

Assessment of Genetic Gain Trends for Yield in IRRI Rice Varieties in the Philippines Using “era” Trial Studies and Implications for Future Rice Breeding

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2 **Philippines using “Era” trial studies and implications for future rice breeding**

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15 **Abstract**

16 Rice is a staple crop for 3.5 billion people in the world. To meet the challenges of the rice production for
17 food security and demand due to population increase, yield improvement due to a rice variety’s genetic
18 characteristics is imperative. Two studies presented in this paper were undertaken at the International
19 Rice Research Institute (IRRI) in the Philippines, to assess genetic gains for yield in rice varieties bred over
20 the past 50 years. These studies are called as “Era” studies as the varieties used for trials were released
21 during long and distinct periods. Due to the differences in time periods of studies, varieties and locations,
22 the studies were treated separately so as to not to compromise the data analyses. The studies
23 demonstrated that IRRI developed varieties have achieved genetic gains and levels of genetic gains were

24 dependent on correction or otherwise for maturities. In Study 1, the highest level of genetic gain was
25 0.70% at about a 23 kg ha⁻¹ annual yield increase when not corrected for maturity followed by a genetic
26 gain of 0.62% when corrected for maturity. In Study 2, the highest level of genetic gain was 0.74% at about
27 a 19 kg ha⁻¹ annual yield increase when corrected for maturity followed by 0.66% genetic gain when not
28 corrected for maturity. Implications for breeding programs are discussed, however, the studies were not
29 intended to compare genetic gains achieved through different breeding methods nor to compare genetic
30 gains achieved using plot trials versus realized genetic gains for crops grown under farmers' management.

31

32 **Key words:** Rice varieties, genetic gain, Era trials

33 **1. BACKGROUND**

34 Rice is the staple food for 3.5 billion people. Within this decade, an estimated 40 million tons of additional
35 rice will be needed to feed the increasing global population (OECD/FAO, 2019). Climate change is
36 projected to decrease yield in major rice-exporting countries, such as Thailand and Vietnam in Southeast
37 Asia (Teng et al., 2016), impacting on the regional importing countries and rice reserves needed for food
38 security. In the 1960s, the International Rice Research Institute (IRRI) led the “Green Revolution” in rice
39 with the release of variety IR8, especially in South Asia (Khush, 1999). New rice varieties have since been
40 released over the past 55 years in many countries of Asia and other parts of the world to increase rice
41 production through improved yield combined with resistance to and/or tolerance of pests and diseases.

42

43 It is projected that, due to climate change—especially temperature increases—there will be a direct
44 impact due to reduction in the yields of four major crops including in rice (Zhao et al., 2017). Rice will be
45 globally impacted indirectly due to increased water demand (Zhao et al., 2017). Due to population growth,
46 consumption of rice is set to increase to over 578 million tons with the average consumption of rice
47 globally projected to increase by 1 kg to reach 55 kg annually during 2019–28, while the area utilized for

48 rice production within this decade is expected to increase by only 1% (OECD/FAO 2019) Therefore,
49 increase in rice production on the farm needs to be achieved by yield improvement of new varieties
50 coupled with optimum agronomic practices. It is imperative that plant breeding programs release new
51 climate-resilient varieties that have higher yields under variable climatic conditions to overcome the
52 challenges to food security.

53 Genetic gains for key traits, including yield, are imperative for successful adoption of varieties at the farm
54 gate level for sustainability and farmer profitability. Genetic gain can be simply defined as the amount of
55 increase in performance that is achieved annually through artificial selection (Xu et al., 2017). To
56 understand the progress made in plant breeding of the crops through artificial selection and breeding,
57 assessment of genetic gain for yield has been conducted in different major crops. Assessment of genetic
58 trends in varieties can provide an estimate of the rate of genetic gain for yield achieved in the breeding
59 program where the varieties originated.

60 Various researchers have used different methods to analyze and estimate genetic gain using a large
61 number of historical datasets generated and collated from different breeding programs that have
62 conducted trials using different varieties. In the United States, it was estimated that maize breeding
63 resulted in genetic gains of 63 kg ha⁻¹ year⁻¹ for the first 30 years between 1930 and 1960 and of 110 kg
64 ha⁻¹ year⁻¹ over the next 40 years (Duvick, 2005a). In the wheat breeding program at the International
65 Maize and Wheat Improvement Center (CIMMYT) in Mexico, Crespo-Herrera et al. (2018) assessed genetic
66 gain and showed that, for grain yield, it was 38 kg ha⁻¹ year⁻¹ (1.8%) in low-yield environments and 57 kg
67 ha⁻¹ year⁻¹ (1.4%) in medium-yielding environments; across the environments, yield gain was 48 kg ha⁻¹
68 year⁻¹ (1.6%). In Germany, Laidig et al. (2014), who analyzed genetic trends for yield improvement across
69 environments for 12 different crop type varieties released over 30 years from different breeding
70 programs, concluded that all the improved crops species tested showed improvements in yield; the lowest

71 genetic trend for dry matter yield was in Italian ryegrass (0.16% annually) and the highest was for winter
72 canola or winter oil seed rape (1.86% annually).

