POST-ANTHESIS LEAF HEALTH AS POTENTIAL DETERMINANT OF YIELD IN EARLY EVALUATION OF DIVERSE WHEAT GENOTYPES

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CERTIFICATE

This is to certify that the thesis/dissertation entitled "POST-ANTHESIS LEAF HEALTH

AS POTENTIAL DETERMINANT OF YIELD IN EARLY EVALUATION OF DI-

VERSE WHEAT GENOTYPES" submitted in partial fulfillment of the requirements

for the degree of MASTER OF SCIENCE IN AGRICULTURE (PLANT BREEDING

AND GENETICS) of the Agriculture and Forestry University, Nepal is a record of orig-

inal research carried out by Mr. Deependra Dhakal Id. No. PLB-03M-2015, under my

supervision, and no part of the thesis/dissertation has been submitted for any other degree or

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Abstract

Recent advances in phenotypic and statistical techniques have enabled exploration of a large number of trait in a greater depth. Current study undertaken in Agriculture and Forestry University, Rampur, Chitwan highlights an experimental-design technique and a mixed modeling approach to uncovering association of wheat crops' yield with the traits characterizing it's leaf health (resistance to foliar disease, flag leaf greenness, leaf area under greenness (LAUG) and flag leaf relative chlorophyll content (SPAD)).

Randomization based experiment design constrained for replicated checks balanced within blocks was generated and employed for estimation of augmented treatment genotypes' effects – also known as augmented row-column design for a small number of checks. The design allows for the recovery of intra-block information to adjust for environmental effects, which was used for studying plant traits that have potential yield determining effect. Observations on crop specific responses were made through metric devices such as ruler, digital scale whereas those traits that could not be physically assessed, such as leaf chlorophyll content (SPAD), and canopy temperature depression (CTD), were measured with equipments (SPAD meter and infrared thermometer).

Even though a significant proportion of variation was heritable (Both entry and check genotypes' effects were significant), substantial variation in a yield component and several crop architecture traits (thousand kernel weight, plant height, flag leaf area, canopy temperature depression, days to heading and days to anthesis) was due to spatial components. Major leaf health traits that influenced yield were the leaf greenness and relative chlorophylly content traits. BLUP estimation and adjustment for fixed effects dictate that 68.9% of the genotypic variation of entry genotypes in yield was explained by the full model, that having both spatial and leaf health components. This has revealed that augmenting leaf health traits

with structured blocking factors leads to enhanced yield estimation. Field associated traits like weed associated crop damage and lodging loss occurring late in the season, on the other hand, did not have significant impacts in the yield determination.

Apart from the check variety Aditya, which gave lower yields, all other check varieties were similar. However, they had more contrasting differences in the yield component trait Number of tillers m^{-2} . Recommendation for inclusion of some of the entry genotypes in breeding programs are made based their random effects adjusted estimates. TRCH/SRTU-//KACHU/3/-KINGBIRD #1, WHEAR/-SOKOLL/4/-PASTOR//MILAN/-KAUZ/3/B... and MUNAL #1*2/4/-HUW234+LR34/PRINIA-//PBW3... are amongst the top three high yielding entry genotypes. Findings suggest for inclusion of these genotypes, having leaf traits characterizing good health and with high yielding characters, in advanced evaluation trials.

Dedication

I dedicate this dissertation to my parents, who provided me the opportunity to further my pursuit of knowledge, and to my lovely sister, who has always cheered me up, even at desperate times.

Chapter 1

Introduction

1.1 Background

The bread Wheat (*Triticum aestivum* L.), as a crop that ranks first globally in cereal protein value, and that provides for majority of staple carbohydrates in the developed world, is gaining attention in improvement of both quality and quantity. This is expected to meet the need, in a substantial portion, of feeding the burgeoning population and for achieving a state of food self-sufficiency in Nepal.

Agricultural statistics published by the Ministry of Agriculture Development states that the country imported food items worth around 10 billion NPR in the year 2015/016 (GoN, 2014). Despite appreciable amounts of committments from the government for agriculture, as evident from the recent trends in fiscal year allocation of budget for agriculture sector, a forerunner indicative of vision of ADS to achieve food self-sufficiency status is yet to be seen (GoN, 2017).

Wheat is one among a few major food crops grown worldwide. It is often times credited for having had a central role to the beginning of agriculture (Harlan, 1981). The Bread Wheat has one of the largest known genome size among cultivated crops. It is a hexaploid with a genome thought to be composed of two or more distinct species' genomes following a series of allopolyploidization events in the past. Bread wheat has been traced to be composed of *T. urartu*(A), the *Sitopsis* section of *Aegilops*(B), and *Aegilops tauschii*(D) genomes (Dvorak,

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Luo, Yang, & Zhang, 1998).

The population structure of wheat can be classified into three distinct gene pools characterized by the ease at which hybridization events occur among them. The primary gene pool consists of common cultivated and landraces of hexaploid wheat, cultivated and landraces of tetraploid wheat, wild *T. diccoides* and diploid donors of the A and D genomes to durum and bread wheats. With exception of from few rare cross combinations requiring embryo rescue, most individual of this gene pool readily hybridize. The secondary gene pool is composed of the polyploid *Triticum* and *Aegilops* spp.. The hybrids produced within this gene pool show poor chromosome pairing. The tertiary gene pool is comprised of individuals whose genomes are non-homologous. Diploid and polyploid species occur in this pool.

Based on kernel color, endosperm hardness, and other grain quality characteristics spring wheat is classified into three distinct classes: Hard red spring wheat, Soft white wheat, and Hard white spring wheat. The spring wheat commonly grown in terai region of Nepal is of Hard red type.

Improving crop ideotype, an important avenue of plant breeding, has motivated extensive research into phenotype and genetics of wheat so that ever long trend of increasing yield can be maintained. Studies have reported that highest yielding so called "Super wheat" have robust semi-dwarf stems, broad leaves, large spikes with a greater number of grains per head, and higher grain weights (Swaminathan, 2007).

Crop breeding programs frequently target yield component traits rather than the yield *per se*, with the hope to have positive effect on yield (G. Slafer, 2003). Several candidate traits set themselves upto inventory of potentially desirable for yield improvement, but only a few of them can actually be reliably used in planned improvement programmes. The problem with candidate traits usually is that underlying mode of trait influencing the yield is not well established, and even if the biological pathway is well established, problem may lie if the variation associated with the trait is not stably inhertited (Saadalla, 1994); even acclaimed traits such as number of spikes per plant are asserted as not being an effective selection criterion (McNeal, Qualset, Baldridge, & Stewart, 1978). Thus, the experiments fail to produce expected results under differing field conditions (Okuyama, Federizzi, & Barbosa

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Neto, 2005). Characterization of effects of leaf health in yield and component traits, however, require measurement of stably expressed traits developmental and morpho-physiological traits, which are but only largely affected by a few attributable environmental factors, such as temperature or soil moisture these traits (Joshi et al., 2007).

Post-anthesis foliar health could be a proxy for determining final grain yield. It has a direct implication on grain filling behavior of a plant. Ideally, the contribution of preanthesis photosynthetic reserves to the final grain yield has been found to stall around 5% to 10% (R. Sharma, 1992). A major fraction of total yield, therefore, is dependent upon the photosynthesis that occurs post-anthesis. On the other hand, The yield synthesis post-anthesis is mainly a derivative of leaf area duration, which in turn is a function of leaf area index at anthesis and leaf longevity (J. Bingham, 1969). Terminal leaves, particularly the flag leaf and the one directly below it, have significant contribution to genotypic variation in yield (Aslam & Hunt, 1978) with respect to provisioning of assimilate to functionally active parts of plants including the storage organ(grain). This variation has been ascribed largely to the difference in the leaf longevity (Lupton, 1972).

Similarly, it is well known that various biotic agents, primarily diseases, sustain reduction in yield with pathways that relate to the effective photosynthetic surface area (Lopes & Reynolds, 2012). In a study that tested diverse lines of wheat from Nepal, Bangladesh and Philippines for tolerance to Helminthosporium leaf blight(HLB) disease, various wheat lines differed significantly for days to heading and maturity, plant height, tiller number, grain yield, area under disease progress curve (AUDPC), and the highest disease score (HDS). There was also a weak negative correlation between yield and AUDPC (r = -0.20) and HDS (r = -0.40) (N. K. C. Sharma R. C. & Shrestha, 1994).

An effective photosynthetic surface area, especially that of uppermost leaves, is probably one of the prime component contributing assimilate to the grain filling. Therefore, a range of conditions that have a measurable effect on leaf health may have implications in yield determination. Specifically, the stay green trait(indicative of leaf longevity and expressed as decline in photosynthetically active chlorophyllous surface), the foliar-disease resistance trait(expressed as levels of necrotic or chlorotic appearance), the leaf area index (LAI),

the leaf waxyness and glaucousness, and other traits associated with the canopy structure that potentially predispose leaf to environmental factors, all of which prolong post-anthesis photosynthetic activity might be traits determining crop's field level yield.

1.2 Statement of problem

Due to intrinsic nature of agricultural land and unattended sources of heterogenity, breeding materials are often tested under limited precision. Improved randomization based experimental designs with multidimensional blocking structure that accommodate large number of treatment entries in a study, could provide an efficient alternative to traditional block designs for the estimation of effects.

While contextualizing on early testing problem, new experimental design framework provides an opportunity for reliably discriminating the variation among genotypes of different genetic backgrounds. The recovered intra-block information, will be used to study traits that contribute to post-anthesis leaf health and finally to the grain yield. The study aims to make model comparison based inference on the suitability of selecting certain crop traits over others and uses multivariate techniques in making recommendation for certain genotypes. Finally, the best among the set of candidate mixed models will be presented for yield prediction with model characteristics.

Accurate field level prediction of yields is essential for making informed decisions about genotypic performance and in order to increase effectiveness of phenotype based selection, especially during early stages of genotype testing. This invariably means a better preparedness and a potential addition of useful germplasm to the national inventory for fighting challange of hunger ellimination and for tackling global climate shift.

1.3 Objectives

• To identify post-antheis leaf health traits associated with yield

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• To analyse association between the post-anthesis leaf health and yield and yield component, crop morphological and phenological traits

- To formulate yield prediction model based on post-anthesis leaf health
- To identify superior genotypes with favorable variation for leaf health

Chapter 2

Literature review

2.1 Use of unreplicated designs in field trials

Early testing is a problem of how to optimize the use of resources. For a plant breeding program, the subject quickly bumps into the problem of having to reliably discriminate between the germplasm resources. The outcome of any planned crossing programme and it's advanced generations—the seed propagule, has to qualify a test or more in order to proclaim itself valuable, and to noticeably proceed to further generations, eventually to make its way being released as a cultivar.

A replicated trial generally forms the basis of estimation of error variance that arises due to heterogeneous experimental conditions, also the inference about the treatment effects should acknowledge largely the role of error variance that characterizes most biological traits (Machado & Petrie, 2006), crop's agro-morphological traits being no exception. While it is customary to implement blocking in replicated trials of the genotype under evaluation with the inclusion of certain check cultivars for local control, the option comes to no avail when the replication of the genotype under test is not feasible due to scarcity of the material. The constraint in the amount of seed of a particular genotype and abundance of number of such genotypes is often what constitutes most early generation testing program. As described by statisticians and field researchers elsewhere, a possible solution here is to augment a randomized design trial having a certain fraction of the plots allocated to check cultivar with entries at limited quantity in some designed setup with partial or even no replication at all

2.1. Use of unreplicated designs in field trials

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(Walter T. Federer & others, 1994; H.-P. Piepho & Williams, 2016). It is in this constrained

scenario that the applicability of Incomplete Block designs (Yates, 1936) is realized.

Some of the well known augmented designs and variants of it include randomized complete

block and resolvable incomplete block designs for addressing one way heterogeniety, early

row-colum designs consisting of equal number of rows and columns defined with systematic

arrangement of treatments including that for checks (W. T. Federer, Nair, & Raghavarao,

1975; Lin & Poushinsky, 1983). Each of these designs can be used to estimate the error

variance in unreplicated trials with variying efficiencies.

In general, two common variants of augmented designs exist: Augmented block design

and Augmented row-column design. As the name suggests, the former class of designs are

useful in elliminating heterogeniety in one direction (in which blocks are laid), while those

of latter class are effective in elliminating in two or more way field heterogenity.

2.1.1 **Augmented block designs**

Implementing augmented block design in study, a researcher is concerned with estimating

block effects and the new varieties' (here-on also referred to as entry or new entry) effects.

This class of designs was first described by (W. Federer, 1956). Since the checks are the only

units repeated in the field, the portions of field that constitute a block from the combination

of checks provide information required to estimate the block effects. Only those designs

are selected where, the block effects are variance-balanced (i.e., variances of difference

between blocks effects are all equal). This is a condition to be met by design to be regarded

as optimal.

Design principles

A simple augmented block design addresses unidirectional heterogeniety in the field. In

augmented designs where checks are laid in randomized complete blocks, following design

parameters need to be specified:

Number of checks: ν ($i = 1, 2, 3, ..., \nu$)

Number of blocks: r (j = 1, 2, 3, ..., r)

Number of new varieties(entries): ν^* ($i = 1, 2, ..., \nu^*$)

Check varieties are randomly alloted to the experimental units within each of the r blocks which are complete for the check varieties, and let the yield of the i^{th} check variety in the j^{th} block be y_{ij} . Let the ν^* new varieties be randomly arranged such that l_i new varieties occur in the j^{th} block, let $\sum_{i=1}^r = \nu^*$, and let z_{ij} be the yield of the i^{th} new variety in the j^{th} block in which it occurs. For simplicity, we are assuming that the block effects are non-random effects. Note that for the z_{ij} yields, j=1,2,...,r and $i=1,2,...,\nu^*$. Using the check variety yields alone, run the statistical analysis for a randomized complete block design, and let EMS be the estimated error mean sum square obtained from this analysis. Then for any pair of those new varieties, say i and i', occuring together in the same block, the difference in their effects is estimated by $z_{ij}-z_{i'j}$ with standard error of $\sqrt{2EMS}$; the difference between two new varieties, i and i', occuring in blocks j and j', respectively, is estimated by $z_{ij}-z_{i'j}-(\bar{y}_i-\bar{y}_{i'})$ with a standard error of $\sqrt{2(\nu+1)EMS/\nu}$, where \bar{y}_i is the mean of the j^{th} block computed on check variety yields only.

Statistical analysis

The statistical analysis may be done in following way:

Step 1

The trial on $v + v^*$ varieties may be analyzed using standard methods for disproportionate numbers in the subclasses; then, contrasts among the check varieties, among the new varieties, and among the checks and new varieties may be made.

Step 2

A statistical analysis is performed on the check variety yields only, and blocking effects, a general mean effect, and check variety effects are estimated; an estimate of the experimental error variance is obtained. Then, the estimated new variety means or effects are obtained and the varietal contrasts are made as in Step 1.

2.1. Use of unreplicated designs in field trials

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2.1.2 Augmented row-column design for small number of check varieties

Improvements have been made in early block form of augmented designs in the past.

With the description of row-column designs in early half of 1970, modification of original

systematic-treatment-placement designs such as augmented lattic square designs (Walter T.

Federer & Nguyen, 2002), and α - α designs (E. Williams & John, 2003) were introduced.

Augmented row-column designs for small number of checks were, however, a much latter

addition.

This class of augmented designs are more flexible, in that they allow for arbitrarily large

number of new cultivars, the number of rows and columns, and even allow for dealing with

additional form of three way field heterogenity.

Design principles

Following design parameters are required to initialize a search for an efficient row-column

design:

Number of rows: k

Number of columns: s

Number of checks: ν_c

Number of rowgroups: g_k ($rg = 1, 2, ..., g_k$)

Number of colgroups: g_s ($cg = 1, 2, ..., g_s$)

Number of new entries: ν_e

Number of plots allocated to checks per block: $\nu_{q_kq_s}$

H.-P. Piepho & Williams (2016) has provided a framework for generation of this form of

design. The steps are mentioned as follows:

1. Blocks are the units formed by the intersection of adjacent rowgroups and colgroups

 $(g_k g_s)$. Define the number of plots for each block $(g_k.g_s)$ that are to be allocated to

checks. Note that, this number need not be a constant.

- 2. Allocate check cultivars (ν_c) to check plots based on the following model for design factors: $g_k + g_s + g_k \cdot g_s$.
- 3. In each row group (i.e., g_k), consider the classification of rows (k) by column groups (i.e., g_s). From step 2, we have a design for ν_c -by- g_s classification. Consider both check cultivar and g_s of this design as factors and optimize the allocation of rows so that effects for rows can be estimated with good efficiency. This produces a row-column group design for check cultivar.
- 4. In each column group (i.e., g_s), consider the classification of row groups (i.e., g_k) by columns (s). From step 2, we have a design for the check ν_c -by- g_k classification. Consider both check cultivar and rowgroups of this design as block factors and optimize the allocation of columns so that effects for columns can be estimated with good efficiency. This produces a row group-column design for check cultivars.
- 5. Merge the two designs obtained in steps 3 and 4 to obtain the final design for checks.
- 6. Allocate entries to free plots in completely randomized order.

Statistical analysis

The responses sampled from a row-column design may be analysed using linear models with effect for both the row and column factors and, effects for genotypes. Futher effects(rowgroup, columngroup, and rowgroup-columngroup interaction) that are associated with with block also need to be specified, due to restricted randomization imposed in allocation of checks within blocks. Thus, the full model for a response y can be stated as show in Equation (2.1).

$$y = \mu + rowgroup + colgroup + rowgroup.colgroup + rowgroup.row + colgroup.col + genotype + e$$
(2.1)

where, μ is the common intercept and e is the residual plot error.

All block effects, with exception of effects for genotype, may be taken as random. Fur-

thermore, the linear model may also be adjusted to fit a separate terms for fixed effects for checks and a random effects for new entries.

2.2 Crop development and yield formation

Process attributing to yield formation, mainly growth and phenological processes, transcend the level of inputs and agricultural sophistication as evidenced by remarkable similarity of trend in yield despite differences in average yields across different countries for the past century (Satorre & Slafer, 1999).

Crop development is defined as the sequence of phenological events conditioned by external factors resulting in changes in the morphology and/or function of some organs (Landsberg, 1977). Internal or external morphological changes can, thus, describe the crop development. Duration of each phase of development and the developmental pathway of each subsequent phase, which is dictated by the intitiation of primodial structures during earlier phase, is determined by the interaction between genetic and environmental factors. Among several methods those used to describe major developmental events, some which are non destructive in sampling are also in use, one of such being the growth scoring code postulted by J. C. Zadoks (Zadoks, Chang, & Konzak, 1974).

As complex a trait the crop yield is, it is explained well when explored in relation to developmental phase of crop. Dynamics of multitude of factors are involved since the very beginning of a life cycle of crop, as early as the beginning of embryonic activities when inside the seed.

2.2.1 Leaf development

By the time wheat crop has grown to a seedling, five to seven leaf primordia will have been developed (McMaster, 1997), meaning that two to three leaf primordia will have already developed well before the tip of the first leaf appears. Duration through which the apical primary primordia persists depends on genotype and environmental factors.

Plastochron(thermal time between the initiation of two consecutive leaf primordia) is usually a constant value for a given environment (Wilhelm & McMaster, 1995). The maximum number of leaves is determined when the apex changes the development from vegetative to reproductive with the floral initiation. A model devised by Kirby, Appleyard, & Fellowes (1985) relates leaf and spike initiation phases with number of leaves that appear in main shoot.

2.2.2 Tiller development

The phytomer contains axillary tiller buds, each of which is able to differentiate into leafy tillers. The relationship between the number of phyllochrons and the number of primary tillers that appear approaches perfectly linear, with an intercept at 3 units in horizontal axis when plotted the number of primary tillers approximated by phyllochrons. Similar pattern follows in tillers that develop from secondary and tertiary tillers. This relationship is only valid only to the extent provided resources are unlimited, in field, however, the potential tiller emergence rate is slower than that predicted by Fionacci series (Masle, 1985), as competition occurs within a single plant system and also between multiple of those.

Alongside the emergence and growth of the tiller, in environmental conditions, tiller mortality is also a defining factor for overall crop stand due maturity. This occurs in the order reverse to that in which the tillers appeared. Phenomena of tiller mortality is reasoned to accelerate at around stem elongation phase due concomitant increase in demand of assimilates to the developing shoots.

2.2.3 Spikelet and floret development

Spikelet intitiation begins with the appearance of double ridge in the apical region of a shoot. This is followed by the floral initiation and finally terminates with the initiation of terminal spikelet in the apical meristem. It is also shown that the rate of spikelet initiation is negatively correlated total number of leaves on a shoot.

The floret development progresses towards the ends starting with the signs first visible in

the lower middle portion of a spike head. Similarly, accumulation of the Growing Degree Days(GDD) provides an indication of for how long floret initiation within each spikelet continues. It continues approximately untill a GDD period of 200-300°C above 0°C, with the latter being reference temperature, before the appearance of the flag leaf ligule. It is postulated that floret death too occurs due to increased competition for assimilates, once stem and spikes growth surges, alike tiller death. The competition phenomena that determines the proportion of florets liable to maintain normal rate of development has been validated in studies that compared the genotypes having different partitioning between spikes and stems ratios (Brooking & Kirby, 1981).

2.2.4 Grain filling

Wheat is a kleistogamus plant. So, once the pollination occurs inside a floret, it begins to develop as a potential grain after it has been are fertilised.

Grain development in wheat passes through a series of distinct well characterized phases. In the initial "lag phase", the grain does not gain much in weight. Although the phase generally spans almost $1/4^{th}$ of the total postanthesis period, only 5 to 10 percent of total grain weight is gained during this. However the term lag phase ascribed to earlier phase, the latter stage of grain growth— which substantially adds to the grain weight, can be well modelled both by in a logistic function as well as bilineal function.

This stage of development too is strictly regulated by the thermal time, in absence of severe environmental stresses however, in which case permature termination of grain filling leads to sub-optimal dry weight (Nicolas, Gleadow, & Dalling, 1984).

