | bcde |
| 40 |  |  | 465 | 102.3 | bcde |
| 41 | 5 | 13 | 25 | 97.4 | bcdef |
| 42 |  |  | 26 | 95.9 | bcdef |
| 43 |  |  | 27 | 93.5 | cdef |
| 44 |  |  | 31 | 92.7 | cdef |
| 45 |  |  | 32 | 95.7 | bcdef |
| 46 |  | 142 | 414 | 90.3 | def |
| 47 |  |  | 415 | 88.0 | ef. |
| 48 |  |  | 416 | 101.9 | bcde |
| 49 |  |  | 418 | 100.2 | bcdef |
| 50 |  |  | 420 | 102.7 | bode |

Means with the same letter are not significantly different at $5 \%$

Duncan's multiple range test for relative growth rate on $5 \%$ upper asymptote


Means with the same letter are not significantly different at $5 \%$

Duncan's multiple range test for relative growth rate at $50 \%$ upper asymptote


Means with the same letter are not significantly different at $5 \%$

Duncan's multiple range test for relative growth rate at 95\% upper asymptote


Means with the same letter are not significantly different at $5 \%$


[^0]:    * Significant at 5\% probability level
    ** Significant at 1\% probability level

[^1]:    * Significant at 5\% probability level
    ** Significant at 1\% probability level

[^2]:    * Significant at 5\% probability level
    ** Significant at $1 \%$ probability level

[^3]:    Means with the same letter are not significantly different at 5\%

[^4]:    Means with the same letter are not significantly different at $5 \%$

[^5]:    Means with the same letter are not significantly different at $5 \%$

[^6]:    Means with the same letter are not significantly different at $5 \frac{3}{\circ}$

[^7]:    Means with the same letter are not significantly different at $5 \%$