73 In rice, there have been very limited genetic gain studies conducted, especially in Asia where rice is the
74 main staple grain. The only study prior to this one in Southeast Asia was done by Peng et al. (2000) at the
75 International Rice Research Institute, Los Banos, Philippines in 1996 with an estimated 1% yield (75-81
76 kg/ha) improvement per year. In Brazil, Breseghello et al. (2011) conducted genetic gain analyses for 25
77 years of upland rice breeding and concluded that annual gain for grain yield was not significant between
78 1984 and 1992. However, it was 15.7 kg ha⁻¹ year⁻¹ (0.53%) between 1992 and 2002; and nearly tripled
79 between 2002 and 2009 with a gain of 45.0 kg ha⁻¹ year⁻¹ (1.44%). More recently in India, Kumar et al.
80 (2020) reported genetic gains of 0.68–1.9% for grain yield in rice varieties grown between 2005 and 2014
81 under different levels of moisture stress conditions.

82 The IRRI breeding program over the decades has had defined objectives in each decade of the program
83 (Peng and Khush 2003). In the first decade (1960-1969) due to lodging among tall rice varieties leading to
84 lower yield, focus was on incorporation of gene conferring lower height (dwarfism) commencing with IR8
85 which led to higher yield. During 1970–1979, emphasis was on multiple disease and insect resistances.
86 During 1980–1989, emphasis was on good grain quality that resulted in IR64. During 1990–1999, hybrids
87 and new plant-type breeding incorporated yield gain with biotic stress resistance and gain quality. By
88 2000–2009, more emphasis was given to molecular breeding approaches to enhance various desired traits
89 and yield improvement and, later, abiotic stress tolerances. IRRI breeders focused on developing high-
90 yielding varieties with multiple stresses, pests and disease tolerances in predominantly irrigated and
91 rainfed environments. Therefore, assessing genetic gain is necessary in such a mature breeding program
92 to understand the progress made during the life of the program and to ensure that higher-yielding
93 varieties continue to be released in the future.

94 Genetic gain for yield and other traits can be assessed by utilizing historical data from breeding programs
95 although this can be labor- and resource-intensive and requires collation of large sets of consistent
96 standardized historical data. In maize, long-term studies of sequential series of successful hybrids were
97 assessed by adding data from additional new varieties (Duvick et al., 2005a). In their review, Smith et al.
98 (2014) refer to the term “Era trials used by Duvick (2005b) for the method to assess yield improvements
99 in maize crops by including varieties released over different years using data from evaluations conducted
100 in single or across multiple environments (Multi-Environment Trials or MET). Although there are several
101 methods with their own limitations to assess genetic gain for historically-released varieties, Rutkoski
102 (2019) showed that “Era” trials can be used to estimate rates of genetic gain in a shorter timeframe and
103 genetic performance of varieties can be assessed with this method in lieu of large sets of interconnected
104 historical data.

105 In rice, it is critical that genetic gain for yield is achieved to meet increasing demands as mentioned in
106 OECD/FAO (2019). It is important to understand if increased yields have been achieved through plant
107 breeding and selections (Duvick, 2005a) and thereby drive the changes needed in breeding strategies and
108 methods to achieve targeted yield gain outcomes. Therefore, to understand genetic gain achieved in IRRI’s
109 rice breeding program over the past 50 years, two “Era” studies were undertaken during different time
110 periods using different number and combinations of varieties with the second study including later-
111 released varieties. It was expected that these studies would contribute to learning more about the genetic
112 gains achieved in rice thus far at IRRI. Additionally, it was expected that the new knowledge would lead
113 to further exploration of breeding strategies at IRRI to improve genetic gain and provide a clear
114 understanding of the gains yet to be achieved.