2.3 Crop yield and yield component traits

In modern wheat, gains in yield increases are due to increased harvest index, shorter period from sowing to anthesis, and more number of grains per unit area. Therefore, spikelet initiation and formation stage, which occurs before anthesis, and has direct relation to yield.

This asserts that total number of grains/kernels per spike head(a component trait of yield) is largely a function of pre-flowering vegetative stage. It has also been concluded that genetic yield increases resulted from an increase in the number of kernels produced rather than an increase in kernel size (McCaig & DePauw, 1995). This suggests that bread wheat has been sink-limited at grain filling.

From the crop physiology perspective, yield component trait (i.e., for wheat, plants per unit area, spikes per plant, grains per spike, and grain weight) approach to understand yield is one but largely unreliable approach (Fischer, 1996; G. Slafer, Calderini, & Miralles, 1996). Owing to the fact that individual yield components are negatively correlated with each other, improvement in one trait will inevitably result in compensative negative effects in some other (G. Slafer, 2003). But understanding of exact nature of relationships between component traits might, enable breeders to manipulate them correctly (G. Slafer et al., 1996). However, given that there exists a strong negative correlation among mostly vegetative stage defined yield component traits (summarized here as the number of grain kernels), it might be reasonable to seek to manipulate a different component trait such as grain yield. The grain weight is probably a much a simpler component, considering its distinctness from other correlated yield component traits in the pathway to yield determination; all other component traits could be summarized to number of grains for given unit of measure (usually, per unit land area).

2.4 Factors affecting wheat growth and development

The major influencer of wheat growth and development is temperature. Every stage of development is, however, to a different degree depending upon various factors (G. Slafer & Rawson, 1995), sensitive to temperature. This is unlike most other environmental growth affecting factors including moisture and photoperiod, which only exhibit their effects either when factors prevail at their extremes or are range in their effects based on the crop's growth stage.

2.4.1 Temperature

Generally, with higher temperature, the rate of development is faster and thus certain phase of crop growth becomes shorter. From among various temperature dependent growth models proposed to predict the growth phase, the thermal time model is the one most widely accepted (Squire, Marshall, Terry, & Monteith, 1984). The simplified form of the model is shown in Equation (2.2).

$$thermal\ time = \frac{mean\ daily\ temperature - base\ temperature(^{o}C)}{growth\ rate(kg\ day^{-1})} \tag{2.2}$$

Conceptually, thermal time is the inverse of change in rate of growth rate per unit temperature, from base to optimum temperature. The concetp assumes that the relationship between rate of development and temperature in linear. Similarly related are the leaf initiation and appearance and, the temperature. Emperically, The slope of the relationship is the rate of these processes per degree day, and the reciprocal of these slopes are the plastochron and phyllochron (Slafer & Rawson, 1995). So, at the least, the parameters of thermal time are specific for a given genotype and developmental stage. This way, the model can be linearized, as it remains unaffected by nuances of relationship change caused solely due to phasic development. In practice, however, thermal time is calculated as the accumulated daily mean temperature above a base temperature (Angus, Mackenzie, Morton, & Schafer, 1981; Rawson & Bagga, 1979).

Studies have highlighted the role of mean air temperature during booting and ear emergence (Calderini, Abeledo, Savin, & Slafer, 1999; Calderini, Savin, Abeledo, Reynolds, & Slafer, 2001). In a different study aimed at understanding the relationship between temperature dependent growth processes and natural variation associated with the temperature, mostly those conspicious during vegetative to reproductive transition of wheat crop, authors investigated the vernalization requirements among Nepalese wheat and barley. Wheats from villages of high altitudes tended to require more vernalization than those from lower altitudes suggestive of local adaptation. This also leads to inference that seed exchange among Nepalese farmers is either poor or well provisoned with a sound knowledge base about local varietal adaptation (Rao & Witcombe, 1977).

Unlike atomospheric temperature, which is mostly influenced by incident solar radiation, Soil or root zone temperature is influenced several factors. The factors include intensity, the quality, and the duration of solar radiation, air temperature, surface vegetation, and the color and the thermal conductivity of the soil. The optimum soil temperature reported for maximum yield of the Bread wheat is $20^{\circ}C$ (Whitfield & Smika, 1971).

The cardinal air temperature for growth of wheat is summarized below:

Table 2.1: Table of cardinal of temperature of wheat growth (Briggle & Curtis, 1987)

	Temperature
Minimum temperature	$3^{o}C$ - $4^{o}C$
Optimun temperature	$25^{o}C$
Maximum temperature	$30^{o}C$ - $32^{o}C$

2.4.2 Soil moisture

Water has it's role in translocation of mineral and anabolic reactions inside plant system. Both high and low levels of moisture can reduce the crop yield, as excess moisture, above all, reduces soil aeration and thus the supply of Oxygen to roots. This can, in turn, have negative effect on soil microbial dynamics. Moisture stress causes both the reduction in cell division and cell elongation, thus in overall growth.

Wheat can be grown in regions where annual precipitation ranges from 250 to 1750 mm. Water associated stress has detrimental effect in all yield associated components (Abayomi & Wright, 1999). In a study that correlates effects of different levels of soil water tension measured near surface, authors point out that component traits like number of spikes per square meter, the number of spikelets per spike, and the number of grains per spike are among the most affected by increasing levels of soil-water-tension (H. Zhang & Oweis, 1999). Root characterstic is mostly associated with the genetic variation that confers drought resistance (Hoad, Russell, Lucas, & Bingham, 2001).

Wheat grown in medium textured soil is unlikely to have simple moisture deficit(occurs

when root zone profile exausts of available water) during early stage of crop growth since wheat is usually planted under favourable soil moisture conditions. However, more serious consequences are faced in grain yield when water shortage occurs, in general, after ear initiation. One of the conditions which emphasizes role of moisture stress is due tiller mortality, which is dependent on pre-anthesis water availability (Aspinall & Husain, 1970; Begg & Turner, 1976). The potential grain number which is directly related to the production of fertile gametes and fertilization is other most influential yield-determining process affected by moisture stress during critical periods of pre-anthesis and anthesis (J. Bingham, 1966). Similarly, post-anthesis drought causing a reduction in thousand grain weight has been described by some authors as other major determinant of yield (Asana, Saini, & Ray, 1958; A. Day & Intalap, 1970; W. Day et al., 1978).

2.4.3 Solar radiation

The intensity of solar radiation varies with the latitude and the season. Tropical to Sub tropical regions occurring within lattitude $20^{\circ}N$ - $30^{\circ}N$ observe the highest incidence of solar radiation.

Solar radiation is converted through fixative cycles into biomass or the crop's usable produce. An emperical framework to describe the relationship between the absorbed Photosynthetically Active Radiation (PAR_A), the radiation use efficiency and the factors accounting for the canopy structure, which is represented by leaf area index (LAI) and the radiation extinction coefficient (k), is stated by Beer's law (Goyne, Milroy, Lilley, & Hare, 1993). The relationship is presented in Equation (2.3).

$$PAR_A = PAR_0[1 - e^{-kLAI}] (2.3)$$

where PAR_0 is the above canopy or incident radiation.

Radiation use efficiency is the slope of a linear function having dry matter accumulation per unit time, at the rate unconstrained by stress causing extraneous factors like soil moisture, nutrients, temperature, and the number of actively growing points, as response approximated by the absorbed photosynthetically active radiation (PAR_A) .

Wheat is not the most efficient utiliser of PAR_A , with only 3.7% of total incident radiation utilized for dry matter accumulation. Sparing the energy efficiency factor, wheat has, however, a better growth rate compared to the C_4 plant Corn, reported when compared across similar growing conditions of temperate region (Fageria, Baligar, & Jones, 2010). In wheat, most of the between cultivar differences in growth duration is accounted for by photoperiod and vernalization sensitivity factor associated with the genotypes (Satorre & Slafer, 1999). Besides that, there is general agreement that a variety should have a high leaf area index and erect leaves, so as to make the best use of incident solar energy.

Damania, Jackson, & Porceddu (1985) measured morphological traits and heading date in wheat and barley landrace samples from Nepal and Yemen grown at an experiment station. They found considerable variation within and between regions. In the case of barley from Nepal and wheat from Yemen this variation correlated significantly with the altitude of origin.

The correlations of Grain Filling Period (GFP) with other traits indicated that improvement in GFP should also improve grain yield, harvest index and kernel weight (R. Sharma, 1992). Findings from the study have asserted that selecting for simultaneously longer GFP and a shorter Vegetative Growth Period (VGP) which is related to earliness of heading, will lead to improved yields. In addition, the author mentions that each of the periods expressed as proportions is negatively correlated with the other which adds to the importance of precisely identifying the growth periods during selection. N. K. C. Sharma R.C. & Mishra (1993) also reported that biomass yield, harvest index, kernel weight, and durations for the vegetative growth (VGP) and grain filling (GFP) were related to the grain yield.

R. Sharma (1987) reported that the seed rate affected the grain yield, tiller number, grains/spike, 1000 kernel weight, days to heading, and plant height at harvest. For both reduced and higher seed rates than the recommended (120 t ha^{-1}), there was an adverse effect on all the above-mentioned traits. The effect of dew on wheat yield and soil water balance at Rampur showed that grain yield was significantly higher for irrigated treatment (4.02 t ha^{-1}) followed by rainfed (3.03 t ha^{-1}), dew (2.60 t ha^{-1}) and control (2.58 t ha^{-1}) treatments. The moisture meter readings and soil moisture content analysis also showed

that dew did not significantly contribute to the evapo-transpiration requirement of wheat (P. Sharma, Khakural, Chhetri, & Chhetri, 1984).

In a study using different intensities of leaf defoliation as treatment in wheat to study relationship among genotypes with different levels of foliar pathogen resistance, investigators reported the removal of flag leaf as having significant effects on both the yield and yield component traits (Rosyara, Sharma, Shrestha, & Duveiller, 2005). Generalizing over the fact that the loss of most photosynthetically active organ causes reduction in yields, though at various levels of severity based on the genotypic reaction to disease, characterization of candidate traits that actively interact in the field grown wheat to influence yields is the subject matter of due concern.

Chapter 3

Materials and methods

3.1 Germplasm

The genotypes used in the current study consisted of 104 diverse lines of wheat that have been advanced as representative lines bearing variable resistance to diseases. Four check varieties (Described in Table 3.1), released at different periods, adapted to a range of environmental conditions in Nepal were included alongside in the study. The genotypes were tested for their phenotypic traits that have potential implications for prolonging the leaf health during post anthesis period. Table 3.5 enlists trait observations that were taken during the study in chronological order.

Table 3.1: A summary of check varieties used in the study

Characteristics	Tilottama ¹	Bhrikuti ²	Gautam ³	Aditya ⁴
Year	2014	1994	2002	2010
Breeding	Nepal Agriculture Research	Nepal Agriculture Research	Nepal Agriculture Research	Nepal Agriculture Research
program	Council	Council	Council	Council
Synonym	NL 1073; NL1073; NL-1073; FRAN- COLIN; FRANKOLIN		BL 1887; BL-1887, BL1887	
Growth Habit	Spring	Spring	Spring	Spring
Variety Type	Bread Wheat	Bread Wheat	Bread Wheat	Bread Wheat
Semidwarf (Rht gene) Grain color	Yes	Yes	Yes	White grain
Grain hardiness				· ·

Ihttp://wheatatlas.org/varieties/detail/26413

²http://wheatatlas.org/varieties/detail/22378

³http://wheatatlas.org/varieties/detail/22383

⁴http://wheatatlas.org/varieties/detail/22373

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Pedigree	WAXWING*2/VIVITSI	CONTESTAD- O/COCORAQUE F 75/3/POLLO//F- URY/ANAHUAC F 75	SIDDHARTHA/NG 8319//NL 297 or SIDHHARTHA/NG 8319//NL 297	GS348/NL746//NL748
Selection history		NL 623 FR 2308-2F-1F-0F	NC 1838-4B-020B-020B- 2B-0B	
CID (Cross ID) SID (Selection ID)		251774 0		
Origin CIMMYT	Advanced line from CIMMYT FRANCOLIN	Advanced line from CIMMYT	Cross made in the country, one CIMMYT parent	Cross made in the country, no CIMMYT parents
breeding line Genes	TRANCOLIN			
Selection Code		Line from CIMMYT, further local selection within country		
Ug99	Yes	No	No	No
Stem Rust Yellow Rust	Yes Yes			
Leaf Rust	Yes			
ME1 ME2 ME3		Spring Wheat Spring Wheat	Spring Wheat Spring Wheat	Spring Wheat
Production	Irrigated and semi-irrigated		Irrigated, both normal and	Terai, Tars and Lower
Conditions	areas of Terai and inner Terai		late sown condition of whole Terai, Taar and	Valleys up to 500 500 masl; maturity 118 days, yield
			foot-hills (<500 m); maturity 105-115; potential yield - 5t/ha	potential 4.8 t/ha
Suitable megaen- vironments	ME1	ME1, ME5	ME1, ME5	ME1
Notes	Resistant to spot blotch. High tillering. Early maturing, stay green, excellent for bread and chapati.			
Cultivated area	<u>F</u>	153,055 (2011/12)		38263 (2011/12)

The trial was conducted under the condition of natural exposure to environmental factors (biotic and abiotic). The field is located at agronomy Farm of Agriculture and Forestry University(AFU), Chitwan, Nepal. A brief geographical context of the experimental site, and a glimpse at history of prevailing weather over the growing season, based on previous year data, are shown in Table 3.2⁵ and Figure 3.1.

Table 3.2: Site of study factsheet

Particulars	Detail
Institution	Agriculture and Forestry University
Farm	Rampur

⁵http://wheatatlas.org/station/NPL/22506/

Particulars Detail Chitwan State 84.4^{o} Longitude 27.62^{o} Latitude Altitude 191 meters Environment ME₁ Dominant Soil(FAO classification) Cambisols (Dystric Cambisols) Surface pH Rice-wheat Cropping system

Table 3.2: Site of study factsheet

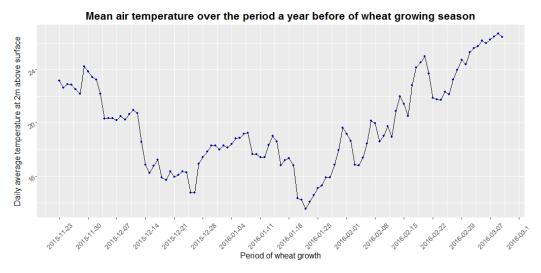


Figure 3.1: Mean air temperature 2m above surface at the site of study a year before expected wheat growing season

3.2 Planting and management

Field was sown in Nov 24, 2016. Seeds were continuously sown within a row while maintaining a between row distance of 25 cm. Seeding depth of 3-5 cm was maintained. Soil sampling (results not shown) show that the dominant soil texture type occuring in the study site is medium to heavy loamy soil. Weeding was carried out twice, first 40 days after sowing and second 70 days after sowing. Blanket recommendation of fertilizer application was followed; Initially, full dose of K_2O (60 $kg\ ha^{-1}$) and P_2O_5 (60 $kg\ ha^{-1}$) were applied at the time of sowing, with Nitrogen (fed as Urea) in split doses(60 kg N ha^{-1} as basal and remaining 60 kg N ha^{-1} top dressed after irrigation). Total lengths of the field row and column were 18 m and 19.5 m, respectively. Individual plots of

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size $0.94~m^2$, occupying a Total area of $351~m^2$ (Net Plot Area: $225~m^2$) were laid in rectangular grids of regular shapes.

Irrigation was limited to once-only throughout the growing period. It is expected that the field will have ample amount of residual moisture during crops' establishment period to start with, because the site usually stays logged with water during the months of August and extended halves of September. The irrigation will be scheduled to best avoid pre-antheis moisture stress, as this has been implied in largest losses resulting in number of fertile florets and the final grain weight (Innes & Blackwell, 1981). Water will be sourced from a deep tubewell and the field will be moistened untill saturated field condition will have been achieved.

Application of insecticidal sprays and disease checking sprays were avoided. This is to promote natural epiphytotic conditions. Observations on disease severity were mostly taken in reference to Zadok's scale of growth scoring.

3.3 Design, treatments specification and layout

Current study, using a small number of check varieties and sparing amounts of test entries, is adapted to fit a problem domain of early genration varietal testing applying row-column design for smaller number of checks. This allows for the estimation of design effects as spatial components, as well as for specification of custom spatial error model, if required.

- Number of rows (*k*): 12
- Number of columns (s): 20
- Number of checks (ν_c): 4
- Number of rowgroups (g_k) : 3
- Number of colgroups (g_s) : 4
- Number of new entries (ν_e): 104
- Number of plots allocated to checks per block ($\nu_{g_kg_s}$): variable

An augmented design that accomodates $\nu_e = 104$ unreplicated entries and $\nu_c = 4$ replicated check cultivars with the number of rows k = 12 and the number of columns s = 20 was been generated. Each row and column is a complete block with respect to checks. A block $(g_k g_s)$ specified by intersection of a rowgroup and a colgroup contains complete but varying number of check plots. A field layout of the the randomized design has been shown in Figure 3.2.

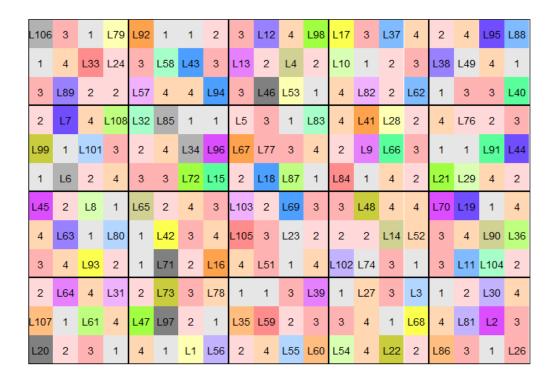


Figure 3.2: Randomized layout of Augmented Row-Column Design

3.4 Sampling and statistical analysis

Individual plots were sampled for making observations. At least four plant hills, except for visual scoring and making observations on plant-environment interacting response (eg. canopy temperature depression, soil temperature and soil moisture traits) where distinct protocol exists, were randomly selected and a few representative tiller from each hill were duely analysed for their relevent traits (enlisted in Table 3.5).

As described earlier in Section 2.1.2, a mixed linear model of the form as shown (with modification), in Equation (2.1) was fitted for each of the response variables y. The genotype term was further

partitioned to separate out Check cultivars and entries (H. Piepho, Williams, & Fleck, 2006). An assumption regarding entry genotypes that these arise from a common set of germplasm and that as a whole represent all variation in a breeding population of interest was made.

All model terms except for the Check variety effects, which was considered as fixed because they are purposely selected, were expressed with random intercept terms for obtaining model parameters (Wolfinger et al., 1997). Model fitting was done through lmer function of **lme4** package. The model formula was fitted for obtaining REML estimates of the parameters by a single call to for both fixed and random effects terms. A general matrix structure representation in a linear mixed model with conditional distribution of y given $\mathcal{B} = \beta$ can be represented as shown in Equation (3.1),

$$(y|\mathcal{B}) \approx \mathcal{N}(X\beta + Zb, \sigma^2 W^{-1})$$
 (3.1)

where Z is the $n \times q$ model matrix for the q-dimensional vector-valued random effects variable, \mathcal{B} , whose value we are fixing at b. The unconditional distribution of \mathcal{B} is also multivariate normal with mean zero and a parameterized $q \times q$ variance-covariance matrix (Bates, Mächler, Bolker, & Walker, 2015).

For multiple comparison of fixed effects means and stepwise variable selection procedure, least squares means and confidence intervals were calculated by Satterthwaite approximation of degrees of freedom (Kuznetsova, Brockhoff, & Christensen, 2017).

Likewise, Relative Efficiency (RE) of alternative model over null model was calculated to augment the decision about model selection. Formula (3.2) was used to compute RE.

$$RE = \frac{Error\ variance\ of\ Null\ Model}{Error\ variance\ of\ Alternative\ model} \tag{3.2}$$

All exploratory and analytical works were accomplished in "R" (an open source computing environment), with open source statistical packages. An extensive list of all packages used in the course of modeling and inference is provided in Appendix B.

3.5 Leaf health assessment

Leaf health trait was explained as a composite of two componenet traits- leaf greenness trait and foliar pathogen associated leaf disease score- each of which, on their own, are more readily observable phenotypic traits.

3.5.1 Flag leaf greenness assessment

A two approach method has been adopted to describe the SG feature. This feature was defined as:

- 1. Leaf Area Under Greenness(LAUG) approach
 - difference for 0-9 visual score of green coloration (chlorophyll) of flag leaf at anthesis (Zadoks stage 65), medium milk (Zadoks stage 75) and soft dough (Zadoks stage 85) stages.

2. SPAD approach

• difference in SPAD values(recorded at Zadoks stage 65 and 85) of flag leaf.

The leaf greenness scores was considered to group genotypes into classess shown in Table 3.3:

Table 3.3: Leaf greenness score intrepretation

Greenness score	Greenness rating
0-2.5	non-green
2.5-5.0	moderately non-green
5.0-7.5	moderately green
7.5-9.9	green

A similar method has been outlined by (Joshi, Chand, & Arun, 2002) for SG trait assessment in wheat. But, current method employs emperical basis outlined in (Joshi et al., 2007) for LAUG calculation. The Leaf Area Under Greenness (LAUG) based on greenness scores over time (PLANK & others, 1963) was estimated from the Equation (3.3)

$$LAUG = \sum_{i=1}^{a} \left[\left\{ \frac{Y_i + Y_{(i+1)}}{2} \right\} \times (t_{i+1} - t_i) \right]$$
 (3.3)

3.5.2 Foliar blight complex associated leaf disease scoring

Disease score observations of individual plots were made for foliar disease pathogens (*Cochliobolus sativus*, *Pyrenophora tritici-repentis*) with attributable symptoms (E. Duveiller & Sharma, 2012) in a way descibed by (Saari & Prescott, 1975). Presence of the pathogen complex was revealed

upon microscopic studies, during early stage diagnostics of the disease. A image of *Cochliobolus sativus* seen under the microscopic in a slide slide is shown in Figure 3.3 as confirmatory diagnosis of foliar blight pathogen as causative agent of Foliar blight disease.. A single digit scoring system was adopted for scoring 5 random plants on each plot the flag leaf and the other directly lower to it. The score categories explaining the response to disease is interpreted in Table 3.4.