115 **2. MATERIALS AND METHODS**

116 This paper describes two “Era” studies conducted at IRRI in Los Baños, Philippines, utilizing its bred
117 varieties in field trials. The varieties used were released from 1966 through to those released at the

118 respective times that the two studies were conducted. Plot trials conducted at the locations were used to
 119 collect the data. Study 1 used 44 varieties and lines available at the time (Table 1) and was undertaken
 120 during two dry seasons and two wet seasons during 2013 and 2015. Study 2 used 22 varieties (Table 2)
 121 was undertaken during one dry season and two wet seasons during 2016 and 2017. Different sets of
 122 varieties were used in Study 1 and Study 2. Only 12 entries were common between Study 1 and Study 2
 123 and therefore data for Study 1 and 2 was not combined for analysis.as that would only lead to a lower
 124 number of common lines being taken into account for analyses.

125 For Study 2, IRRI Multi Environment Trials (MET, Supplementary Table 1) additional data was included in
 126 the analyses for some varieties common between “Era” and MET trials adding to the data robustness.
 127 These MET trials were conducted in wet season in 2016 and two dry seasons in 2016 and 2017, in one
 128 common location Los Banos between “Era” trial and in additional locations in the Philippines. These
 129 additional locations in the Philippines included Midsayap, Muñoz, Musuan, Agusan, Ubay, and San Mateo.

130 **Variety selection for Study 1**

131 Forty-four IRRI-developed entries, released between 1966 and 2013 commencing with IR8 in 1966 to IRRI
 132 168 in 2013, were included in Study 1. These varieties were selected based on the area to which they were
 133 grown by farmers during their period of maximum use in Southeast Asia (Brennan & Malabayabas, 2011).
 134 Thirty-eight varieties had been released for irrigated environments. Two rainfed varieties (IRRI 119 and
 135 IRRI 164) and four hybrid varieties were included (Table 1).

136 **TABLE 1** Study 1 variety list by type and in chronological order by year of release

Serial number	Variety designation	Pedigree of the line	Category of variety type	Year of release
1	IR8	PETA/DEEGEOWOGEN	Irrigated	1966
2	IR20	IR262-24-3/TKM6	Irrigated	1969
3	IR24	IR8/IR127-2-2	Irrigated	1971

4	IR26	IR24/TKM6	Irrigated	1973
5	IR32	IR20*2/ONIVARA//CR94-13	Irrigated	1975
6	IR36	IR1561-228-1-2/IR1737//CR94-13	Irrigated	1976
7	IR42	IR1561-228-1-2/IR1737//CR94-13	Irrigated	1977
8	IR50	IR2153-14-1-6-2/IR28//IR36	Irrigated	1979
9	IR58	IR28/KWANGCHANGAI//IR36	Irrigated	1983
10	IR60	IR4432-53-33/PTB33//IR36	Irrigated	1983
11	IR62	ARIKARAI///IR24/TKM6//IR20*4/ONIVARA/4/IR 1561-228-1-2/IR1737//CR94-13	Irrigated	1984
12	IR64	IR5657-33-2-1/IR2061-465-1-5-5	Irrigated	1985
13	IR66	IR13240-108-2-2-3/IR9129-209-2-2-2-1	Irrigated	1987
14	IR70	IR19660-73-4/IR54//IR9828-36-3	Irrigated	1988
15	IR72	IR19661-9-2-3-3/IR15795-199-3-3//IR9129-209- 2-2-2-1	Irrigated	1988
16	IR74	IR19661-131-1-2/IR15795-199-3-3	Irrigated	1988
17	IRRI 102	IR4215-301-2-2-6/BG90-2//IR19661-131-1-2	Irrigated	1991
18	IRRI 103	IR4547-4-1-2/IR1905-81-3-1//IR25621-94-3-2	Irrigated	1991
19	IRRI 104	IR33021-39-2-2/IR32429-47-3-2-2	Irrigated	1992
20	IRRI 105	IR24594-204-1-3-2-6-2/IR28222-9-2-2-2-2	Irrigated	1994
21	IRRI 106	IR35293-125-3-2-3/IR32429-47-3-2-2//IRRI 103	Irrigated	1994
22	IRRI 108	IR28239-94-2-3-6-2/IR64	Irrigated	1995
23	IRRI 109	IR72/IR24632-34-2	Irrigated	1995
24	IRRI 115	IR48613-54-3-3-1/IR28239-94-2-3-6-2	Irrigated	1997
25	IRRI 116	IR72/IR48525-100-1-2	Irrigated	1997
26	IRRI 118	IRRI102/IR39292-142-3-2-3	Irrigated	1997
27	IRRI 119	IR43581-57-3-3-6/IR26940-20-3-3-3- 1//KHAODAWKMALI105	Rainfed	1997
28	IRRI 121	IR58025A/IR34686-179-1-2-1R	Hybrid	1997