Figure 3.3: An image of slide under microscope showing *Cochliobolus* spp.

For assessing foliar pathogen associated leaf score, scoring was limited only upto Zadok's stage S77(Late milk stage). This is to acknowledge the fact that, after this stage the natural cycle of senescence confounds the effects that follow due to pathogen activity (Neupane et al., 2007).

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Table 3.4: Foliar pathogen associated disease score interpretation

Disease score	Score rating
0-2.5	very less affected
2.5-5.0	less affected
5.0-7.5	much affected
7.5-9.9	very much affected

3.6 Observation

Data were recorded from the field for the following response variables.

3.6.1 Yield, morphology, phenology, soil and atmospheric conditions

The record of all observations made on field, those including crops' yield, morphology, phenology traits, and fields' soil and atmospheric conditions has been shown in Table 3.5.

Table 3.5: Field records of crop yield, morphology, phenology, and soil and atmospheric conditions

Stage recorded
Tillering (Zadoks stage 21-29)
Jointing and Booting (Zadoks stage 37 and 40)
when approximately 50% of the plants in a plot are at
Zadoks stage 55
Anthesis (Zadoks 60)
Anthesis (Zadoks stage 65)
Anthesis (Zadoks stage 65)
Anthesis and soft dough (Zadoks stage 65 and 85)
Anthesis and Late milk (Zadoks stage 65 and 77)
when approximately 50% of the plants in a plot are at
Zadoks stage 65
Medium milk (Zadok stage 75)
Soft dough (Zadoks stage 85)
Ripening (Zadoks stage 90)
Ripening (Zadoks stage 92)

Days to maturity Ripening (Zadoks stage 92)

Number of grains per panicle After harvested

Thousand kernel weight After harvested and dried Grain yield After harvested and dried

3.7 Missing values treatment

Missing values are largely a result of omitting improbable and extreme values due to biases in observation. In particular, two plots have been systematically left out from the observation record and further analysis because of less than optimal plant population stand. Nevertheless, mixed modeling procedure employed in inference offers fairly robust measures against such occurances (West, Welch, & Galecki, 2014). For other procedures like correlation analysis (Shown in Table 4.29) pairwise complete data were used in computing the correlation coefficients. Similarly, in multivariate procedures such as PCA and cluster analysis, only complete cases were used in obtaining distance matrix.

Chapter 4

Results

Observed response variables are summarised under three major aspects of a crop trait—Yield and yield component traits, Leaf health traits and Crop morphological, phenological and architectural traits.

4.1 Modeling yield and yield component traits

4.1.1 Model estimates of yield and yield component traits

Table 4.1: Model adjusted genotypic mean responses of yield and yield component traits

encoding	Genotype	Yield	Effective tillers	Thousand kernel weight	Grain per panicle	Panicle lengtl
1	Aditya	3.02	26.7	36.3	27.1	16.3
2	Bhrikuti	3.55	30.2	37.7	26.8	16.8
3	Gautam	3.33	27.1	38.5	28.3	17.5
4	Tilottama	3.42	33.7	36.1	27.4	16.6
L1	MUNAL #1	4.07	27.0	35.9	28.8	16.6
L10	MUNAL/3/HUW234+LR34/PRINIA//PFAU/WEAVER/4	/	36.0			16.6
L101	PFAU/MILAN/5/CHEN/AEGILOPS SQUARROSA (TAUS)/	3.28	30.0	43.6	27.7	16.4
L102	PREMIO/5/TUI//2*SUNCO/SA1166/3/TUI/4/FINSI/	2.88	35.0	39.1	21.1	13.7
L103	PREMIO/BAVIS		47.0			15.0
L104	CHIYAK//ND643/2*WAXWING/3/ND643/2*WAXWIN	G	30.0			
L105	BABAX/LR42//BABAX*2/4/SNI/TRAP#1/3/KAUZ*2/	4.15	38.0	41.5	26.9	17.1
L106	PFAU/MILAN//TROST/3/MUNAL #1/4/PFAU/MILAN//	4.46	27.0	44.2	29.4	16.6
L107	YUK/AE.SQUARROSA (217)//2*PANDORA	2.51	30.0	39.4	22.3	13.8
L108	SUP152*2//ND643/2*WAXWING	3.44	36.0	41.5	28.1	12.9
L11	SWSR22T.B./5/KAUZ//ALTAR 84/AOS/3/KAUZ/4/		28.0			
L12	WHEAR/VIVITSI//WHEAR/3/PANDORA	3.68	33.0	46.5	30.7	15.7
L13	WBLL1*2/KURUKU//TACUPETO F2001*2/BRAMBLING	4.17	28.0	49.6	29.5	17.9
L14	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN/OTUS//	2.85	39.0	33.8	26.9	15.9
L15	FRANCOLIN #1*2/ND643/2*WBLL1	2.84	35.0	37.7	24.6	14.7
L16	KIRITATI/ 2*WBLL1/3/TAM200/PASTOR//TOBA97/4/	3.83	31.0	44.1	30.6	18.6
L17	SUP152/KENYA SUNBIRD		33.0			15.9
L18	SUP152/KENYA SUNBIRD	3.43	37.0	37.7	26.5	13.3
L19	SUP152/KENYA SUNBIRD	3.74	35.0	32.0	30.9	14.2
L2	KACHU #1	3.27	39.0	41.8	29.3	15.7
L20	WBLL1*2/BRAMBLING/5/BABAX/LR42//BABAX*2/4	1 /	26.0			
L21	ND643/2*WBLL1/3/KIRITATI//2*PRL/2*PASTOR/4/		34.0			16.3
L22	BETTY/3/CHEN/AE.SQ//2*OPATA/4/BAVIS	3.20	29.0	41.5	24.7	18.0
L23	BAVIS/VORB/5/CROC_1/AE.SQUARROSA(205)//	2.58	23.0	45.8	23.4	15.0
L24	BABAX/LR42//BABAX/3/ER2000/4/NAVJ07		39.0			17.7
L26	NING-MAI-96035/FINSI//HEILO/3/NAVJ07		26.0			
L27	WBLL1*2/BRAMBLING//TAM200/TUI/3/		39.0			17.1

Table 4.1: Model adjusted genotypic mean responses of yield and yield component traits (*continued*)

encoding	Genotype	Yield	Effective tillers	Thousand kernel weight	Grain per panicle	Panicle lengt
L28	KISKADEE	4.10	47.0	39.8	22.6	14.9
	#1/5/KAUZ*2/MNV//KAUZ/3/MILAN/4/					
L29	MERCATO//PARUS/PASTOR		31.0			16.5
L3	KACHU/BECARD//WBLL1*2/BRAMBLING	3.10	40.0	29.4	23.5	19.0
L30	PREMIO//PARUS/PASTOR	4.15	32.0	42.8	30.3	20.1
L31	ND643/2*WBLL1//ND643/2*WAXWING	3.81	37.0	46.9	23.9	16.3
L32	ND643/2*WBLL1//ND643/2*WAXWING	3.22	41.0	41.8	24.6	14.1
L33	ND643/2*WBLL1//ND643/2*WAXWING		41.0			16.7
L34	SNB//CMH79A.955/3*CN079/3/ATTILA/4/CHEN/		32.0			
L35	KIRITATI/ 2*WBLL1/8/SHA7/ PRL/VEE#6/ 3/		32.0			13.5
	FASAN/					
L36	SERI.1B*2/3/KAUZ*2/BOW//KUZ/4/CIRCUS/5/	2.50	28.0	40.6	22.3	14.3
L37	WHEAR/KUKUNA/3/C80.1/3*BATAVIA//2*WBLL1/4/	3.62	43.0	29.8	31.6	18.5
L38	BAJ #1*2/BECARD		24.0			16.5
L39	KACHU*2/MUNAL #1	2.72	28.0	46.8	23.5	14.1
L4	PRL/2*PASTOR/3/PFAU/WEAVER*2//CHAPIO		38.0			
L40	KACHU*2/MUNAL #1	3.20	27.0	36.0	21.8	14.4
L41	KACHU/2*MUNAL #1		34.0			
L42	KACHU/2*MUNAL #1		33.0			
L43	KACHU*2/PANDORA	4.36	50.0	48.0	23.5	17.0
L44	SUP152/2*MUNAL #1	2.43	24.0	47.9	24.0	13.5
L45	FRNCLN*2/BECARD		35.0			17.1
L46	WBLL1*		39.0			17.1
	2/BRAMBLING//SAAR/2*WAXWING/4/					
L47	CHIBIA//PRLII/CM65531/3/MISR 2*2/4/	3.45	30.0	42.4	26.5	15.2
L48	CHIBIA//PRLII/CM65531/3/MISR 2*2/4/		27.0			14.3
L49	WAXWING/KRONSTAD F2004//2*FRNCLN	3.49	36.0	29.4	31.9	17.4
L5	DANPHE/CHONTE	2.42	33.0	34.6	21.9	15.5
L51	MUNAL		27.0			13.3
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/					
L52	MUNAL	3.07	30.0	42.4	23.4	12.3
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/					
L53	MUNAL #1/2*FRNCLN		28.0			
L54	MUNAL #1/2*FRNCLN	2.89	34.0	36.9	29.2	17.5
L55	MUNAL #1*2//WHEAR/SOKOLL	2.72	31.0	45.8	24.9	13.9
L56	MUNAL #1*2/KINGBIRD #1	4.39	32.0	46.5	28.0	14.5

Table 4.1: Model adjusted genotypic mean responses of yield and yield component traits (*continued*)

encoding	Genotype	Yield	Effective tillers	Thousand kernel weight	Grain per panicle	Panicle lengt
L57	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//		31.0			15.9
L58	#1/4/OTUS// PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	2.74	35.0	40.9	27.5	15.4
L59	KAUZ*2/BOW//KAUZ/3/W98.6.38/5/SABUF/4/		26.0			16.4
L6	SUP152/FRNCLN		25.0			
L60	LERKE/5/KAUZ/3/MYNA/VUL//BUC/FLK/4/MILAN	/6/.217	24.0	54.5	25.1	16.5
L61	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN/OTUS		30.0			14.2
L62	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN/OTUS		26.0	40.8	25.0	16.9
L63	PFAU/MILAN//TROST/3/PBW65/2*SERI.1B*2/4/	2.83	27.0	37.9	29.3	18.0
L64	PFAU/MILAN//TROST/3/PBW65/2*SERI.1B*2/4/		37.0			
L65	DOY1/AE.SQUARROSA (447)/3/KA/NAC//TRCH/4/	3.44	31.0	39.8	25.7	14.9
L66	MON/IMU//ALD/PVN/3/BORL95/4/OASIS/2*BORL9	5/	32.0			16.8
L67	WHEAR/SOKOLL/4/PASTOR//MILAN/KAUZ/3/BAV	92 4.79	39.0	44.9	27.5	14.6
L68	SOKOLL/WBLL1/4/D67.2/PARANA 66.270//	3.14	27.0	36.1	30.8	19.1
L69	MILAN//PRL/2*PASTOR/4/CROC_1/AE.SQUARROS	A	33.0			17.4
	(2					
L7	MUU/FRNCLN	2.40	25.0	44.9	26.3	18.9
L70	TRCH/SRTU//KACHU/3/KINGBIRD #1	3.58	28.0	44.6	23.7	15.8
L71	TRCH/SRTU//KACHU/3/KINGBIRD #1	3.98	47.0	42.7	22.8	15.2
L72	TRCH/5/BAV92//IRENA/KAUZ/3/HUITES/4/DOLL		26.0			15.6
L73	BECARD #1/3/PBW343*2/KUKUNA//PBW343*2/KUKUNA	3.87	35.0	39.7	28.8	16.4
L74	DANPHE/3/PBW343*2/KUKUNA//PBW343*2/KUKU	NA3.28	27.0	37.1	24.5	16.0
L76	PBW343*2/KUKUNA//PBW343*2/KUKUNA/6/WBLI		33.0	42.7	25.7	14.3
L77	BECARD/6/FRET2*2/4/SNI/TRAP#1/3/KAUZ*2/TRA		36.0	45.1	26.3	17.6
L78	KACHU/6/YAR/AE.SQUARROSA (783)/4/GOV/AZ//	2.87	28.0	42.7	23.7	11.7
L79	FRANCOLIN #1/NELOKI	4.12	37.0	39.4	31.4	22.1
L8	PRL/ 2*PAS-	3.90	28.0		21.6	16.3
	TOR/4/CHOIX/STAR/3/HE1/3*CNO79//					
L80	PAURAQ/8/NG8201/KAUZ/4/SHA7//PRL/VEE#6/3/		27.0	42.5	27.1	19.7
L81	PRL/2*PASTOR//SUNSTATE/3/GRACK	1.05	24.0	33.5	24.3	16.7
L82	PICAFLOR #1/NELOKI	3.63	40.0	37.9	24.4	16.8

Table 4.1: Model adjusted genotypic mean responses of yield and yield component traits (*continued*)

encoding	Genotype	Yield	Effective tillers	Thousand kernel weight	Grain per panicle	Panicle length
L83	PICAFLOR		32.0			15.7
	#1/8/NG8201/KAUZ/4/SHA7//PRL/VEE#6/					
L84	NL971*2/MUU	3.83	46.0	35.5	25.2	15.6
L85	MUNAL	4.79	40.0	44.2	27.8	17.7
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/					
L86	MUNAL	4.15	26.0	41.6	26.0	15.7
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/					
L87	MUNAL	3.67	37.0	46.1	27.1	15.9
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/					
L88	MUNAL		23.0			16.5
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/					
L89	BECARD #1*2/KINGBIRD #1		41.0			
L9	DANPHE #1*2/SHORTENED SR26	3.14	36.0	34.9	28.9	16.3
	TRANSLOCATION					
L90	BECARD*2/PFUNYE #1	2.91	41.0	38.2	28.0	16.2
L91	PBW343/PASTOR//OTUS/TOBA97*2/3/PICAFLOR	1.27	28.0	35.1	24.9	15.9
	#1					
L92	ROLF07/KINGBIRD #1//MUNAL #1	4.68	34.0	38.2	29.2	16.4
L93	MUNAL	4.83	30.0	43.1	26.2	15.3
	#1/3/PBW343*2/KUKUNA*2//YANAC					
L94	SERI.1B*2/3/KAUZ*2/BOW/KAUZ/4/PBW343*2/	2.07	37.0	27.0	24.2	15.4
L95	KAUZ//ALTAR		28.0			15.4
	84/AOS/3/MILAN/KAUZ/4/SAUAL/5/					
L96	PBW343*2/KHVAKI//PARUS/3/PBW343/PASTOR/5/	1.88	18.0	37.1	22.7	15.5
L97	FRET2*2/4/SNI/TRAP#1/3/KAUZ*2-	1.46	27.0	32.5	26.4	15.9
	TRAP//KAUZ*2/					
L98	ATTILA*2/PBW65*2//JUCHI/3/KINGBIRD		27.0			19.5
	#1/4/					
L99	MUNAL #1*2//WBLL1*2/BRAMBLING	3.63	36.0	49.3	25.7	15.1

4.1.2 Mixed model summary of fixed effects terms of yield and yield component traits

			Dependent variable	e:	
	Yield	Number of effective tillers	linear mixed-effects Thousand kernel weight	Grains per pani- cle	Panicle length
	(1)	(2)	(3)	(4)	(5)
Bhrikuti	$0.52^{***} (0.14)$ p = 0.0003	$3.34^{**} (1.35)$ p = 0.02	1.17 (0.85) p = 0.17	-0.22 (0.49) p = 0.66	0.51 (0.33) p = 0.13
Gautam	$0.31^{**} (0.14)$ p = 0.03	0.33 (1.36) p = 0.81	$2.06^{**} (0.85)$ $p = 0.02$	$1.33^{***} (0.49)$ p = 0.01	$1.22^{***} (0.33)$ $p = 0.0003$
Tilottama	$0.40^{***} (0.14)$ p = 0.005	$6.94^{***} (1.36)$ p = 0.0000	-0.32 (0.85) p = 0.72	0.38 (0.49) p = 0.45	0.27 (0.33) p = 0.43
Aditaya (Constant)	$3.02^{***} (0.61)$ p = 0.0000	$26.80^{***} (2.45)$ $p = 0.00$	$36.40^{***} (4.32)$ $p = 0.00$	$27.00^{***} (1.88)$ $p = 0.00$	$16.30^{***} (1.09)$ $p = 0.00$
Observations	202	238	201	202	226
Log Likelihood	-209.00	-752.00	-575.00	-452.00	-414.00
Akaike Inf. Crit.	443.00	1,528.00	1,174.00	928.00	853.00
Bayesian Inf. Crit.	482.00	1,570.00	1,213.00	968.00	894.00

Note:

*p<0.1; **p<0.05; ***p<0.01

Yield

Genotypes show highly significant differences for the yield trait. The model with both genotypes (checks and entries) outperforms in comparison to the one without in predicting the real yield ($\chi^2 = 14.4$ with df = 4).

Pairwise comparison of fixed effects estimate indicate that check variety Aditya has the lowest yield amongst all varieties while other have similar yields. Average yield of the lowest yielder is $4.53 \ tons \ ha^{-1} \ (\pm \ 0.60)$ and Bhrikuti yields the highest with average yield of $5.04 \ tons \ ha^{-1} \ (\pm \ 0.60)$.

Similarly, entry genotypes show considerable amount of heritable variation as evident from a large measure of variance in random effects term. ANOVA table (A.1) and the dotplot (Figure 4.2a) showing random effects estimates of the entry genotypes illustrate the relationship. Blocking structure, however, do not result in improvement of the effect estimates of the genotypes; blocking factors do not have significant effect on the yield trait.

Table 4.3: Mean differences in Yield of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	-0.516	0.139	101.0	-3.71	-0.792	-0.240	0.000
Aditya-Gautam	-0.305	0.140	99.9	-2.18	-0.583	-0.027	0.032
Aditya-Tilottama	-0.400	0.140	98.8	-2.86	-0.677	-0.122	0.005
Bhrikuti-Gautam	0.211	0.140	99.9	1.51	-0.066	0.489	0.134
Bhrikuti-Tilottama	0.116	0.140	99.0	0.83	-0.161	0.394	0.407
Gautam-Tilottama	-0.095	0.141	99.8	-0.67	-0.374	0.185	0.503

Number of effective tillers

Genotypes show highly significant differences for the Number of effective tillers trait. LRT shows that both checks and entries possess heritable variation that could potentially be exploited in further selection of the germplasm. The model comparison indicates a lower BIC value and χ^2 statistic of 35.2 with 4 df.

Pairwise comparison of fixed effects estimate indicate that except for Aditya and Gautam varieties, which are similar in Number of effective tillers, all other check varieties differ for the trait. Highest

number of effective tillers was found in Tilottama (33.73 \pm 2.46) and the least in Aditya (26.80 \pm 2.45) varieties.

In contrast, entry genotypes are not significantly different for the variability they contribute in the number of effective tillers. This can be seen as small spread in genotype estimates of the random effects term. ANOVA table (A.2) and the dotplot (Figure 4.3) showing random effects estimates of the entry genotypes illustrate the relationship. None of the blocking components might add to the benefit of improving the estimates of the genotypic effects for the number of effective tillers trait (based on LRT).

Table 4.4: Mean differences in Number of effective tillers of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	-3.340	1.35	121	-2.47	-6.015	-0.664	0.015
Aditya-Gautam	-0.328	1.36	121	-0.24	-3.026	2.369	0.810
Aditya-Tilottama	-6.939	1.36	120	-5.09	-9.637	-4.242	0.000
Bhrikuti-Gautam	3.011	1.36	120	2.21	0.316	5.707	0.029
Bhrikuti-Tilottama	-3.600	1.36	120	-2.65	-6.293	-0.906	0.009
Gautam-Tilottama	-6.611	1.37	120	-4.82	-9.325	-3.897	0.000

Thousand kernel weight

Genotypes are significantly different (at p=0.05) for the Thousand kernel weight trait. LRT shows that trait variation has the genetic roots linked to the genotype (for both checks and entries). Relative advantage of including genotype in the model terms is also highlighted by a lower BIC value and χ^2 statistic of 10.279 with $4 \, df$.

Pairwise comparison of fixed effects estimate indicate that Gautam variety has the highest thousand kernel weight (38.48 \pm 4.32 gm) and Tilottama has the least (36.103 \pm 4.32 gm) among check varieties.

Likewise, entry genotypes possessing a considerable amount of heritable variation can be evident from the ANOVA table (A.3) and the dotplot (Figure 4.2c showing random effects estimates of the entry genotypes illustrate the relationship. Besides genotype, blocking along the columns nested inside the column groups had a significant role in determination of Thousand kernel weight.

0.316

-0.894

1.486

2.380

2.009

0.800

3.176

4.085

0.712

0.297

0.084

0.007

Contrast Estimate Standard Error DF t-value Lower CI Upper CI p-value 90.6 0.514 -1.1700.847 -1.38-2.8530.171 Aditya-Bhrikuti -2.0640.853 90.1 -2.42 -3.759-0.3690.018 Aditya-Gautam

89.2

89.9

89.1

89.8

0.37

-1.05

1.75

2.77

-1.377

-2.588

-0.204

0.675

Table 4.5: Mean differences in Thousand kernel weight of check varieties (post-hoc comparison using Tukey procedure)

0.852

0.852

0.851

0.858

Number of grains per panicle

Aditya-Tilottama

Bhrikuti-Gautam

Bhrikuti-Tilottama

Gautam-Tilottama

Genotypes show significant differences for the Number of grains per panicle trait. LRT shows that both checks and entries possess significant (χ^2 statistic=11.59 with 4 df) heritable variation.

Pairwise comparison of fixed effects estimate indicate that check varieties Aditya and Gautam have different mean Number of grains per panicle. Likewise pairwise contrast using same procedure (Tukey) indicates different means for Bhrikuti and Gautam varieties. The highest and the lowest number of grains per panicle was found in Gautam (28.36 ± 1.88) and Bhrikuti (26.80 ± 1.88) varieties, respectively.

Similarly, entry genotypes show considerable amount of heritable variation as evident from a large measure of variance in random effects term. ANOVA table (A.4) and the dotplot (Figure 4.2d) showing random effects estimates of the entry genotypes illustrate the relationship. Likewise, no significant effects were associated with either of the blocking factors.

Table 4.6: Mean differences in Number of grains per panicle of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	0.221	0.488	120	0.45	-0.745	1.187	0.651
Aditya-Gautam	-1.331	0.492	121	-2.70	-2.305	-0.356	0.008
Aditya-Tilottama	-0.377	0.492	121	-0.77	-1.351	0.598	0.446
Bhrikuti-Gautam	-1.552	0.492	120	-3.16	-2.525	-0.578	0.002
Bhrikuti-Tilottama	-0.598	0.491	120	-1.22	-1.571	0.375	0.226
Gautam-Tilottama	0.954	0.496	121	1.92	-0.027	1.935	0.057

Panicle length

Genotypes show highly significant differences for the panicle length trait. LRT shows that both checks and entries possess significant (χ^2 statistic=14.62 with 4 df) heritable variation.

Pairwise comparison of fixed effects estimate indicate that check variety Gautam has significantly longer panicles than (all) Aditya, Tilottama and Bhrikuti. Likewise pairwise contrast using same procedure (Tukey) indicates that no difference exists between other varieties. Longest panicles were found in Gautam (17.53 \pm 1.09cm) and the shortest in Aditya (16.32 \pm 1.09cm) varieties.

Similarly, entry genotypes show considerable amount of heritable variation as evident from a large measure of variance in random effects term. ANOVA table (A.5) and the dotplot (Figure 4.3) showing random effects estimates of the entry genotypes illustrate the relationship. No attributable blocking effects, based on LRT, can be inferred about for the panicle length trait, alike most other yield and yield component traits.

Table 4.7: Mean differences in Panicle length of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	-0.510	0.331	129	-1.54	-1.165	0.146	0.126
Aditya-Gautam	-1.215	0.334	129	-3.64	-1.876	-0.555	0.000
Aditya-Tilottama	-0.266	0.334	129	-0.80	-0.926	0.394	0.427
Bhrikuti-Gautam	-0.705	0.334	129	-2.11	-1.366	-0.045	0.036
Bhrikuti-Tilottama	0.244	0.334	129	0.73	-0.416	0.904	0.466
Gautam-Tilottama	0.949	0.336	129	2.83	0.284	1.614	0.006

4.1.3 Mean comparison of yield and yield component traits of entry genotypes

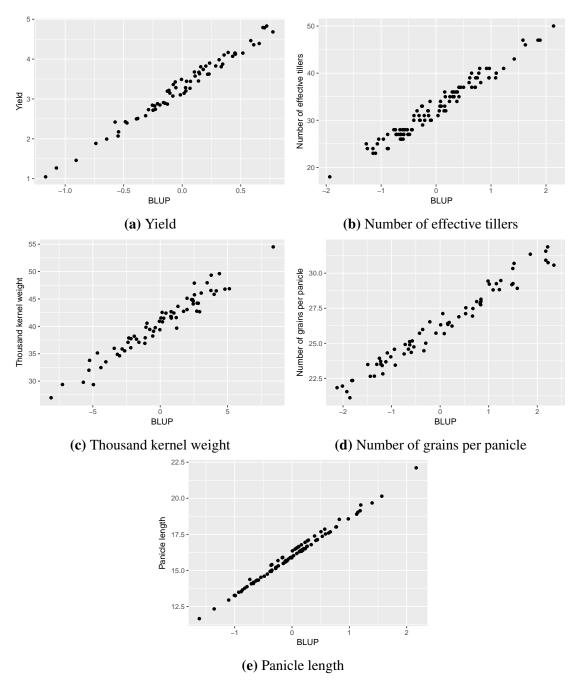


Figure 4.1: Scatterplot of observed versus BLUP values of yield and yield component traits

Scatterplot showing observed mean values of yield and yield component traits plotted against their respective genotypic BLUP values is presented in Figure 4.1.

The coefficient of determination, as computed by squared correlation coefficient between the observed and fitted values of random effects model are 0.61, 0.36, 0.71, 0.55, 0.48, respectively for the Yield, Number of effective tillers, Thousand kernel weight, Number of grains per panicle and Panicle length.

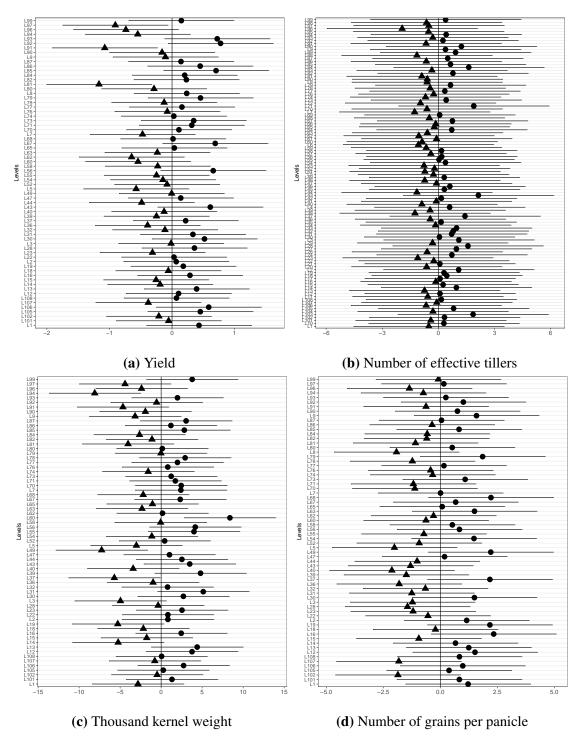


Figure 4.2: Random effect estimates of yield and yield component traits of entry genotypes

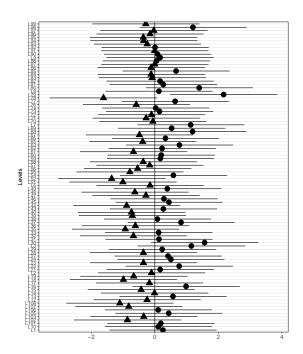


Figure 4.3: Random effect estimates of yield and yield component traits of entry genotypes (...continued)

4.2 Modeling leaf health traits

4.2.1 Model estimates of leaf health traits

Table 4.8: Model adjusted genotypic mean responses of leaf health traits

encoding	Genotype	Disease (Zadok's 75)	Greenness (Zadok's 65)	Greenness (Zadok's 75)	Greenness (Zadok's 85)	LAUG (Zadok's 65-75)	LAUG (Zadok's 75-85)
1	Aditya	2.86	8.07	7.33	5.83	110.8	92.89
2	Bhrikuti	3.32	8.01	6.66	5.20	108.4	86.28
3	Gautam	2.45	7.92	7.18	5.14	109.8	84.78
4	Tilottama	3.36	7.95	6.51	4.89	107.2	77.56
L1	MUNAL #1	4.00	8.13	5.63	5.42	111.3	68.27
L10	MUNAL/3/HUW234+LR34/PRINIA//PFAU/	WEAV E(RO) 4/	8.84	3.98	3.53	101.3	56.60
L101	PFAU/MILAN/5/CHEN/AEGILOPS SQUARROSA (TAUS)/	2.00	8.84	4.89	4.68	93.2	47.19
L102	PREMIO/5/TUI//2*SUNCO/SA1166/3/TUI/4	/FINS # .00	8.75	7.01	6.52	103.2	95.55
L103	PREMIO/BAVIS	2.50	7.73	5.81	3.98	99.6	55.47
L104	CHIYAK//ND643/2*WAXWING/3/ND643/2	*WAX 6 WOONG	7.45	3.43	0.65	83.5	30.94
L105	BABAX/LR42//BABAX*2/4/SNI/TRAP#1/3/	KAUZ#: 5 0	7.80	5.41	4.21	104.6	65.47
L106	PFAU/MILAN//TROST/3/MUNAL #1/4/PFAU/MILAN//	1.33	8.35	7.96	6.97	120.1	92.18
L107	YUK/AE.SQUARROSA (217)//2*PANDORA	6.00	7.92	6.36	5.01	102.7	92.90
L108	SUP152*2//ND643/2*WAXWING	4.00	8.84	4.06	4.02	100.4	42.28
L11	SWSR22T.B./5/KAUZ//ALTAR 84/AOS/3/KAUZ/4/	2.00	8.84	5.97	6.06	113.1	86.15
L12	WHEAR/VIVITSI//WHEAR/3/PANDORA	1.00	7.86	4.46	3.34	98.2	53.31
L13	WBLL1*2/KURUKU//TACUPETO F2001*2/BRAMBLING	1.50	7.79	7.56	5.17	113.3	98.78
L14	PASTOR/3/VORONA/CN079//KAUZ/4/MIL	AN/OTAU89/	8.60	5.77	5.51	107.2	72.10
L15	FRANCOLIN #1*2/ND643/2*WBLL1	3.50	8.67	4.23	3.97	90.3	60.03
L16	KIRITATI/ 2*WBLL1/3/TAM200/PASTOR//TOBA97/4/.	5.00	8.10	5.01	4.04	94.2	63.07
L17	SUP152/KENYA SUNBIRD	2.00	7.23	4.77	0.94	90.5	20.16
L18	SUP152/KENYA SUNBIRD	5.50	8.67	3.19	1.64	82.8	32.26
L19	SUP152/KENYA SUNBIRD	2.50	8.00	3.83	2.26	88.7	32.11
L2	KACHU #1	7.00	7.77	7.48	4.43	110.5	86.48
L20	WBLL1*2/BRAMBLING/5/BABAX/LR42//F	BABA K.02 /4/	8.59	5.77	5.56	112.9	78.65
L21	ND643/2*WBLL1/3/KIRITATI//2*PRL/2*PA	STOR2400.	8.49	6.49	6.31	111.9	85.33
L22	BETTY/3/CHEN/AE.SQ//2*OPATA/4/BAVIS	7.00	8.13	4.77	4.49	103.8	58.23
L23	BAVIS/VORB/5/CROC_1/AE.SQUARROSA	(205)/600	7.50	1.84	0.57	62.6	9.13
L24	BABAX/LR42//BABAX/3/ER2000/4/NAVJ07	7 1.50	8.67	4.03	3.73	92.4	42.57

Table 4.8: Model adjusted genotypic mean responses of leaf health traits (*continued*)

encoding	Genotype	Disease (Zadok's 75)	Greenness (Zadok's 65)	Greenness (Zadok's 75)	Greenness (Zadok's 85)	LAUG (Zadok's 65-75)	LAUG (Zadok's 75-85)
L26	NING-MAI-	7.00	7.60	6.61	3.61	107.5	73.19
	96035/FINSI//HEILO/3/NAVJ07						
L27	WBLL1*2/BRAMBLING//TAM200/TUI/3/	1.50	8.13	3.19	2.30	83.1	40.10
L28	KISKADEE	3.50	8.54	4.17	4.15	98.6	75.38
	#1/5/KAUZ*2/MNV//KAUZ/3/MILAN/4/						
L29	MERCATO//PARUS/PASTOR	1.50	8.40	7.56	7.60	125.2	100.35
L3	KACHU/BECARD//WBLL1*2/BRAMBLING	1.50	8.57	6.71	6.58	118.5	76.82
L30	PREMIO//PARUS/PASTOR	1.00	8.13	5.27	5.26	98.7	88.29
L31	ND643/2*WBLL1//ND643/2*WAXWING	2.50	8.74	6.81	6.66	115.8	96.52
L32	ND643/2*WBLL1//ND643/2*WAXWING	2.50	8.59	5.93	5.53	103.1	81.34
L33	ND643/2*WBLL1//ND643/2*WAXWING	2.50	8.84	6.23	6.07	109.1	95.29
L34	SNB//CMH79A.955/3*CN079/3/ATTILA/4/CH	EN/2.50	8.67	2.32	2.18	80.4	13.88
L35	KIRITATI/ 2*WBLL1/8/SHA7/ PRL/VEE#6/ 3/ FASAN/	7.00	8.01	5.06	4.54	98.9	59.09
L36	SERI.1B*2/3/KAUZ*2/BOW//KUZ/4/CIRCUS/	5/3.50	7.69	3.51	1.38	81.4	35.51
L37	WHEAR/KUKUNA/3/C80.1/3*BATAVIA//2*W	BL1L10/04/	7.99	3.04	1.20	88.3	29.41
L38	BAJ #1*2/BECARD	6.50	8.47	5.94	5.79	97.5	82.33
L39	KACHU*2/MUNAL #1	2.00	8.22	6.41	6.32	104.4	93.39
L4	PRL/2*PASTOR/3/PFAU/WEAVER*2//CHAPIO	O 4.00	8.34	4.74	4.66	99.0	61.64
L40	KACHU*2/MUNAL #1	1.00	8.50	7.27	5.94	112.4	100.07
L41	KACHU/2*MUNAL #1	3.50	8.00	4.86	4.26	92.4	60.84
L42	KACHU/2*MUNAL #1	6.00	8.60	6.05	6.04	109.8	81.77
L43	KACHU*2/PANDORA	2.00	7.84	5.81	4.72	107.6	69.27
L44	SUP152/2*MUNAL #1	4.50	7.79	5.81	4.30	87.7	74.82
L45	FRNCLN*2/BECARD	3.50	8.63	5.46	5.12	103.6	78.43
L46	WBLL1* 2/BRAM-	2.00	7.93	4.46	3.79	89.8	60.63
	BLING//SAAR/2*WAXWING/4/						
L47	CHIBIA//PRLII/CM65531/3/MISR 2*2/4/	3.00	8.36	6.67	6.03	111.3	90.87
L48	CHIBIA//PRLII/CM65531/3/MISR 2*2/4/	2.00	8.13	5.17	4.59	97.7	74.98
L49	WAXWING/KRONSTAD F2004//2*FRNCLN	3.00	8.34	5.26	5.06	100.0	64.64
L5	DANPHE/CHONTE	1.00	8.16	3.23	3.14	74.3	40.74

Table 4.8: Model adjusted genotypic mean responses of leaf health traits (continued)

encoding	Genotype (Disease Zadok's 75)	Greenness (Zadok's 65)	Greenness (Zadok's 75)	Greenness (Zadok's 85)	LAUG (Zadok's 65-75)	LAUG (Zadok's 75-85)
L51	MUNAL	6.00	8.40	4.46	3.09	95.1	38.65
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/						
L52	MUNAL	4.00	8.42	4.43	3.96	89.9	53.01
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/						
L53	MUNAL #1/2*FRNCLN	2.50	8.67	5.89	5.65	111.2	86.07
L54	MUNAL #1/2*FRNCLN	4.00	7.90	5.57	5.04	100.2	67.48
L55	MUNAL #1*2//WHEAR/SOKOLL	3.00	8.14	3.99	3.53	88.2	51.74
L56	MUNAL #1*2/KINGBIRD #1	6.00	8.25	4.06	3.76	96.4	53.13
L57	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	3.50	8.67	4.87	4.85	99.0	65.52
L58	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	5.50	8.50	5.66	5.46	103.6	66.97
L59	KAUZ*2/BOW//KAUZ/3/W98.6.38/5/SABUF/4	/ 3.00	8.09	5.77	5.62	107.2	70.23
L6	SUP152/FRNCLN	8.00	7.85	6.21	4.42	106.4	71.68
L60	LERKE/5/KAUZ/3/MYNA/VUL//BUC/FLK/4/N	MIL 5\00 6/	8.13	5.66	4.89	93.4	95.86
L61	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN	/OTAL089/	8.19	3.83	3.46	92.3	63.58
L62	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN	/OT2L58)/	8.60	5.97	4.66	108.4	79.00
L63	PFAU/MILAN//TROST/3/PBW65/2*SERI.1B*2	2/4/.6.00	8.53	5.77	5.52	102.1	81.67
L64	PFAU/MILAN//TROST/3/PBW65/2*SERI.1B*2	2/4/.300	8.51	5.91	5.85	103.0	96.05
L65	DOY1/AE.SQUARROSA (447)/3/KA/NAC//TRCH/4/	3.00	8.34	3.94	4.08	96.9	47.40
L66	MON/IMU//ALD/PVN/3/BORL95/4/OASIS/2*F	BORL5905/	8.69	5.61	5.55	108.6	61.30
L67	WHEAR/SOKOLL/4/PASTOR//MILAN/KAUZ/	/3/BA \$ @2	8.50	6.78	4.95	118.6	95.13
L68	SOKOLL/WBLL1/4/D67.2/PARANA 66.270//	8.00	8.38	5.57	5.48	107.6	70.25
L69	MILAN//PRL/2*PASTOR/4/CROC_1/AE.SQUA (2	ARROMA	7.95	4.54	3.79	98.5	61.01
L7	MUU/FRNCLN	2.50	8.84	3.17	3.10	82.2	47.39
L70	TRCH/SRTU//KACHU/3/KINGBIRD #1	2.50	8.56	5.51	5.10	107.8	64.83
L71	TRCH/SRTU//KACHU/3/KINGBIRD #1	6.00	7.71	5.03	4.53	122.3	58.03
L72	TRCH/5/BAV92//IRENA/KAUZ/3/HUITES/4/D	OL b .00	8.13	5.26	4.80	96.3	77.78
L73	BECARD #1/3/PBW343*2/KUKUNA//PBW343*2/KUKU	2.50 NA	7.75	6.21	4.78	105.2	80.47
L74	DANPHE/3/PBW343*2/KUKUNA//PBW343*2/	/KU4KOONA	8.60	4.31	3.69	94.4	62.11
L76	PBW343*2/KUKUNA//PBW343*2/KUKUNA/6	5/W BБ0 .1*2/	8.75	3.99	3.85	91.4	31.34

Table 4.8: Model adjusted genotypic mean responses of leaf health traits (continued)

encoding	Genotype	Disease (Zadok's 75)	Greenness (Zadok's 65)	Greenness (Zadok's 75)	Greenness (Zadok's 85)	LAUG (Zadok's 65-75)	LAUG (Zadok's 75-85)
L77	BECARD/6/FRET2*2/4/SNI/TRAP#1/3/KAV	UZ*2/T1R0A0P/	7.73	5.26	2.84	100.7	60.67
L78	KACHU/6/YAR/AE.SQUARROSA (783)/4/GOV/AZ//	3.00	8.14	5.49	5.19	105.7	81.74
L79	FRANCOLIN #1/NELOKI	1.83	7.66	5.17	2.03	100.4	25.22
L8	PRL/ 2*PAS-	7.00	4.67	2.00	0.80	62.1	10.86
T 00	TOR/4/CHOIX/STAR/3/HE1/3*CNO79//	VEE 11/2 1/20100	0.00	5.24	5.10	100.7	70.72
L80	PAURAQ/8/NG8201/KAUZ/4/SHA7//PRL/V		8.80	5.34	5.19	100.7	70.73
L81	PRL/2*PASTOR//SUNSTATE/3/GRACK	4.00	7.97	3.35	0.75	81.9	34.31
L82	PICAFLOR #1/NELOKI	0.50	8.45	4.23	3.02	98.7	62.43
L83	PICAFLOR #1/8/NG8201/KAUZ/4/SHA7//PRL/VEE#6/.	2.50	8.51	5.44	5.32	104.0	72.11
L84	NL971*2/MUU	3.00	8.84	6.95	6.14	124.6	102.69
L85	MUNAL	2.00	8.50	4.84	4.68	105.3	66.69
T 06	#1*2/4/HUW234+LR34/PRINIA//PBW343*		7.70	5.66	4.00	104.5	02.10
L86	MUNAL #1*2/4/HUW234+LR34/PRINIA//PBW343*.	7.00 2/	7.79	5.66	4.09	104.5	92.19
L87	MUNAL	1.50	8.24	4.56	4.46	103.3	75.77
	#1*2/4/HUW234+LR34/PRINIA//PBW343*	2/					
L88	MUNAL	2.00	8.80	5.91	5.65	104.9	81.74
	#1*2/4/HUW234+LR34/PRINIA//PBW343*	2/					
L89	BECARD #1*2/KINGBIRD #1	3.00	8.84	4.37	4.31	101.1	45.62
L9	DANPHE #1*2/SHORTENED SR26 TRANSLOCATION	4.50	8.50	4.46	4.01	104.4	56.80
L90	BECARD*2/PFUNYE #1	6.00	8.38	4.36	4.31	96.0	56.78
L91	PBW343/PASTOR//OTUS/TOBA97*2/3/PIC		8.84	0.70	0.20	71.5	21.37
271	#1		0.0.	0.70	0.20	71.0	21.07
L92	ROLF07/KINGBIRD #1//MUNAL #1	4.17	7.63	3.27	0.69	91.1	30.59
L93	MUNAL	5.00	8.75	4.33	3.53	98.3	47.50
	#1/3/PBW343*2/KUKUNA*2//YANAC						
L94	SERI.1B*2/3/KAUZ*2/BOW/KAUZ/4/PBW	343*2/2.50	8.67	3.43	1.54	97.1	21.87
L95	KAUZ//ALTAR	3.00	8.06	3.43	2.09	92.1	43.18
	84/AOS/3/MILAN/KAUZ/4/SAUAL/5/						
L96	PBW343*2/KHVAKI//PARUS/3/PBW343/PA	ASTOR 25 00.	8.67	4.26	4.14	86.2	53.83
L97	FRET2*2/4/SNI/TRAP#1/3/KAUZ*2- TRAP//KAUZ*2/	8.00	8.09	2.94	1.10	84.5	16.52

Table 4.8: Model adjusted genotypic mean responses of leaf health traits (continued)

encoding	Genotype	Disease (Zadok's 75)	Greenness (Zadok's 65)	Greenness (Zadok's 75)	Greenness (Zadok's 85)	LAUG (Zadok's 65-75)	LAUG (Zadok's 75-85)
L98	ATTILA*2/PBW65*2//JUCHI/3/KINGBIRD #1/4/	0.50	7.76	2.95	0.70	78.3	5.25
L99	MUNAL #1*2//WBLL1*2/BRAMBLING	1.50	8.84	1.91	0.57	84.6	11.86

4.2.2 Mixed model summary of fixed effects terms of leaf health traits

			Depender	nt variable:		
				near l-effects		
	Disease score zadok 75	Greenness1	Greenness2	Greenness3	LAUG1	LAUG2
	(1)	(2)	(3)	(4)	(5)	(6)
Bhrikuti	0.47* (0.26) p = 0.07	-0.07 (0.08) p = 0.37	$-0.67^{***} (0.17)$ p = 0.0001	$-0.67^{**} (0.26)$ p = 0.02	-2.62 (1.77) p = 0.14	-6.97** (3.33) p = 0.04
Gautam	-0.33(0.26)	$-0.15^* (0.08)$ $p = 0.06$	-0.17 (0.17) p = 0.31	$-0.71^{***} (0.26)$ $p = 0.01$	-1.23(1.79)	-8.35**(3.36)
Tilottama	$p = 0.20$ $0.50^* (0.26)$	-0.12(0.08)	-0.83^{***} (0.17)	-0.93**** (0.26)	$p = 0.49$ $-3.76^{**} (1.79)$	$p = 0.02$ $-15.70^{***} (3.36)$
Aditaya (Constant)	p = 0.06 $2.85^{**} (1.30)$ p = 0.03	p = 0.11 8.07*** (0.43) p = 0.00	$p = 0.0000$ $7.33^{***} (1.18)$ $p = 0.00$	p = 0.0005 5.83^{***} (1.23) p = 0.0000	$p = 0.04$ $111.00^{***} (9.21)$ $p = 0.00$	p = 0.0000 93.00*** (19.70) p = 0.0000
Observations	238	238	238	238	238	238
Log Likelihood	-414.00	-128.00	-328.00	-410.00	-859.00	-1,013.00
Akaike Inf. Crit. Bayesian Inf. Crit.	852.00 894.00	280.00 322.00	680.00 722.00	844.00 886.00	1,742.00 1,784.00	2,050.00 2,092.00

Note:

*p<0.1; **p<0.05; ***p<0.01

Disease score at Zadok's stage 75

Genotypes show highly significant differences for the disease resistance trait. LRT shows that both checks and entries possess significant (χ^2 statistic=13.85 with 4 df) heritable variation.