29	IRRI 122	IR50401-77-2-1-3/IR42068-22-3-3-1-3	Irrigated	2000
30	IRRI 123	IR47761-27-1-3-6/IRRI108	Irrigated	2000
31	IRRI 135	IR1561*3/HABIGANJDW8//5*IR64	Irrigated	2002
32	IRRI 136	IR1561*2/OBARTHII//4*IR64	Irrigated	2002
33	IRRI 138	IR68897A/IR60819-34-2R	Hybrid	2002
34	IRRI 141	IR44625-139-2-2-3/IR32822-94-3-3-2-2	Irrigated	2003
35	IRRI 144	IR68897A/IR71604-4-1-4-4-2-2-2R	Hybrid	2006
36	IRRI 145	IR44699-21-1-3-4/IR66438-167-3-3-2-3	Irrigated	2007
37	IRRI 146	IRRI134/IR70479-45-2-3//IR64680-81-2-2-1-3	Irrigated	2007
38	IRRI 150	IR68077-82-2-2-2-3/IR59548-122-1-4-1	Irrigated	2007
39	IRRI 151	IR00A112/IRRI106	Irrigated	2009
40	IRRI 154	IR73012-137-2-2-2/IRRI104	Irrigated	2009
41	IRRI 156	IR72870-102-2-2-1-3/IR72870-19-2-2-3	Irrigated	2011
42	IRRI 158	IR80559A/IR60819-34-2R	Hybrid	2011
43	IRRI 164	IR43/IR65564-22-2-3//IR68	Rainfed	2011
44	IRRI 168	IR72904-65-1-3-3/IR73012-137-2-2-2	Irrigated	2013

137

138 **Variety selection for Study 2**

139 The entries for Study 2 were selected based on their year of release starting from the release of IR8 in
140 1966. Entries included IRRI varieties (Table 2) developed for irrigated rice in Southeast Asia. Therefore, 22
141 varieties released up to 2016 for irrigated environments were selected. However, unlike in the Study 1,
142 rainfed entries were used in separate trials (data not presented here) as rainfed varieties performance
143 under irrigated conditions may be very different under non-water limited conditions. Hybrid entries were
144 not used along with inbred entries as they are generally considered to yield higher than inbred varieties.
145 Additionally, data from varieties common to all “Era” trials and the MET (Supplementary Table 1) trials
146 conducted in seven locations including Los Baños, in the Philippines (2016–2017) were included in the

147 analysis. This was done to increase data robustness as it would involve several environments other than
 148 the one environment used in the “Era” trial.

149 **TABLE 2** Study 2 variety list by type and in chronological order by year of release

Serial number	Variety designation	Pedigree of the line	Category of variety type	Year of release
1	IR8	PETA/DEEGEOWOOGEN	Irrigated	1966
2	IR24	IR8/IR127-2-2	Irrigated	1971
3	IR34	BPI 76/CHIANUNG YU 280	Irrigated	1975
4	IR36	IR1561-228-1-2/IR1737//CR94-13	Irrigated	1976
5	IR52	BPI 76*2/KAOHSIUNG 68	Irrigated	1980
6	IR64	IR5657-33-2-1/IR2061-465-1-5-5	Irrigated	1985
7	IR 72	IR19661-9-2-3-3/IR15795-199-3-3//IR9129-209-2-2-2-1	Irrigated	1988
8	IRRI 104	IR33021-39-2-2/IR50404	Irrigated	1992
9	IRRI 105	IR 24594-204-1-3-2-6-2/IR 28222-9-2-2-2-2	Irrigated	1994
10	IRRI 108	IR 28239-94-2-3-6-2/IR64	Irrigated	1995
11	IRRI 115	IR48613-54-3-3-1/IR28239-94-2-3-6-2	Irrigated	1997
12	IRRI 123	IR47761-27-1-3-6/IRRI108	Irrigated	2000
13	IRRI 141	IR44625-139-2-2-3/IR32822-94-3-3-2-2	Irrigated	2003
14	IRRI 143	IR64 (WH)/ADAY SEL//3*IR 64	Irrigated	2006
15	IRRI 154	IR73012-137-2-2-2/PSB RC10 (IR50404-57-2-2-3)	Irrigated	2009
16	IRRI 174	IR00A117/IR 64	Irrigated	2013
17	IRRI 179	IR73008-138-2-2-2/IR68544-29-2-1-3-1-2//IR72870-19-2-2-3	Irrigated	2014
18	IRRI 180	ADDAY LOCAL SEL	Irrigated	2014
19	IRRI 181	IR80894-18-2-2-3	Irrigated	2014