Pairwise comparison of fixed effects estimate indicate that check variety Gautam is relatively more asymptomatic (2.51 (\pm 1.30)) than Bhrikuti(3.32 (\pm 1.30)) and Tilottama (3.35 (\pm 1.30)) varieties to foliar pathogen. Although, no distinction could be made on the severity scale of the disease (All check varieties exhibited similarly less affected (2.5-5.0) symptom at the time of scoring).

Entry genotypes also show considerable amount of heritable variation evident from a large measure of random effects variance contributed by the genotypes. ANOVA table (A.6) and the dotplot (Figure 4.5a) showing random effects estimates of the entry genotypes illustrate the relationship. Similarly, blocking factor nested column within columgroup has a significant contribution to the variation in the disease score trait.

Table 4.10: Mean differences in Disease score at zadok's stage 75 of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	-0.473	0.256	97.7	-1.85	-0.981	0.034	0.067
Aditya-Gautam	0.333	0.258	97.0	1.29	-0.179	0.844	0.200
Aditya-Tilottama	-0.502	0.258	96.0	-1.95	-1.013	0.009	0.054
Bhrikuti-Gautam	0.806	0.258	96.7	3.13	0.295	1.317	0.002
Bhrikuti-Tilottama	-0.029	0.257	96.1	-0.11	-0.539	0.481	0.911
Gautam-Tilottama	-0.835	0.259	96.7	-3.22	-1.349	-0.321	0.002

Leaf greenness score at Zadok's stage 65

The LRT indicates that fixed effects due to check varieties have no significant effects on the leaf greenness score (p-value for χ^2 statistic > 0.05).

Entry genotypes are significantly variable for the trait. ANOVA table (A.7) and the dotplot (Figure 4.6) showing random effects estimates of the entry genotypes illustrate the relationship. Blocking factors, on the other hand, do not have significant effects in the variation for trait.

Table 4.11: Mean differences in Leaf greenness at zadok's stage 65 of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	0.069	0.076	107	0.91	-0.082	0.220	0.367
Aditya-Gautam	0.148	0.077	107	1.92	-0.005	0.300	0.058
Aditya-Tilottama	0.124	0.077	107	1.62	-0.028	0.277	0.108
Bhrikuti-Gautam	0.078	0.077	108	1.02	-0.074	0.231	0.310
Bhrikuti-Tilottama	0.055	0.077	107	0.72	-0.097	0.208	0.473
Gautam-Tilottama	-0.023	0.077	107	-0.30	-0.176	0.130	0.766

Leaf greenness score at Zadok's stage 75

Genotypes show highly significant differences in the greenness score measured measured at Zadok's stage 75. LRT shows that both checks and entries possess significant (χ^2 statistic=32.828 with 4 df) heritable variation.

Pairwise comparison of fixed effects estimate indicate that check variety Gautam and Bhrikuti are significantly different for the green appearence of the flag leaf. The lowest scores are for the check variety Gautam (6.51 (\pm 1.76)) and the highest are for the Tilottama (7.34 (\pm 1.76)) variety. Although, based on the qualitative description (Table 3.3), both can be characterized as having retained moderately green leaf coloration.

Similarly noteworthy is the variation among the entry genotypes (significant at p < 0.01). A measure of random effects variance contributed by the genotypes is shown in ANOVA Table (A.8) and the dotplot (Figure 4.5c) showing random effects estimates of the entry genotypes. Unlike entry effects, controlling for exraneous factors with blocking did not result in significant reduction in residual variance.

Table 4.12: Mean differences in Leaf greenness at zadok's stage 75 of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	0.674	0.166	104	4.07	0.346	1.003	0.000
Aditya-Gautam	0.170	0.167	103	1.02	-0.161	0.501	0.310
Aditya-Tilottama	0.829	0.167	102	4.98	0.499	1.160	0.000
Bhrikuti-Gautam	-0.504	0.167	103	-3.02	-0.835	-0.173	0.003
Bhrikuti-Tilottama	0.155	0.167	102	0.93	-0.176	0.485	0.355
Gautam-Tilottama	0.659	0.168	103	3.92	0.326	0.992	0.000

Leaf greenness score at Zadok's stage 85

Genotypes show highly significant differences in the greenness score measured at Zadok's stage 85. LRT shows that both checks and entries possess significant (χ^2 statistic=14.17 with 4 df) heritable variation.

Pairwise comparison of fixed effects estimate indicate that check variety Aditya (5.832 (\pm 1.23)) has the most green coloration of the flag leaf followed by Bhrikuti and Gautam (both have similar greenness scores) and Tilottama (4.90 (\pm 1.23)). Based on the qualitative description (Table 3.3), both can be characterized as having retained moderate amounts of green color in leaf.

Similarly noteworthy is the variation among the entry genotypes (significant at p < 0.01). A measure of random effects variance contributed by the genotypes is shown in ANOVA Table (A.9) and the dotplot (Figure 4.5d) showing random effects estimates of the entry genotypes. Likewise, effect of blocking in the direction along columnested inside column groups resulted in significant reduction of residual error variance.

Table 4.13: Mean differences in Leaf greenness at zadok's stage 85 of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	0.666	0.262	100.9	2.54	0.145	1.186	0.013
Aditya-Gautam	0.713	0.264	100.0	2.70	0.188	1.238	0.008
Aditya-Tilottama	0.928	0.264	98.9	3.51	0.404	1.453	0.001
Bhrikuti-Gautam	0.048	0.265	99.2	0.18	-0.478	0.573	0.858
Bhrikuti-Tilottama	0.263	0.264	98.8	1.00	-0.261	0.787	0.322
Gautam-Tilottama	0.216	0.266	99.4	0.81	-0.313	0.744	0.420

LAUG between Zadok's stage 65 and 75

The LRT indicates that fixed effects due to check varieties have no significant effects on the LAUG value (p-value for χ^2 statistic > 0.05).

However, entry genotypes do represent a diverse collection with the trait tracing it's roots in genetic make up of the diverse germplam. ANOVA table (A.10) and the dotplot (Figure 4.6a) showing random effects estimates of the entry genotypes illustrate the relationship. To the contrary, no significant effects of blocking factors on the traits LAUG between zadok's stage 65 and 75 was discovered.

Table 4.14: Mean differences in LAUG during zadok's stage 65 to 75 of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	2.62	1.77	101.5	1.48	-0.896	6.14	0.142
Aditya-Gautam	1.24	1.79	100.4	0.69	-2.310	4.78	0.491
Aditya-Tilottama	3.76	1.78	99.3	2.11	0.217	7.30	0.038
Bhrikuti-Gautam	-1.39	1.78	100.3	-0.78	-4.930	2.15	0.439
Bhrikuti-Tilottama	1.14	1.78	99.5	0.64	-2.401	4.67	0.525
Gautam-Tilottama	2.52	1.80	100.2	1.40	-1.042	6.09	0.163

LAUG between Zadok's stage 75 and 85

Genotypes show highly significant differences in the LAUG values for duration between Zadok's stage 75 and 85 (an indication of how much reduction in photosynthesis has occured do to decline in

green pigmentation). LRT shows that both checks and entries possess significant (χ^2 statistic=21.63 with 4 df) heritable variation.

Pairwise comparison of fixed effects estimate indicate that check variety Aditya suffered largest reduction in green pigmentation (93.03 (\pm 19.70)), while Tilottama had the least (77.35 (\pm 19.71)) loss of all check varieties.

Likewise, variation among the entry genotypes (significant at p < 0.01) for the trait. A measure of random effects variance contributed by the genotypes is shown in ANOVA Table (A.11) and the dotplot (Figure 4.6b) showing random effects estimates of the entry genotypes. Similar to the approximation of LAUG between Zadok's stage 65 to 75, the effects of blocking factors is not realized to be significant either in approximating LAUG during a later period.

Table 4.15: Mean differences in LAUG during zadok's stage 75 to 85 of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	6.97	3.33	103	2.09	0.358	13.58	0.039
Aditya-Gautam	8.35	3.36	103	2.48	1.684	15.02	0.015
Aditya-Tilottama	15.67	3.36	102	4.66	9.006	22.34	0.000
Bhrikuti-Gautam	1.38	3.36	101	0.41	-5.278	8.04	0.682
Bhrikuti-Tilottama	8.70	3.35	101	2.59	2.048	15.36	0.011
Gautam-Tilottama	7.32	3.38	102	2.16	0.613	14.03	0.033

4.2.3 Mean comparison of leaf health traits of entry genotypes

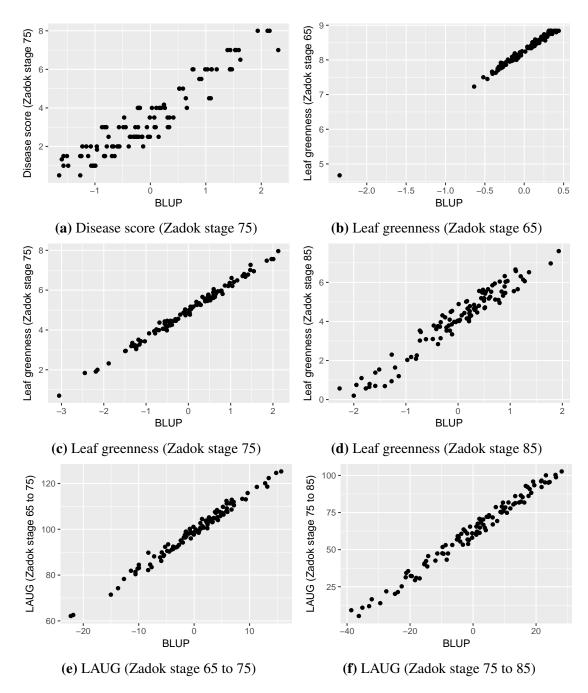


Figure 4.4: Scatterplot of observed versus BLUP values of leaf health traits

Scatterplot showing observed mean leaf health traits plotted against their respective genotypic BLUP values is presented in Figure 4.4.

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The coefficient of determination, as computed by squared correlation coefficient between the observed and fitted values of random effects model are 0.77, 0.73, 0.88, 0.72, 0.75, 0.79, respectively for the Disease score at Zadok's stage 75, Leaf greenness Zadok's stage 65, Leaf greenness Zadok's stage 75, Leaf greenness Zadok's stage 85, LAUG during Zadok's stage 65 to 75 and LAUG during Zadok's stage 75 to 85.

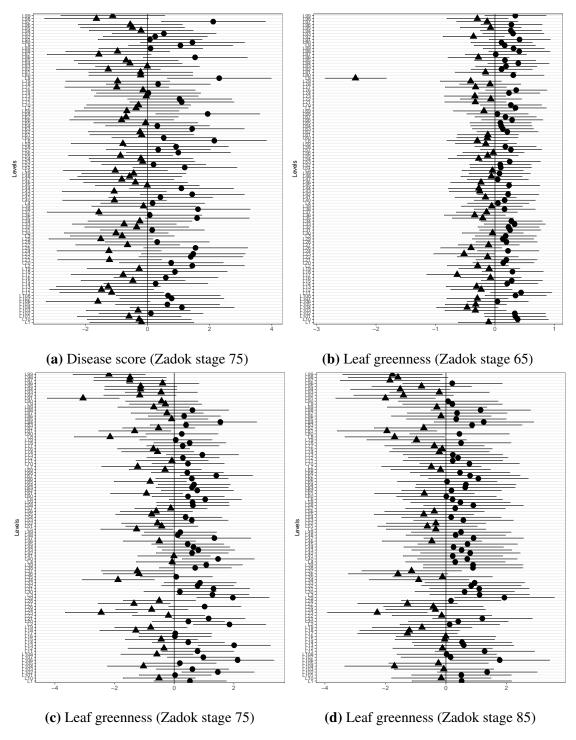


Figure 4.5: Random effect estimates of leaf health traits of entry genotypes

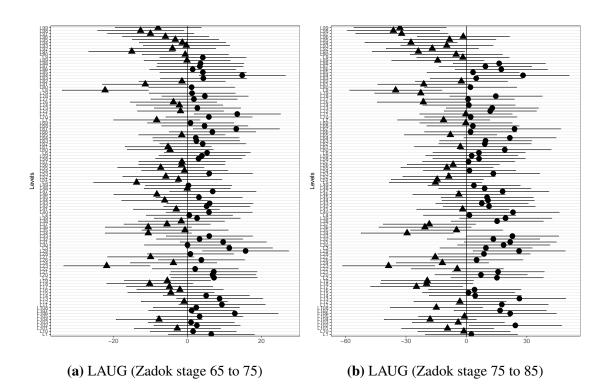


Figure 4.6: Random effect estimates of leaf health traits of entry genotypes (...continued)

- 4.3 Modeling plant morphological, architectural and phenological traits
- 4.3.1 Model estimates of plant morphological, architectural and phenological traits

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits

encoding	Genotype	Plant height	Leaf area	SPADI	SPADII
1	Aditya	103.9	37.5	46.0	38.8
2	Bhrikuti 94.6		38.8	44.2	35.0
3	Gautam	105.1	41.0	44.9	38.4
4	Tilottama	99.2	36.0	44.5	32.3
L1	MUNAL #1	107.5	41.5	48.3	43.8
L10	MUNAL/3/HUW234+LR34/PRINIA//PFAU/	WEA V162R 54/	33.0	43.9	37.4
L101	PFAU/MILAN/5/CHEN/AEGILOPS SQUARROSA (TAUS)/	95.0	25.7	51.8	37.2
L102	PREMIO/5/TUI//2*SUNCO/SA1166/3/TUI/	4/FINS 9 /45	29.2	48.9	42.3
L103	PREMIO/BAVIS	101.0	47.0	42.2	43.0
L104	CHIYAK//ND643/2*WAXWING/3/ND643/2		25.7	43.0	21.5
L105	BABAX/LR42//BABAX*2/4/SNI/TRAP#1/3	/KAU Z®3/0	40.9	42.7	27.5
L106	PFAU/MILAN//TROST/3/MUNAL #1/4/PFAU/MILAN//	95.3	37.6	55.0	44.6
L107	YUK/AE.SQUARROSA (217)//2*PANDORA	97.0	56.7	51.6	38.5
L108	SUP152*2//ND643/2*WAXWING	97.5	31.0	49.2	26.7
L11	SWSR22T.B./5/KAUZ//ALTAR 84/AOS/3/KAUZ/4/	93.0	26.2	42.6	37.3
L12	WHEAR/VIVITSI//WHEAR/3/PANDORA	99.0	33.9	50.6	46.4
L13	WBLL1*2/KURUKU//TACUPETO F2001*2/BRAMBLING	111.0	49.7	46.9	40.8
L14	PASTOR/3/VORONA/CN079//KAUZ/4/MIL	AN/OT0250/	43.7	40.6	40.9
L15	FRANCOLIN #1*2/ND643/2*WBLL1	89.0	50.4	48.9	31.9
L16	KIRITATI/ 2*WBLL1/3/TAM200/PASTOR//TOBA97/4/	101.0	55.0	46.6	39.2
L17	SUP152/KENYA SUNBIRD	99.0	40.0	45.4	44.5
L18	SUP152/KENYA SUNBIRD	95.0	41.5	39.8	23.3
L19	SUP152/KENYA SUNBIRD	96.5	40.3	42.2	39.2
L2	KACHU #1	91.0	40.1	42.4	37.4
L20	WBLL1*2/BRAMBLING/5/BABAX/LR42//	BABA % 3*: 0 /4/	35.8	44.6	32.9
L21	ND643/2*WBLL1/3/KIRITATI//2*PRL/2*PA	ASTORØ#.0	31.9	46.6	42.9
L22	BETTY/3/CHEN/AE.SQ//2*OPATA/4/BAVI	S 101.5	36.5	42.1	34.4
L23	BAVIS/VORB/5/CROC_1/AE.SQUARROSA	(205) // 14.0	23.5	36.8	27.8
L24	BABAX/LR42//BABAX/3/ER2000/4/NAVJ0	7 100.5	36.5	43.5	34.4
L26	NING-MAI- 96035/FINSI//HEILO/3/NAVJ07	101.0	31.0	46.5	40.3

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits *(continued)*

encoding	Genotype	Plant height	Leaf area	SPADI	SPADII
L27	WBLL1*2/BRAMBLING//TAM200/TUI/3/	103.5	41.9	41.6	39.5
L28	KISKADEE	92.5	56.1	44.9	39.0
	#1/5/KAUZ*2/MNV//KAUZ/3/MILAN/4/				
L29	MERCATO//PARUS/PASTOR	106.5	29.1	43.9	41.4
L3	KACHU/BECARD//WBLL1*2/BRAMBLING	106.0	36.9	50.1	39.0
L30	PREMIO//PARUS/PASTOR	95.5	51.7	51.5	43.3
L31	ND643/2*WBLL1//ND643/2*WAXWING	96.5	44.4	46.6	35.6
L32	ND643/2*WBLL1//ND643/2*WAXWING	97.0	32.9	46.6	33.9
L33	ND643/2*WBLL1//ND643/2*WAXWING	95.0	32.6	47.0	41.4
L34	SNB//CMH79A.955/3*CN079/3/ATTILA/4/CF	IEN/9.7.5	30.3	42.3	36.9
L35	KIRITATI/ 2*WBLL1/8/SHA7/	106.0	39.2	38.8	32.3
	PRL/VEE#6/ 3/ FASAN/				
L36	SERI.1B*2/3/KAUZ*2/BOW//KUZ/4/CIRCUS	/5/95.5	25.5	43.0	26.3
L37	WHEAR/KUKUNA/3/C80.1/3*BATAVIA//2*V	VBL 9d/3 l/	47.0	45.7	38.1
L38	BAJ #1*2/BECARD	96.5	36.5	39.9	21.0
L39	KACHU*2/MUNAL #1	99.0	41.0	42.2	39.1
L4	PRL/2*PASTOR/3/PFAU/WEAVER*2//CHAP	IO 95.0	42.5	46.6	43.9
L40	KACHU*2/MUNAL #1	97.0	36.8	45.9	45.6
L41	KACHU/2*MUNAL #1	98.0	36.8	45.0	27.0
L42	KACHU/2*MUNAL #1	106.5	32.2	45.6	28.4
L43	KACHU*2/PANDORA	95.5	40.2	45.4	34.7
L44	SUP152/2*MUNAL #1	100.0	35.1	45.6	24.1
L45	FRNCLN*2/BECARD	97.0	28.4	41.7	38.3
L46	WBLL1* 2/BRAM-	90.0	47.9	45.4	37.1
	BLING//SAAR/2*WAXWING/4/				
L47	CHIBIA//PRLII/CM65531/3/MISR	101.0	49.2	45.5	38.7
L48	2*2/4/ CHIBIA//PRLII/CM65531/3/MISR 2*2/4/	107.5	31.1	45.9	39.9
L49	WAXWING/KRONSTAD F2004//2*FRNCLN	96.5	41.8	41.1	38.3
L5	DANPHE/CHONTE	109.5	34.9	45.0	39.4
L51	MUNAL	105.0	24.3	42.4	32.3
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/				
L52	MUNAL #1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/	102.5	24.4	42.5	36.1

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits *(continued)*

encoding	Genotype	Plant height	Leaf area	SPADI	SPADII
L53	MUNAL #1/2*FRNCLN	85.0	46.1	44.2	20.2
L54	MUNAL #1/2*FRNCLN	91.5	31.4	50.0	38.3
L55	MUNAL #1*2//WHEAR/SOKOLL	94.0	38.3	42.6	29.7
L56	MUNAL #1*2/KINGBIRD #1	93.0	29.1	47.4	43.5
L57	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	98.5	36.4	43.5	24.9
L58	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	101.0	40.8	42.2	32.4
L59	KAUZ*2/BOW//KAUZ/3/W98.6.38/5/SABU	JF/4/105.0	54.3	47.8	33.9
L6	SUP152/FRNCLN	107.0	45.5	44.0	22.7
L60	LERKE/5/KAUZ/3/MYNA/VUL//BUC/FLK	Z/4/MIL9 9N 96/	51.3	48.8	35.9
L61	PASTOR/3/VORONA/CN079//KAUZ/4/MII	LAN/O T0985 /	40.0	42.0	35.1
L62	PASTOR/3/VORONA/CN079//KAUZ/4/MII	LAN/O Þ9:5 //	46.8	47.0	40.5
L63	PFAU/MILAN//TROST/3/PBW65/2*SERI.1	B*2/4/9.7.5	35.6	45.2	39.5
L64	PFAU/MILAN//TROST/3/PBW65/2*SERI.1	B*2/4/104.0	43.2	51.9	49.5
L65	DOY1/AE.SQUARROSA (447)/3/KA/NAC//TRCH/4/	115.5	43.1	36.4	27.0
L66	MON/IMU//ALD/PVN/3/BORL95/4/OASIS	/2*BORI395/	51.5	49.8	39.2
L67	WHEAR/SOKOLL/4/PASTOR//MILAN/KA	UZ/3/ B AM. 9 2	40.1	47.1	38.8
L68	SOKOLL/WBLL1/4/D67.2/PARANA 66.270//	98.5	46.8	41.6	28.4
L69	MILAN//PRL/2*PASTOR/4/CROC_1/AE.S0 (2	QUAR ®9.5 A	38.0	45.0	33.0
L7	MUU/FRNCLN	94.5	33.4	49.4	24.3
L70	TRCH/SRTU//KACHU/3/KINGBIRD #1	101.5	33.5	44.4	39.0
L71	TRCH/SRTU//KACHU/3/KINGBIRD #1	97.5	32.5	43.7	33.2
L72	TRCH/5/BAV92//IRENA/KAUZ/3/HUITES/	/4/DOLII04.0	36.7	44.4	39.1
L73	BECARD #1/3/PBW343*2/KUKUNA//PBW343*2/KU	102.5 JKUNA	47.2	46.5	36.3
L74	DANPHE/3/PBW343*2/KUKUNA//PBW34	3*2/K UKUN A	30.5	50.2	37.1
L76	PBW343*2/KUKUNA//PBW343*2/KUKUN	NA/6/WBLD1*2/	32.2	47.9	40.3
L77	BECARD/6/FRET2*2/4/SNI/TRAP#1/3/KA	UZ*2/ T05A0 P/	55.9	47.6	47.2
L78	KACHU/6/YAR/AE.SQUARROSA (783)/4/GOV/AZ//	94.5	37.8	39.8	29.8
L79	FRANCOLIN #1/NELOKI	101.0	48.7	45.7	40.4

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits *(continued)*

encoding	Genotype	Plant height	Leaf area	SPADI	SPADII
L8	PRL/ 2*PAS-	98.5	32.9	39.6	29.7
	TOR/4/CHOIX/STAR/3/HE1/3*CNO79//				
L80	PAURAQ/8/NG8201/KAUZ/4/SHA7//PRL/V	EE#6/B0/0.5	46.2	45.5	36.2
L81	PRL/2*PASTOR//SUNSTATE/3/GRACK	104.0	39.0	38.7	34.9
L82	PICAFLOR #1/NELOKI	111.5	43.5	43.6	44.2
L83	PICAFLOR	91.5	25.6	42.7	19.3
	#1/8/NG8201/KAUZ/4/SHA7//PRL/VEE#6/				
L84	NL971*2/MUU	104.5	39.8	44.9	42.0
L85	MUNAL	98.5	28.2	49.2	35.3
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2	2/			
L86	MUNAL	92.5	27.6	41.4	25.3
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2	2/			
L87	MUNAL	97.0	26.0	50.7	35.1
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2	2/			
L88	MUNAL	100.5	33.8	50.0	38.1
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2	2/			
L89	BECARD #1*2/KINGBIRD #1	98.0	35.0	40.8	29.5
L9	DANPHE #1*2/SHORTENED SR26	99.5	31.6	42.2	25.8
	TRANSLOCATION				
L90	BECARD*2/PFUNYE #1	105.0	44.7	48.1	43.2
L91	PBW343/PASTOR//OTUS/TOBA97*2/3/PIC	AFLO98.0	28.5	34.5	32.8
	#1				
L92	ROLF07/KINGBIRD #1//MUNAL #1	93.0	46.4	50.7	31.2
L93	MUNAL	95.0	32.9	42.0	31.4
	#1/3/PBW343*2/KUKUNA*2//YANAC				
L94	SERI.1B*2/3/KAUZ*2/BOW/KAUZ/4/PBW3	343*2/99.5	39.3	38.0	37.4
L95	KAUZ//ALTAR	99.0	39.5	48.0	37.6
	84/AOS/3/MILAN/KAUZ/4/SAUAL/5/				
L96	PBW343*2/KHVAKI//PARUS/3/PBW343/PA	STOR95/0.	34.0	52.0	29.0
L97	FRET2*2/4/SNI/TRAP#1/3/KAUZ*2-	107.0	42.5	39.0	27.0
	TRAP//KAUZ*2/				
L98	ATTILA*2/PBW65*2//JUCHI/3/KINGBIRD	98.0	38.6	49.4	43.8
	#1/4/	02.5	10.7		
L99	MUNAL #1*2//WBLL1*2/BRAMBLING	93.5	18.7	47.5	41.1

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits (...continued)

encoding	Genotype	CTD	Days to heading	Days to anthesis
1	Aditya	8.82	67.7	76.2
2	Bhrikuti	9.63	64.6	73.6
3	Gautam	9.39	67.7	74.9
4	Tilottama	9.52	62.7	71.7
L1	MUNAL #1	11.28	74.0	85.0
L10	MUNAL/3/HUW234+LR34/PRINIA//PFAU/WI	EAVER //0 //7.1	74.0	84.0
L101	PFAU/MILAN/5/CHEN/AEGILOPS SQUARROSA (TAUS)/	10.59	73.0	80.0
L102	PREMIO/5/TUI//2*SUNCO/SA1166/3/TUI/4/FI	NSI/11.77	74.0	82.0
L103	PREMIO/BAVIS	11.69	75.0	84.0
L104	CHIYAK//ND643/2*WAXWING/3/ND643/2*W	AXWI NG 3	73.0	80.0
L105	BABAX/LR42//BABAX*2/4/SNI/TRAP#1/3/KA	AUZ*21.3.68	74.0	80.0
L106	PFAU/MILAN//TROST/3/MUNAL #1/4/PFAU/MILAN//	2.60	74.0	82.0
L107	YUK/AE.SQUARROSA (217)//2*PANDORA	8.54	68.0	78.0
L108	SUP152*2//ND643/2*WAXWING	10.37	71.0	78.0
L11	SWSR22T.B./5/KAUZ//ALTAR 84/AOS/3/KAUZ/4/	10.77	75.0	86.0
L12	WHEAR/VIVITSI//WHEAR/3/PANDORA	11.30	74.0	84.0
L13	WBLL1*2/KURUKU//TACUPETO F2001*2/BRAMBLING	10.69	74.0	82.0
L14	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN	/OTUS/128	74.0	84.0
L15	FRANCOLIN #1*2/ND643/2*WBLL1	9.03	73.0	81.0
L16	KIRITATI/ 2*WBLL1/3/TAM200/PASTOR//TOBA97/4/	9.83	73.0	80.0
L17	SUP152/KENYA SUNBIRD	10.15	71.0	78.0
L18	SUP152/KENYA SUNBIRD	11.34	63.0	73.0
L19	SUP152/KENYA SUNBIRD	11.72	63.0	73.0
L2	KACHU #1	9.62	71.0	80.0
L20	WBLL1*2/BRAMBLING/5/BABAX/LR42//BA	BAX*2 7 495.	70.0	78.0
L21	ND643/2*WBLL1/3/KIRITATI//2*PRL/2*PAST	OR/4/10.75	74.0	80.0
L22	BETTY/3/CHEN/AE.SQ//2*OPATA/4/BAVIS	9.50	70.0	78.0
L23	BAVIS/VORB/5/CROC_1/AE.SQUARROSA(20	5)// 12.89	65.0	73.0
L24	BABAX/LR42//BABAX/3/ER2000/4/NAVJ07	4.80	68.0	76.0

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits (...continued) (continued)

encoding	Genotype	CTD	Days to heading	Days to anthesis
L26	NING-MAI-	9.97	68.0	78.0
	96035/FINSI//HEILO/3/NAVJ07			
L27	WBLL1*2/BRAMBLING//TAM200/TUI/3/	11.53	75.0	86.0
L28	KISKADEE	12.74	74.0	80.0
	#1/5/KAUZ*2/MNV//KAUZ/3/MILAN/4/			
L29	MERCATO//PARUS/PASTOR	11.19	74.0	85.0
L3	KACHU/BECARD//WBLL1*2/BRAMBLING	13.35	74.0	81.0
L30	PREMIO//PARUS/PASTOR	11.68	75.0	85.0
L31	ND643/2*WBLL1//ND643/2*WAXWING	9.71	71.0	81.0
L32	ND643/2*WBLL1//ND643/2*WAXWING	10.74	73.0	80.0
L33	ND643/2*WBLL1//ND643/2*WAXWING	5.30	72.0	80.0
L34	SNB//CMH79A.955/3*CN079/3/ATTILA/4/CHEN/	8.62	75.0	85.0
L35	KIRITATI/ 2*WBLL1/8/SHA7/	10.03	73.0	78.0
	PRL/VEE#6/ 3/ FASAN/			
L36	SERI.1B*2/3/KAUZ*2/BOW//KUZ/4/CIRCUS/5/	11.72	72.0	78.0
L37	WHEAR/KUKUNA/3/C80.1/3*BATAVIA//2*WBLL1	/40.25	75.0	85.0
L38	BAJ #1*2/BECARD	10.52	70.0	78.0
L39	KACHU*2/MUNAL #1	10.05	75.0	85.0
L4	PRL/2*PASTOR/3/PFAU/WEAVER*2//CHAPIO	10.75	74.0	85.0
L40	KACHU*2/MUNAL #1	12.14	75.0	84.0
L41	KACHU/2*MUNAL #1	10.45	65.0	73.0
L42	KACHU/2*MUNAL #1	5.30	68.0	74.0
L43	KACHU*2/PANDORA	6.42	72.0	78.0
L44	SUP152/2*MUNAL #1	10.86	64.0	73.0
L45	FRNCLN*2/BECARD	10.65	75.0	84.0
L46	WBLL1*	11.27	75.0	86.0
	2/BRAMBLING//SAAR/2*WAXWING/4/			
L47	CHIBIA//PRLII/CM65531/3/MISR 2*2/4/	8.90	73.0	82.0
L48	CHIBIA//PRLII/CM65531/3/MISR 2*2/4/	11.64	73.0	84.0
L49	WAXWING/KRONSTAD	10.96	73.0	82.0
	F2004//2*FRNCLN			
L5	DANPHE/CHONTE	11.05	75.0	85.0
L51	MUNAL	12.11	73.0	78.0
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/			
L52	MUNAL	11.85	73.0	80.0
	#1/5/2*PRL/2*PASTOR/4/CHOIX/STAR/3/			

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits (...continued) (continued)

encoding	Genotype	CTD	Days to heading	Days to anthesis
L53	MUNAL #1/2*FRNCLN	10.22	70.0	78.0
L54	MUNAL #1/2*FRNCLN	9.53	75.0	85.0
L55	MUNAL #1*2//WHEAR/SOKOLL	9.82	74.0	84.0
L56	MUNAL #1*2/KINGBIRD #1	10.65	73.0	80.0
L57	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	6.60	71.0	78.0
L58	PRL/2*PASTOR//SUNSTATE/3/MUNAL #1/4/OTUS//	4.72	72.0	78.0
L59	KAUZ*2/BOW//KAUZ/3/W98.6.38/5/SABUF/4/	10.70	73.0	80.0
L6	SUP152/FRNCLN	9.38	70.0	78.0
L60	LERKE/5/KAUZ/3/MYNA/VUL//BUC/FLK/4/MIL/	N869.7.	75.0	85.0
L61	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN/OT	U S%.63	74.0	80.0
L62	PASTOR/3/VORONA/CN079//KAUZ/4/MILAN/OT	U SI/18 8	74.0	84.0
L63	PFAU/MILAN//TROST/3/PBW65/2*SERI.1B*2/4/	7.30	74.0	81.0
L64	PFAU/MILAN//TROST/3/PBW65/2*SERI.1B*2/4/	9.88	75.0	86.0
L65	DOY1/AE.SQUARROSA (447)/3/KA/NAC//TRCH/4/	9.00	71.0	80.0
L66	MON/IMU//ALD/PVN/3/BORL95/4/OASIS/2*BOR	L9151/.9.9	74.0	82.0
L67	WHEAR/SOKOLL/4/PASTOR//MILAN/KAUZ/3/BA	AV91272	74.0	85.0
L68	SOKOLL/WBLL1/4/D67.2/PARANA 66.270//	9.04	73.0	80.0
L69	MILAN//PRL/2*PASTOR/4/CROC_1/AE.SQUARRO (2	OSIA.34	75.0	85.0
L7	MUU/FRNCLN	10.11	72.0	78.0
L70	TRCH/SRTU//KACHU/3/KINGBIRD #1	12.15	75.0	86.0
L71	TRCH/SRTU//KACHU/3/KINGBIRD #1	9.81	75.0	84.0
L72	TRCH/5/BAV92//IRENA/KAUZ/3/HUITES/4/DOLL	8.93	75.0	85.0
L73	BECARD #1/3/PBW343*2/KUKUNA//PBW343*2/KUKUNA	8.71	73.0	82.0
L74	DANPHE/3/PBW343*2/KUKUNA//PBW343*2/KU	KUNA8	75.0	85.0
L76	PBW343*2/KUKUNA//PBW343*2/KUKUNA/6/WF	BLL11.1521/	75.0	82.0
L77	BECARD/6/FRET2*2/4/SNI/TRAP#1/3/KAUZ*2/TI	RAP/.53	73.0	84.0
L78	KACHU/6/YAR/AE.SQUARROSA (783)/4/GOV/AZ//	9.69	67.0	77.0
L79	FRANCOLIN #1/NELOKI	5.78	70.0	78.0

Table 4.16: Model adjusted genotypic mean responses of plant's morphological, architectural and phenological traits (...continued) (continued)

encoding	Genotype	CTD	Days to heading	Days to anthesis
L8	PRL/ 2*PAS-	9.72	75.0	86.0
	TOR/4/CHOIX/STAR/3/HE1/3*CNO79//			
L80	PAURAQ/8/NG8201/KAUZ/4/SHA7//PRL/VEE#6/3/		75.0	85.0
L81	PRL/2*PASTOR//SUNSTATE/3/GRACK	8.81	75.0	86.0
L82	PICAFLOR #1/NELOKI	12.38	75.0	86.0
L83	PICAFLOR	9.82	69.0	78.0
	#1/8/NG8201/KAUZ/4/SHA7//PRL/VEE#6/			
L84	NL971*2/MUU	11.45	74.0	84.0
L85	MUNAL	9.76	69.0	78.0
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/			
L86	MUNAL	9.93	67.0	78.0
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/			
L87	MUNAL	11.75	73.0	78.0
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/			
L88	MUNAL	10.22	72.0	80.0
	#1*2/4/HUW234+LR34/PRINIA//PBW343*2/			
L89	BECARD #1*2/KINGBIRD #1	4.55	72.0	78.0
L9	DANPHE #1*2/SHORTENED SR26	10.55	73.0	80.0
	TRANSLOCATION			
L90	BECARD*2/PFUNYE #1	11.32	73.0	82.0
L91	PBW343/PASTOR//OTUS/TOBA97*2/3/PICAFLOR	11.70	75.0	86.0
	#1			
L92	ROLF07/KINGBIRD #1//MUNAL #1	6.34	74.0	80.0
L93	MUNAL	9.73	72.0	80.0
	#1/3/PBW343*2/KUKUNA*2//YANAC			
L94	SERI.1B*2/3/KAUZ*2/BOW/KAUZ/4/PBW343*2/	9.47	74.0	84.0
L95	KAUZ//ALTAR	9.62	70.0	78.0
	84/AOS/3/MILAN/KAUZ/4/SAUAL/5/			
L96	PBW343*2/KHVAKI//PARUS/3/PBW343/PASTOR/5	/.8.88	74.0	84.0
L97	FRET2*2/4/SNI/TRAP#1/3/KAUZ*2-	10.18	74.0	82.0
	TRAP//KAUZ*2/			
L98	ATTILA*2/PBW65*2//JUCHI/3/KINGBIRD	10.50	75.0	86.0
	#1/4/			
L99	MUNAL #1*2//WBLL1*2/BRAMBLING	11.07	75.0	86.0

4.3.2 Mixed model summary of fixed effects terms of plant morphological, architectural and phenological traits

	Dependent variable:							
	Plant height	linear mixed-effects Leaf area SPAD I		SPAD II	Canopy temperature depression			
	(1)	(2)	(3)	(4)	(CTD)			
Bhrikuti	-9.46*** (1.17)	1.51 (1.79)	-1.74*** (0.60)	-3.78** (1.61)	0.52** (0.24)			
Gautam	p = 0.00 $1.21 (1.17)$	$p = 0.40$ $3.28^* (1.81)$	$p = 0.004 \\ -1.06^* (0.61)$	$p = 0.02 \\ -0.38 (1.62)$	$p = 0.04 \\ 0.20 (0.24)$			
Tilottama	$p = 0.31$ $-4.92^{***} (1.17)$	p = 0.07 -1.49 (1.80)	$p = 0.08 \\ -1.38^{**} (0.61)$	$p = 0.82 -6.46^{***} (1.62)$	$p = 0.42 \\ 0.43^* (0.24)$			
Aditaya (Constant)	$p = 0.0001$ $104.00^{***} (2.23)$ $p = 0.00$	$p = 0.41$ 37.60^{***} (2.39) $p = 0.00$	$p = 0.03$ $45.90^{***} (2.73)$ $p = 0.00$	$p = 0.0001$ $38.80^{***} (1.37)$ $p = 0.00$	p = 0.08 $9.07^{***} (0.79)$ p = 0.00			
Observations	238	238	238	238	238			
Log Likelihood	-730.00	-821.00	-600.00	-782.00	-378.00			
Akaike Inf. Crit. Bayesian Inf. Crit.	1,484.00 1,525.00	1,666.00 1,707.00	1,224.00 1,265.00	1,587.00 1,629.00	780.00 821.00			

*p<0.1; **p<0.05; ***p<0.01

Dependent variable:			
linear			
mixed-effects			
Days to heading	Days to anthesis		
(1)	(2)		

Bhrikuti	-3.22*** (0.37)	-2.75*** (0.45)
	p = 0.00	p = 0.00
Gautam	-0.19 (0.38) p = 0.62	$-1.59^{***} (0.46)$ p = 0.001
Tilottama	-5.07^{***} (0.38)	$-4.65^{***} (0.46)$
	p = 0.00	p = 0.00
Aditaya (Constant)	67.80*** (2.40)	76.30*** (3.10)
	p = 0.00	p = 0.00
Observations	238	238
Log Likelihood	-509.00	-560.00
Akaike Inf. Crit.	1,042.00	1,145.00
Bayesian Inf. Crit.	1,084.00	1,186.00
Nota:	*n <0.1. **n	<0.05: ***n<0.01

Note: *p<0.1; **p<0.05; ***p<0.01

Plant height

Highly significant difference exist among the genotypes for Plant height trait based on the significane of LRT using χ^2 statistic (59.5 with 4 df).

A pairwise contrast of fixed effects estimate shows that all check varieties differ from one another except for Aditya (mean: $103.00~cm~(\pm~1.4)$) and Gautam (which are similar at 0.05 level of significance), implying that plant height is an important trait to differentiate the genotypes (indicated by a p-value of less than 0.01). With an average height of $94.53~cm~(\pm~1.4)$, Bhrikuti is the shortest variety. Likewise, the tallest variety (Gautam) has the mean height of $102.86~cm~(\pm~1.41)$.

Meanwhile, random effects estimates for entry genotype levels do not show as much of difference. Also, the non-significance of LRT comparing models with and without random terms for entry genotypes clearly indicates that no relationship exists between plant height and entry genotypes. ANOVA table (A.12) and the dotplot (Figure 4.10a) showing random effects estimates of the entry genotypes illustrate the relationship.

However, blocking factors – row nested within rowgroup and superimposed blocks arising from the intersection between rowgroup and colgroup – have contributed significantly to the variation in plant height.

Table 4.19: Mean differences in Plant height of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	9.46	1.17	106	8.10	7.14	11.77	0.000
Aditya-Gautam	-1.21	1.18	106	-1.03	-3.54	1.12	0.305
Aditya-Tilottama	4.92	1.17	105	4.19	2.59	7.24	0.000
Bhrikuti-Gautam	-10.67	1.17	106	-9.08	-13.00	-8.34	0.000
Bhrikuti-Tilottama	-4.54	1.17	105	-3.87	-6.87	-2.22	0.000
Gautam-Tilottama	6.13	1.18	106	5.18	3.78	8.47	0.000

Flag leaf surface area

No significant difference exist among the genotypes for Flag leaf surface area trait based on the non significance of LRT using χ^2 statistic (ANOVA Table A.13).

A pairwise contrast of fixed effects estimate however reveals that check varieties Gautam (mean:

 $41.0~cm^2~(\pm~1.81)$) and Tilottama (mean: $36.0~cm^2~(\pm~1.80)$) had different flag leaf surface area, indicated by a p-value of less than 0.05.

However, effect of blocking factor – row nested within rowgroup – can be confirmed as significant source of environmental variability in the flag leaf area trait (based on LRT).

Table 4.20: Mean differences in Flag leaf area of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	-1.51	1.79	110	-0.84	-5.070	2.041	0.400
Aditya-Gautam	-3.28	1.81	110	-1.82	-6.864	0.299	0.072
Aditya-Tilottama	1.49	1.80	110	0.83	-2.087	5.066	0.411
Bhrikuti-Gautam	-1.77	1.80	109	-0.98	-5.340	1.803	0.329
Bhrikuti-Tilottama	3.00	1.80	109	1.67	-0.567	6.576	0.098
Gautam-Tilottama	4.77	1.82	110	2.63	1.172	8.374	0.010

SPAD at Zadok's stage 65

No significant difference exist among the genotypes SPAD measurement at Zadok's stage 65 based on the non significance of LRT using χ^2 statistic. However, the test statistic of 8.95 at 4 df indicates that the hypothesis of no-difference do not hold at 10% level of significance.

A pairwise contrast of fixed effects estimate, however, reveals that check variety Aditya has a higher mean SPAD value ($45.90~(\pm~2.73)$) than both check varieties Bhrikuti and Tilottama. The variety with the lowest relative chlorophyll content at the given stage is Bhrikuti ($44.20~(\pm~2.73)$).

Likewise, highly variable estimates of random effects (Shown in 4.10c) along with significance of LRT (Shown in A.14) indicates that the entry genotypes vary to a large extent in the leaf chlorophyll content, early in the reproductive phase of the crop.

Meanwhile, blocking factors do not have significant effects associated with the trait at the reference level of significance ($\alpha = 0.05$).

Table 4.21: Mean differences in SPADI (Zadok's stage 65) of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	1.739	0.603	107	2.88	0.544	2.935	0.005
Aditya-Gautam	1.064	0.608	105	1.75	-0.140	2.269	0.083
Aditya-Tilottama	1.380	0.607	104	2.27	0.176	2.584	0.025
Bhrikuti-Gautam	-0.675	0.607	106	-1.11	-1.878	0.528	0.268
Bhrikuti-Tilottama	-0.359	0.606	104	-0.59	-1.561	0.843	0.555
Gautam-Tilottama	0.316	0.611	105	0.52	-0.896	1.528	0.606

SPAD at Zadok's stage 85

LRT of mixed models using χ^2 statistic show that highly significant differences exist among the genotypes for the SPAD measured at Zadok's stage 85.

A pairwise contrast of fixed effects estimate, however, reveals that check variety Aditya $(38.80~(\pm~1.37))$ has the higher mean SPAD values in comparison to check varieties Bhrikuti and Tilottama, although no significant difference exist between the most chlorophyll preserved variety (Aditya) and the Gautam. Similarly, the mean comparison also shows that check variety Gautam also has higher relative chlorophyll content than other two check varieties. The lowest mean SPAD values was recorded for Tilottama $(32.35~(\pm~1.37))$ the given stage.

In contrast, the entry genotypes do not present as much a variation as check cultivars do among each other for SPAD measurement at latter stage of growth. This is evident from LRT test shown in A.15 and also random effects estimate of the entry genotypes (Shown in 4.10d). Similarly, blocking factors do not present an attributable source of variation for the trait.

Table 4.22: Mean differences in SPADII (Zadok's stage 85) of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	3.78	1.61	126	2.35	0.599	6.962	0.020
Aditya-Gautam	0.38	1.62	126	0.23	-2.826	3.586	0.815
Aditya-Tilottama	6.46	1.62	126	3.99	3.257	9.668	0.000
Bhrikuti-Gautam	-3.40	1.62	126	-2.10	-6.605	-0.196	0.038
Bhrikuti-Tilottama	2.68	1.62	126	1.66	-0.522	5.887	0.100
Gautam-Tilottama	6.08	1.63	126	3.73	2.854	9.311	0.000

Canopy temperature depression (CTD)

LRT of mixed models using χ^2 statistic show that genotypes have significant differences for the crop architecture trait Canopy temperature depression (CTD) measured near anthesis stage.

Pairwise comparison, using Tukey procedure, of fixed effects estimate shows that canopy structure of check variety Bhrikuti favors greater reduction in temperature than check variety Aditya $(9.08^{\circ}C~(\pm~0.79))$. Other varieties were at par with both of the check varieties. The

In contrast, the entry genotypes do not present attributable variation in CTD. This is evident from LRT test shown in A.16 and also the variance components decomposition of random effects terms, which shows no heritable variation is found among the tested entries.

Although, multiple blocking factors, comprising effects of row group, nested effects of columns within column group and the effects of superimposed blocks arising from the intersection of individual row group and column group are found significantly high for the variation in canopy temperature depression trait.

Estimate Upper CI Contrast Standard Error DF t-value Lower CI p-value -2.14 -0.998 -0.0410.034 -0.5190.243 206 Aditya-Bhrikuti -0.1990.244 206 -0.82-0.6810.283 0.416 Aditya-Gautam -0.433-1.77 -0.9150.078 Aditya-Tilottama 0.244 206 0.048 Bhrikuti-Gautam 0.320 0.245 207 1.31 -0.1630.803 0.193 Bhrikuti-Tilottama 0.086 0.244 206 0.35 -0.3960.568 0.725

206

-0.95

-0.720

0.252

0.343

Table 4.23: Mean differences in CTD of check varieties (post-hoc comparison using Tukey procedure)

Days to heading

Gautam-Tilottama

-0.234

Highly significant difference exists among the genotypes for days to heading trait based on the significane of LRT using χ^2 statistic (155 with 4 degrees of freedom).

0.246

Pairwise comparison of fixed effects estimate shows that all check varieties differ from one another except for Aditya (late heading type) (with mean of $67.79~(\pm~2.40)$ days) and Gautam, which require similar Days for heading. Mean number of days taken for head to develop in half of the population was lowest in Tilottama ($62.71~(\pm~2.40)$ days) variety.

Similarly, random effects estimates for entry genotype levels also exhibit high variation for the trait, thereby indicating that significant proportion of the variation is heritable. ANOVA table (A.17) and the dotplot (Figure 4.11b) showing random effects estimates of the entry genotypes illustrate the relationship. Amongst blocking factors, the effect of grouping the experimental units in rowgroups has relevance in reducing extraneous source of variability carried to the trait in the field.

Table 4.24: Mean differences in Days to heading of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	3.221	0.373	112	8.63	2.482	3.960	0.000
Aditya-Gautam	0.187	0.376	111	0.50	-0.558	0.932	0.621
Aditya-Tilottama	5.073	0.376	111	13.51	4.329	5.817	0.000
Bhrikuti-Gautam	-3.034	0.376	113	-8.08	-3.779	-2.290	0.000
Bhrikuti-Tilottama	1.852	0.375	112	4.93	1.108	2.596	0.000
Gautam-Tilottama	4.887	0.379	112	12.91	4.137	5.637	0.000

Days to anthesis

Highly significant difference exists among the genotypes for Number of days to anthesis trait based on the significane of LRT using χ^2 statistic (85.8 with 4 df).

Pairwise comparison of fixed effects estimate shows that all check varieties differ from one another, with longest period to anthesis being that of check variety Aditya (76.30 (\pm 3.10) days) and the earliest flowering cultivar being Tilottama with an average of 71.7 (\pm 0.46) days.

Similarly, entry genotype also have a considerable genotypic variation which is reflected in a large measure of variance in random effects term. ANOVA table (A.18) and the dotplot (Figure 4.11c) showing random effects estimates of the entry genotypes illustrate the relationship. Likewise, the effects of rowgroup and colgroup, contributing significantly to the variation in trait, may be safely substracted for obtaining better effects estimates of the entry genotypes.

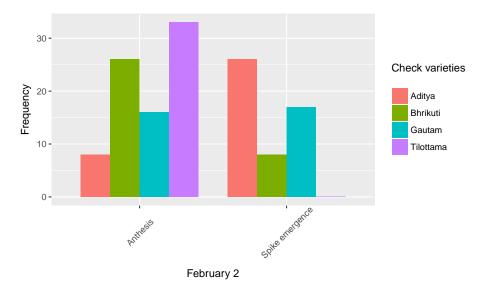


Figure 4.7: Number of checks at different growth stages at February 2

Frequency barplot of different stages of reproductive periods of growth in wheat (Figure 4.7) reaffirms that Tilottama variety flowers the earliest amongst the check varieties and Aditya reaches to the flowering relatively late.

Table 4.25: Mean differences in Days to anthesis of check varieties (post-hoc comparison using Tukey procedure)

Contrast	Estimate	Standard Error	DF	t-value	Lower CI	Upper CI	p-value
Aditya-Bhrikuti	2.75	0.453	105	6.07	1.854	3.65	0.000
Aditya-Gautam	1.59	0.457	105	3.48	0.683	2.50	0.001
Aditya-Tilottama	4.65	0.456	104	10.19	3.747	5.56	0.000
Bhrikuti-Gautam	-1.16	0.456	104	-2.55	-2.067	-0.26	0.012
Bhrikuti-Tilottama	1.90	0.456	104	4.17	0.996	2.80	0.000
Gautam-Tilottama	3.06	0.460	104	6.66	2.151	3.97	0.000

4.3.3 Mean comparison of plant's morphological, architectural and phenological traits of entry genotypes

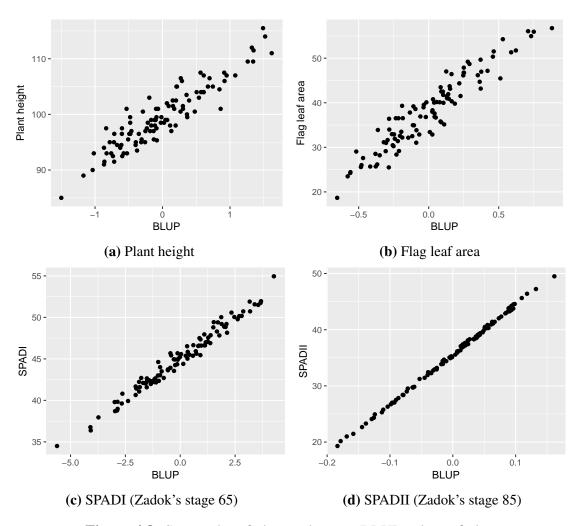


Figure 4.8: Scatterplot of observed versus BLUP values of plant morphological, architectural and phenological traits

Scatterplot showing observed means of plant's morphological, phenological and architectural traits plotted against their respective genotypic BLUP values is presented in Figure 4.8.

The coefficient of determination, as computed by squared correlation coefficient between the observed and fitted values of random effects model are 0.56, 0.33, 0.7, 0.1, 0.81, 0.92, 0.91, respectively for the Plant height, Flag leaf area, SPAD at Zadok's stage 65, SPAD at Zadok's stage 85, Canopy temperature depression, Days to heading and Days to anthesis.

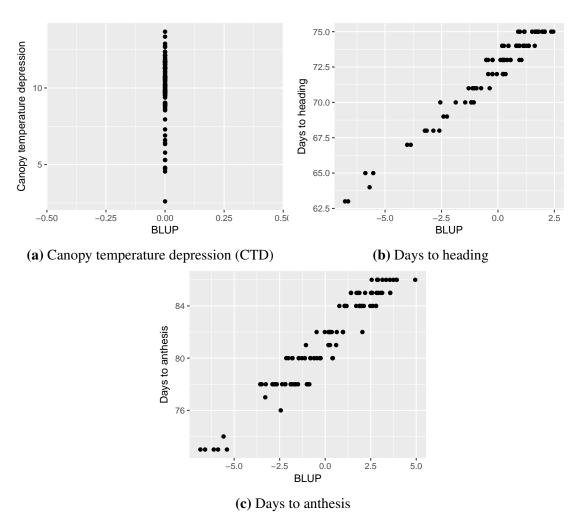


Figure 4.9: Scatterplot of observed versus BLUP values of plant's morphological, architectural and phenological traits (...continued)

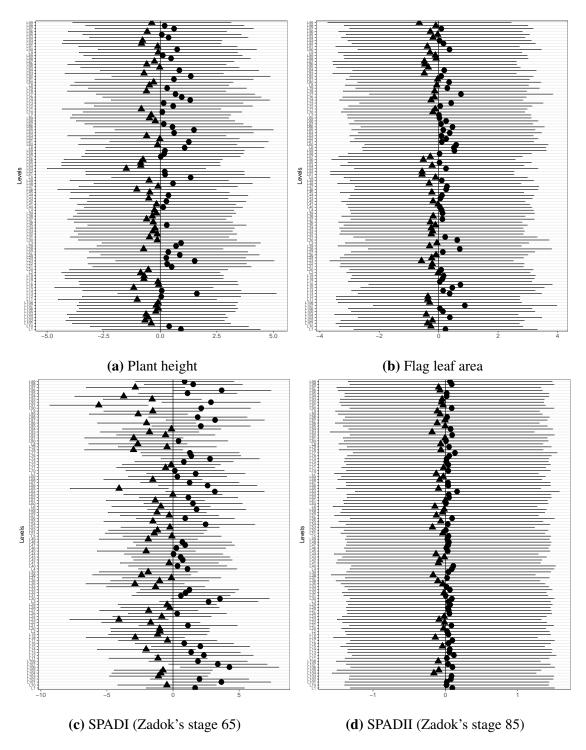
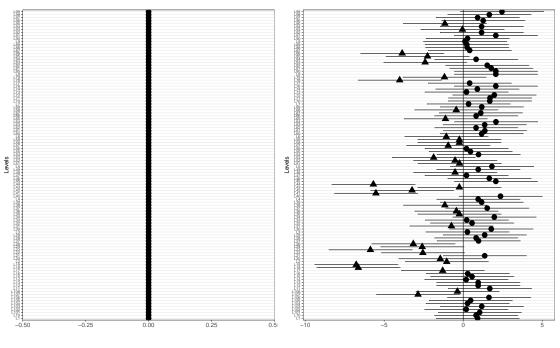
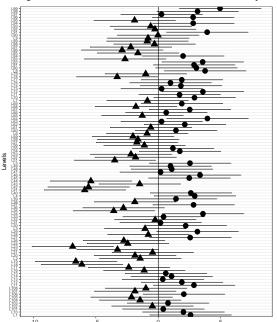


Figure 4.10: Random effect estimates of plant's morphological, architectural and phenological traits of entry genotypes



(a) Canopy temperature depression (CTD)

(b) Days to heading



(c) Days to anthesis

Figure 4.11: Random effect estimates of plant's morphological, architectural and phenological traits of entry genotypes (...continued)

4.4 Yield prediction using leaf health

A model approximating yield as a linear function of fixed effect terms for leaf health traits and random terms for blocking structure was formulated and ran. Along with model summary, effects estimates were obtained and coefficient of determination calculated based on the likelihood-ratio test (improvement from null (intercept only) model to the fitted model) (Magee, 1990).

4.4.1 Stepwise variable selection

For determining influencial variables among the candidates, stepwise variable selection was performed from a set of possible models. Table 4.26 and Table 4.27 summarize which fixed and random effects variables, respectively, had or did not have significant influence in yield determination.

Table 4.26: ANOVA of fixed effect model terms with indication of significance of variables

Trait variables	Sum Sq	Mean Sq	NumDF	DenDF	F.value	elim.num	Pr(>F)
SPADII	0.016	0.016	1	170.3	0.065	1	0.798
Disease Z75	0.120	0.120	1	188.8	0.484	2	0.487
Greenness Z85	0.220	0.220	1	190.0	0.890	3	0.347
Check genotypes	4.416	1.104	4	27.9	4.471	kept	0.006
Greenness Z65	4.385	4.385	1	170.4	17.755	kept	0.000
Greenness Z75	5.322	5.322	1	190.5	21.552	kept	0.000
LAUG 65_75	18.945	18.945	1	188.1	76.717	kept	0.000
SPADI	3.134	3.134	1	187.2	12.689	kept	0.000

Table 4.27: ANOVA of random effect model terms with indication of significance of variables

Random variables	Chi.sq	Chi.DF	elim.num	p.value
Nested col	0.000	1	1	1.000
Rowgroup x Colgroup	0.000	1	2	1.000
Nested row	0.061	1	3	0.804
Rowgroup	1.556	1	4	0.212
Colgroup	8.560	1	kept	0.003
Line genotypes	5.116	1	kept	0.024

Stepwise variable selection and backward elimination of a mixed model with response variable yield suggests that, aside from the genotype effects, fixed effects of leaf health and canopy related traits significantly affect the yield (based on both, P-values calculated from F-test by Sattethwaite's

approximation and AIC (Mazerolle, 2004; Wagenmakers & Farrell, 2004) drop in the single term deletions of fixed effects terms in the model). In particular leaf greenness scores at Zadok's stage 65 and 75, LAUG value during Zadok's stage 65 and 75 and relative leaf chlorophyll content (SPADI) at Zadok's stage 65.

4.4.2 Efficiency of yield prediction

The coefficient of determination of full model (squared correlation coefficient between observed and fitted values) was 0.794. Similarly, accounting for random effects, including all blocking and treatment structures, R^2 value of fixed effects terms (i.e. proportion of variance explained by leaf health traits) is 0.393.

Using relative efficiency measure: indicates the decrease in error variance with additional terms in alternative model (with leaf health traits) over null the model (with random terms only), approximate gain in efficiency with the use of leaf health in estimating yield was almost 51.812%.

Top ten best performing entry genotypes based on fitted yield (BLUPs) ranking with their yield estimates are shown in Table 4.28.

Table 4.28: Top ten best performing genotypes with respect to estimated yields

Entry genotype	Estimated yield	Genotype
L71	4.50	TRCH/SRTU//KACHU/3/KINGBIRD #1
L67	4.38	WHEAR/SOKOLL/4/PASTOR//MILAN/KAUZ/3/BAV92
L85	4.28	MUNAL #1*2/4/HUW234+LR34/PRINIA//PBW343*2/
L106	4.22	PFAU/MILAN//TROST/3/MUNAL #1/4/PFAU/MILAN//
L92	4.08	ROLF07/KINGBIRD #1//MUNAL #1
L87	3.98	MUNAL #1*2/4/HUW234+LR34/PRINIA//PBW343*2/
L43	3.95	KACHU*2/PANDORA
L1	3.94	MUNAL #1
L84	3.93	NL971*2/MUU
L105	3.89	BABAX/LR42//BABAX*2/4/SNI/TRAP#1/3/KAUZ*2/

4.5 Correlation between numeric variables

Table 4.29: Correlation coefficients of numeric response variables observed in study

	Yild	Paen	ThWt	Gran	Efer	Di75	Gr65	Gr75	Gr85	La75	La85	Plht	Laar	Cttd	Spdi	Spii	Daad
Yild																	
Paen	0.13																
ThWt	0.18*	-0.12															
Gran	0.32****	0.41****	-0.06														
Efer	0.56****	-0.06	-0.01	-0.03													
Di75	-0.07	-0.17*	0.03	-0.05	-0.06												
Gr65	-0.10	-0.09	0.06	-0.01	0.10	-0.14*											
Gr75	0.10	0.20**	-0.12	0.18**	-0.19**	-0.18**	-0.07										
Gr85	0.06	0.12	-0.04	0.07	-0.10	-0.15*	0.37****	0.77****									
La75	0.34****	0.19**	-0.10	0.20**	0.07	-0.16*	0.17**	0.82****	0.75****								
La85	0.08	0.12	-0.01	0.02	-0.16*	-0.14*	0.13	0.87****	0.85****	0.76****							
Plht	-0.10	0.09	-0.12	0.05	-0.11	-0.09	0.00	0.11	0.12	0.07	0.06						
Laar	-0.04	0.18**	0.05	0.10	-0.02	-0.08	-0.11	0.10	0.03	0.08	0.07	0.11					
Cttd	0.08	-0.19**	0.07	-0.05	0.13*	-0.10	0.00	-0.20**	-0.14*	-0.17**	-0.15*	0.03	-0.02				
Spdi	0.22**	0.11	0.13	0.16*	-0.03	-0.22**	0.10	0.18**	0.18**	0.19**	0.19**	-0.11	0.04	-0.08			
Spii	0.05	0.26****	-0.02	0.14*	0.02	-0.41****	0.05	0.17**	0.14*	0.14*	0.15*	0.23**	0.11	0.08	0.26****		
Daad	-0.13	-0.07	0.25**	-0.12	0.04	-0.06	0.30****	-0.45****	-0.17**	-0.29****	-0.31****	0.11	0.07	0.18**	0.13*	0.24**	
Dath	-0.13	-0.11	0.23**	-0.18**	0.07	-0.10	0.29****	-0.45****	-0.17**	-0.29****	-0.29****	0.08	0.08	0.24**	0.09	0.26****	0.93****

Note:

p < .0001: ****; p < .001: ***; p < .01: **; p < .05: *

Yild: Yield, Paen: Panicle length,

ThWt: Thousand kernel weight, Gran: Number of grains per panicle,

Efer: Number of effective tillers, Di75: Disease score at Z75,

Gr65: Greenness score at Z65, Gr75: Greenness score at Z75,

Gr85: Greenness score at Z85, La75: LAUG score at Z75,

La85: LAUG score at Z85, Plht: Plant height,

Laar: Leaf area, Cttd: Canopy temperature depression,

Spdi: SPAD score at Z65, Spii: SPAD score at Z85,

Daad: Days to heading

Pairwise test of pearson's correlation coefficients suggest a significant positive association of LAUG value during Zadok's stage 65 and 75 and the yield. Besides that, relative leaf chlorophyll content of the leaves (SPAD) recorded at Zadok's stage 65 are also significantly positively correlated with the yield. To the contrary, LAUG during Zadok's stage 65 was, with high degree of significant positive correlation, associated with the yield. This potentially implies that, a sharp drop in leaf greenness values right after flowering might be beneficial for yield formation, particularly to the number of grains per panicles.

No direct correlation to yield was apparent for other leaf health and traits characterizing canopy architecture. Major yield component traits associated with the yield were (in the decreasing order of significance) Number of effective tillers, Number of grains per panicle and the Thousand kernel weight. In addition, a higher relative chlorophyll content and a larger leaf surface area are implicated in longer panicles, although panicle length on itself is not significantly associated with the yield of the crop.

Several leaf health and canopy related traits had significant correlation with each other, for example, number of heading and flowering days for genotypes showed negative correlation with the decline in leaf greenness values, at both the early and latter stages of the life cycle. This potentially implies that, genotypes which flower early show higher drop in chlorophyll contents and reach to senescence earlier.

Likewise, certain phenological traits of interest in most selection programmes like days to heading and days to anthesis were significantly positively correlated with a particular yield component trait – thousand kernel weight, and the latter is even negatively correlated to the number of grains per panicle.

4.6 Principal component analysis

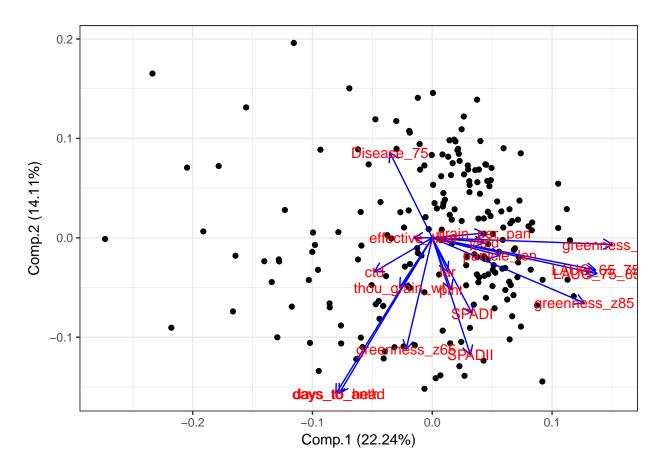


Figure 4.12: PCA biplot showing first two principal components

The principal component analysis of continuous multivariate response shows that first two principal components explain 35.17% variation. This is depicted in the Figure 4.12. Table 4.30 shows summary of principal component analysis (Top ten principal components are shown with corresponding measures of proportionate variance and standard deviation of principal axes).

For the first principal component axis (PC1), most variables have positive values (All except plant height, canopy temperature depression, disease score and the number of effective tillers triats). Thus, most variables are positively correlated between each other. Most leaf health related and phenological traits, including greenness scores of two latter stages, LAUG values and the days to heading and anthesis traits, constrain the PC1 eigenvector (based on a large magnitude of variable eigenvalue loadings) (Table 4.31). Interestingly, the variability accounted for by this PC axis is different from that of the PC2 and PC3 (latter captures 10.3% variation). Traits related to yield have higher factor

variable loadings in PC3, amongst which yield *per se* has the highest, while the number of effective tillers and the number of grains per panicle also contrain the third eigenvector. The scoring for disease contrasts itself in PC2. Also, all yield related traits can be seen as being positively correlated in the PC3 axis. Similarly, the indication of negative correlation between the days to heading and anthesis and the leaf greenness related traits corroborates the earlier speculation that genotypes that flower early are also characterized by earlier signs of drop in chlorophyll contents, and therefore reach to the senescence earlier.

Table 4.30: Summary of principal component analysis with measures of variation explained by top ten principal axes

	PC1	PC2	PC3	PC4	PC5	PC6	PC7	PC8	PC9	PC10
Standard deviation	2.00	1.59	1.36	1.30	1.12	1.03	1.02	0.92	0.87	0.82
Proportion of Variance	0.22	0.14	0.10	0.09	0.07	0.06	0.06	0.05	0.04	0.04
Cumulative Proportion	0.22	0.36	0.47	0.56	0.63	0.69	0.75	0.80	0.84	0.87

Table 4.31: PCA factor loadings of top three principal components on variables for each of the numeric response variables observed in study

Variable	PC1	PC2	PC3
Yield	-0.135	0.014	0.630
Panicle Len	-0.176	0.055	0.154
Grain Per Pan	-0.134	-0.015	0.333
Thou Grain Wt	0.085	0.155	0.134
Effective Tiller	0.048	0.000	0.512
Greenness Z65	0.068	0.349	-0.091
Greenness Z75	-0.468	0.021	-0.114
Greenness Z85	-0.395	0.204	-0.163
Laug 65 75	-0.424	0.100	0.091
Laug 75 85	-0.428	0.112	-0.161
Disease 75	0.110	-0.267	-0.065
Spadi	-0.104	0.237	0.206
Spadii	-0.099	0.366	0.068
Ctd	0.150	0.106	0.148
Plht	-0.049	0.162	-0.176
Lar	-0.040	0.105	-0.010
Days To Head	0.242	0.487	-0.028
Days To Anth	0.250	0.488	-0.042

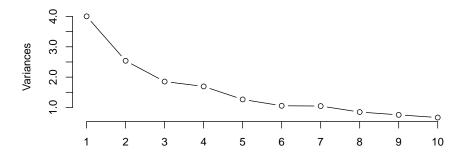


Figure 4.13: Principal component analysis screeplot of plants' trait responses

The PCA screeplot (Figure 4.13) shows the decay of variability (y-axis) with increasing numbers of independent principal components (x-axis) used for decomposing observed variability in the data. A sharp drop in proportion of total variance explained by the PCs is only realized after linear combination of 6-7 PCs. This indicates that data inherently is best resolved into large number of dimensions. This seems to be true and features unique opportunity as well as challenge from the perspective of modeling and analysing of a field experiment while controlling for several extrinsic factors that might otherwise result in the problem of sparsity of data.

Chapter 5

Discussion

5.1 Relationship between leaf health and yield

Four leaf health traits were identified with significant association to yield. Inspection of individual variance components reveals that for yield and yield component traits, blocking of plots into either factors, i.e row, column or superimposed row-colum, generally did not significantly reduce the error variance. However, thousand kernel weight was affected also by the variation along the column nested inside colum-group (increasing of effect along column 1 to 12). The finding is in accordance to the study of Neupane Adhikari, Wu, & Caffe (2016), where significant estimates have been recovered by spatial modeling.

Also, the effect on disease rating of foliar disease complex due to blocking in nested column inside columngroup is characterized by a decreasing gradient along column rank one to rank twelve. This might provide an explaination for significant difference in thousand grain weight also along the the column blocks. As expected, the yield associated trait benefitted by the reduced disease pressure along the column block. In a multi-year study undertaken to identify definitive traits and their relative importance in improving resistance to Helminthosporium blight along with yield related traits, R. Sharma & Duveiller (2003) reported similar negative correlation between resistance to foliar disease and the kernel weight.

However, genotypes which manifested a sharp drop in relative chlorophyll content of the flag leaf just after antheis and before milking stage generally gave higher yields. The drop in chlorophyll content has often been ascribed to drought tolerance, in which case a higher drop in chlorophyll content is associated with drought sensitivity (Khakwani et al., 2013) (authors used chlorophyll fluorescence

extinction measurement as) and irrigated cultivars in general showed higher decline in chlorophyll content, under stress. In comparison, experimental condition of the field where genotypes were tested was irrigated prior to heading. Therefore, this might represent non-water-limiting/irrigated condition. This set of environmental conditions in combination with genotypic features that supress activated drought tolerance mechanism, might enable genotypes with higher LAUG values (120.1 and 92.18, respectively for period between anthesis and medium milk and medium milk and soft dough stages) to better realize their potential yields.

Findings on the association of relative chlorophyll content and the decline in chlorophyll content with the yield in wheat extend that of Khakwani et al. (2013), where significant correlation between yield and the indicator of leaf chlorophyll have been reported. Current study highlights, in particular, the importance of leaf chlorophyll content and the rate of decline of greenness at early stages in reproductive phase of the crop (X. Wu, Tang, Li, Wu, & Huang, 2015). For the leaf greenness trait at Z65, however leaving out the entry genotype "PRL/2*PAS-TOR/4/CHOIX/STAR/3/HE1/3*CNO79//...", which showed signs of severe depletion of green coloration of leaf early on life cycle, significant differences among entry genotypes could be inconclusive had the outlier been trimmed out.

5.2 Relationship between morphological and phenological traits and yield

With apparent positive but mild correlation between the days to heading and days to anthesis and the thousand kernel weight, the results from current study contrasts with observations by Duveiller, Kandel, Sharma, & Shrestha (2005), in which they report a negative association of days to maturity with the thousand kernel weight. Extending to their findings, authors clarify that differences in the trait association were not always significant for early sown wheat (planted between 26 Nov to 11 Dec). Inferring on grounds that current study had sowing date prior to that reported in the study (23 March), it is imperative that negative association do not hold. The study, however, fails to link the relationship between days to heading/anthesis and yield, which has been observed with negative but weak correlation in some studies (Mohammadi, Sharifi, Karimizadeh, & Shefazadeh, 2012).

5.3 Relationship between morphological and phenological traits and leaf health

Canopy architecture traits comprising plant height and flag leaf area had no significant correlation with any of the observed leaf health traits. To the contrast, canopy temperature depression showed weak negative correlation with the leaf greenness at Z75, suggesting that a larger drop in canopy temperature often occurs in genotypes with greenner leaves. Also, greenness of leaves and the relative chlorophyll content of leaves are both negatively associated with foliar disease specific symptoms. This phenomena along with correlated response to days to maturity has been explained as probable resistance mechanism to HLB in a study exploring selection index for HLB reistance in spring wheat (Duveiller et al., 2005).

Likewise, although negative correlation was found between days to heading and the foliar disease prevalence, it's significance of could not be validated. This is in contrast to the finding of Mahto (1999), who reported significant negative albeit weak correlation between the two variables.

5.4 Leaf health as determinant of yield

Decrease in error variance invariably adds to the heritability estimates and relative efficiency of a trait, thus maximizing chances of selecting true to the type genotypes (K. D. Bondalapati, Jenkins, McCarty, & Wu, 2015). Almost 52% reduction in the error variance was obtained by incorporating leaf health traits in the model. Similarly, leaf health traits can be ascribed to upto 39% variation in yield estimates. Among the traits of interest are the LAUG decline during Z65 and Z75, greenness scores at both Z75 and Z65 and the relative chlorophyll content at Z65. In a study by (Benbella & Paulsen, 1998) have also implied significant effect of greenness trait – defined in context of late maturity stage as "Stay green" – in yield determination and its association with foliar blight disease. However, significance of foliar blight complex disease in yield determination in experimental conditions of the present study was not evident. This is probably because environmental/spatial effects of disease were more relevant (as observed from significant columwise blocking) and the heterogeniety between blocks was much higher to directly ascribe yields levels. Also, comparing two distinct yield groups (one with high TKW and the other with low TKW) in a study of evaluating wheat genotypes, authors have reported no significant association of two yield groups and the disease

response (AUDPC) under timely sown conditions (R. Sharma, Tiwary, & Ortiz-Ferrara, 2008).

Application of mixed modeling procedures to field phenotyping in wide set of environmental conditions is justified in several previous studies (Qiao, Basford, DeLacy, & Cooper, 2000; Smith, Cullis, & Gilmour, 2001). Spatial covariate modeling along with leaf health also had beneficial effect on yield approximation, although exact quantification of effect estimates would require more advanced probabilistic techniques such as Jackknife resampling and bootstrapping (J. Wu et al., 2013; J. Wu, Jenkins, & McCarty, 2010).

With respect to estimated yield, all check cultivars were superior to Aditya variety and no distinction could be made between Bhrikuti, Gautam and Tilottama genotypes. While with genotypic BLUP estimates, ranking of genotypes places following three genotypes (on decending order) on top: L71 (TRCH/SRTU-//KACHU/3/-KINGBIRD #1), L67 (WHEAR/-SOKOLL/4/-PASTOR//MILAN/-KAUZ/3/B...) and L85 (MUNAL #1*2/4/-HUW234+LR34/PRINIA-//PBW3...). Pedigree information of the entry genotypes at foremost ranks suggests that genotypes composed of "MUNAL" in their pedigree with higher order crossings or recurrent crossing, preferably as female parent has tendency for better yields.

¹Order of crossing is denoted by a number sandwiched between two backslash ('/')

Conclusions

Yield approximation, having significant role in genotype selection, could be much improved by adding leaf health factor in screening and evaluation programs. Likewise, even in classical breeding traits, i.e. crop architecture such as plant height and canopy temperature depression, accuracy of phenotype based selection could be improved through adoption of better designs, which can address multiple sources of spatial heterogeniety in the field. In turn, these extraneous variation could be accounted for through mixed modeling procedures. Therefore, besides improving the precision of estimation with inclusion of leaf health traits, in particular greenness and relative chlorophyll content measures at early post-anthesis period, certain traits such as thousand kernel weight, plant height, flag leaf area, canopy temperature depression, days to heading and days to anthesis benefitted the most by modeling of spatial covariates.

Appendix A

I

Tables referenced in Chapter 4.1:

Table A.1: ANOVA table with likelihood ratio test for Yield

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	442		-214	428	2.60	1	0.055
Blocking factors + Entry genotypes	8	440	467	-212	424	3.68	1	0.055
Blocking factors +	12	434	474	-205	410	14.41	4	0.006
Check and entry genotypes								

Table A.2: ANOVA table with likelihood ratio test for Number of effective tillers

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors Blocking factors +	7 8		1597 1593		1559 1549	9.73	1	0.002
Entry genotypes Blocking factors + Check and entry genotypes	12	1538	1580	-757	1514	35.15	4	0.000

Table A.3: ANOVA table with likelihood ratio test for Thousand kernel weight

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1216	1239	-601	1202			
Blocking factors +	8	1186	1212	-585	1170	32.0	1	0.000
Entry genotypes								
Blocking factors +	12	1184	1223	-580	1160	10.3	4	0.036
Check and entry								
genotypes								

Table A.4: ANOVA table with likelihood ratio test for Number of grains per panicle

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	946	969	-466	932			
Blocking factors +	8	936	962	-460	920	12.4	1	0.000
Entry genotypes								
Blocking factors +	12	932	972	-454	908	11.6	4	0.021
Check and entry								
genotypes								

Table A.5: ANOVA table with likelihood ratio test for Panicle length

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	869	893	-428	855			
Blocking factors +	8	859	886	-421	843	12.8	1	0.000
Entry genotypes								
Blocking factors +	12	852	893	-414	828	14.6	4	0.006
Check and entry								
genotypes								

Tables referenced in Chapter 4.2:

Table A.6: ANOVA table with likelihood ratio test for Disease score at Zadok's stage 75

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	871	895	-428	857			
Blocking factors +	8	856	884	-420	840	16.6	1	0.000
Entry genotypes								
Blocking factors +	12	850	892	-413	826	13.9	4	0.008
Check and entry								
genotypes								

Table A.7: ANOVA table with likelihood ratio test for Flag leaf greenness score at Zadok's stage 65

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	306	331	-146	292			
Blocking factors +	8	263	291	-123	247	45.50	1	0.000
Entry genotypes								
Blocking factors +	12	266	308	-121	242	4.83	4	0.305
Check and entry								
genotypes								

Table A.8: ANOVA table with likelihood ratio test for Flag leaf greenness score at Zadok's stage 75

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	867	891	-426	853			
Blocking factors +	8	699	727	-341	683	169.8	1	0
Entry genotypes								
Blocking factors +	12	674	716	-325	650	32.8	4	0
Check and entry								
genotypes								

Table A.9: ANOVA table with likelihood ratio test for Flag leaf greenness score at Zadok's stage 85

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	889	914	-438	875			
Blocking factors +	8	848	876	-416	832	43.4	1	0.000
Entry genotypes								
Blocking factors +	12	842	884	-409	818	14.2	4	0.007
Check and entry								
genotypes								

Table A.10: ANOVA table with likelihood ratio test for LAUG between Zadok's stage 65 and 75

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1828	1852	-907	1814			
Blocking factors +	8	1757	1785	-871	1741	72.79	1	0.000
Entry genotypes								
Blocking factors +	12	1759	1801	-867	1735	6.22	4	0.184
Check and entry								
genotypes								

Table A.11: ANOVA table with likelihood ratio test for LAUG between Zadok's stage 75 and 85

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	2174	2199	-1080	2160			
Blocking factors +	8	2088	2116	-1036	2072	88.6	1	0
Entry genotypes								
Blocking factors +	12	2074	2116	-1025	2050	21.6	4	0
Check and entry								
genotypes								

Tables referenced in Chapter 4.3:

Table A.12: ANOVA table with likelihood ratio test for Plant height

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1565	1589	-775	1551			
Blocking factors +	8	1567	1594	-775	1551	0.0	1	1
Entry genotypes								
Blocking factors +	12	1493	1534	-734	1469	82.2	4	0
Check and entry								
genotypes								

Table A.13: ANOVA table with likelihood ratio test for Flag leaf surface area

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors Blocking factors +	7 8	1676 1678	1700 1706	-831 -831	1662 1662	0.00	1	1.000
Entry genotypes Blocking factors + Check and entry	12	1678	1720	-827	1654	7.68	4	0.104
genotypes								

Table A.14: ANOVA table with likelihood ratio test for SPAD at Zadok's stage 65

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1238	1262	-612	1224			
Blocking factors +	8	1230	1258	-607	1214	9.59	1	0.002
Entry genotypes								
Blocking factors +	12	1229	1271	-603	1205	8.95	4	0.062
Check and entry								
genotypes								

Table A.15: ANOVA table with likelihood ratio test for SPAD at Zadok's stage 75

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1608	1632	-797	1594			
Blocking factors +	8	1610	1638	-797	1594	0	1	1
Entry genotypes								
Blocking factors +	12	1597	1639	-786	1573	21	4	0
Check and entry								
genotypes								

Table A.16: ANOVA table with likelihood ratio test for Canopy temperature depression (CTD)

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	794	819	-390	780			
Blocking factors +	8	779	807	-382	763	17.2	1	0.000
Entry genotypes								
Blocking factors +	12	775	817	-375	751	12.4	4	0.015
Check and entry								
genotypes								

Table A.17: ANOVA table with likelihood ratio test for Days to heading

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1393	1417	-689	1379			
Blocking factors +	8	1192	1220	-588	1176	202	1	0
Entry genotypes								
Blocking factors +	12	1045	1087	-511	1021	155	4	0
Check and entry								
genotypes								

Table A.18: ANOVA table with likelihood ratio test for Days to anthesis

term	df	AIC	BIC	logLik	deviance	statistic	Chi.Df	p.value
Blocking factors	7	1427	1451	-706	1413			
Blocking factors +	8	1228	1255	-606	1212	201.2	1	0
Entry genotypes								
Blocking factors +	12	1150	1191	-563	1126	85.9	4	0
Check and entry								
genotypes								

Appendix B

II

Snapshot of all statistical packages and processing environment used in analysis and interpretation of data:

```
R version 3.4.4 (2018-03-15)
Platform: x86_64-w64-mingw32/x64 (64-bit)
Running under: Windows 8.1 x64 (build 9600)
Matrix products: default
locale:
[1] LC_COLLATE=English_United States.1252
[2] LC_CTYPE=English_United States.1252
[3] LC_MONETARY=English_United States.1252
[4] LC_NUMERIC=C
[5] LC_TIME=English_United States.1252
attached base packages:
             graphics grDevices utils datasets methods base
[1] stats
other attached packages:
 [1] ggfortify_0.4.4 bindrcpp_0.2.2 Hmisc_4.1-1
 [4] Formula_1.2-2 survival_2.41-3 lattice_0.20-35
```

[7]	xtable_1.8-2	lmerTest_2.0-36	lme4_1.1-16
[10]	Matrix_1.2-12	forcats_0.3.0	stringr_1.3.0
[13]	purrr_0.2.4	readr_1.1.1	tidyr_0.8.0
[16]	tibble_1.4.2	tidyverse_1.2.1	thesisdowndss_0.0.1
[19]	knitr_1.20	bookdown_0.7	ggplot2_2.2.1
[22]	dplyr_0.7.4	devtools_1.13.5	
loade	ed via a namespace (a	and not attached):	
[1]	nlme_3.1-131.1	lubridate_1.7.4	RColorBrewer_1.1-2
[4]	httr_1.3.1	rprojroot_1.3-2	tools_3.4.4
[7]	backports_1.1.2	R6_2.2.2	rpart_4.1-13
[10]	lazyeval_0.2.1	colorspace_1.3-2	nnet_7.3-12
[13]	withr_2.1.2	gridExtra_2.3	mnormt_1.5-5
[16]	compiler_3.4.4	cli_1.0.0	rvest_0.3.2
[19]	htmlTable_1.11.2	xml2_1.2.0	labeling_0.3
[22]	scales_0.5.0	checkmate_1.8.5	psych_1.8.3.3
[25]	digest_0.6.15	foreign_0.8-69	minqa_1.2.4
[28]	rmarkdown_1.9	base64enc_0.1-3	pkgconfig_2.0.1
[31]	htmltools_0.3.6	MuMIn_1.40.4	htmlwidgets_1.2
[34]	rlang_0.2.0	readxl_1.0.0	rstudioapi_0.7
[37]	bindr_0.1.1	jsonlite_1.5	acepack_1.4.1
[40]	magrittr_1.5	kableExtra_0.8.0	Rcpp_0.12.16
[43]	munsell_0.4.3	stringi_1.1.7	yaml_2.1.18
[46]	MASS_7.3-49	plyr_1.8.4	grid_3.4.4
[49]	parallel_3.4.4	crayon_1.3.4	haven_1.1.1
[52]	stargazer_5.2.1	splines_3.4.4	hms_0.4.2
[55]	pillar_1.2.1	reshape2_1.4.3	codetools_0.2-15
[58]	stats4_3.4.4	glue_1.2.0	evaluate_0.10.1
[61]	latticeExtra_0.6-28	data.table_1.10.4-3	modelr_0.1.1
[64]	nloptr_1.0.4	cellranger_1.1.0	gtable_0.2.0
[67]	assertthat_0.2.0	xfun_0.1	broom_0.4.4
[70]	<pre>viridisLite_0.3.0</pre>	memoise_1.1.0	cluster_2.0.7-1

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