20	IRRI 186	IR02A127/IR 64	Irrigated	2015
21	IRRI 192	Huang-Hua-Zhan*2/OM 1723	Rainfed	2016
22	IRRI 193	IR 68077-82-2-2-2- 3/IR00A117	Irrigated	2016

150

151 **Field trials and environments for Study 1**

152 All trials were conducted on IRRI’s experiment station at Los Baños. The field trials were replicated over 2
 153 years during the 2013 and 2014 wet seasons (WS) and the 2014 and 2015 dry seasons (DS). Generally, in
 154 the Philippines, the monsoon wet season has low solar radiation and high relative humidity and pest and
 155 disease pressure. During the dry season, temperature, evaporative demand, and solar radiation are high
 156 and relative humidity and pest disease pressures are low.

157 **Trial design and data collection for Study 1**

158 Field trials were a Randomized Complete Block Design (RCBD) with four replications. Each plot size was
 159 10.8 m² (5.4m x 2.0 m) with 10 rows (20 cm x 20-cm spacing with two or three plants hill⁻¹). Plot sizes were
 160 as recommended by IRRI statisticians and used in IRRI trials over the years.

161 Trial data were collected and collated for analysis at appropriate stages following IRRI’s standard
 162 evaluation system (SES; IRRI, 2014). Data collected included flowering time (FLW), plant height (PH), and
 163 number of productive tillers (TN); maturity was calculated by adding 30 days to the 50% flowering date.
 164 Grain yield was determined by harvesting 25 hills per row from the six inner rows (harvest area was 6.0
 165 m²), adjusting to 14% moisture content, and calculating yield in kg/ ha⁻¹. Yield day⁻¹ was calculated by
 166 dividing grain yield by the number of days to maturity.

167 **Field trials and environments for Study 2**

168 These “Era” trials represented 2 years 3 seasons during 2016 wet season and 2017 wet season and 2017
 169 dry season at IRRI Los Baños, Philippines. Seeding and transplant were done in accordance with the

170 appropriate seeding and transplanting practices used for the IRRI MET trials. Manual harvesting was
171 conducted; border rows were not harvested. Data were collected on any missing hills and were accounted
172 for in the analysis.

173 Simultaneous with the “Era” trials conducted in a single location at Los Baños, Multi Environment Trials
174 (MET) were also conducted in 2016 and 2017 wet and dry seasons in 7 locations across Philippines in
175 Munoz, Midsayap, Musuan, Augusan, Isabella Ubay and Los Baños.

176 **Trial design and data collection for Study 2**

177 For both “Era” trials and Multi Environment Trials (MET) trials, the trial plot design was Alpha Lattice
178 except for MET level 2 2017 wet season trials which was Partially Balanced Incomplete Block design.
179 Individual plot size was uniformly 5.4 meters x 2 meters with 26 hills for length and 10 hills at width of
180 each plot. Border rows were not harvested to minimize border effects. The data was collected from “Era”
181 trials varieties from one wet season each in 2016 and 2017, and one dry season in 2017. Data was also
182 collected for one dry season in 2017 and two wet seasons in 2016 and 2017 from MET1 and MET 2 level
183 trials conducted in all the Philippines locations including in Los Baños location

184 **Data analyses**

185 Only yield and maturity data were considered for analyses in the two studies. Estimations of genetic gain
186 and analyses are detailed below. Pest and disease and environmental data as co-variates were not
187 considered in these analyses. LS Means for Study 1 are presented in Supplementary Table 2. Trial
188 heritability estimates for Study 1 are in Supplementary Table 3. LS Means for Study 2 are in Supplementary
189 Table 4. Trial heritability estimates for Study 2 are in Supplementary Table 5. Details of methods used for
190 the data analyses are described below. Pedigrees of the varieties were not used for phenotypic data
191 analyses.

192

193

194 **Relationship among varieties**

195 Kinship matrices account for relationships among individuals and the higher the value, the closer the
196 relationship between the varieties or lines, in general. Kinship matrices for varieties in the two studies
197 were done using the pedigrees recorded for the entries. Kinship among the rice varieties/lines was
198 estimated in both studies to understand if the relationships among varieties could contribute either
199 positively or negatively to the estimated genetic gain. Very close relationships among varieties could
200 narrow the genetic pool and might adversely contribute to the estimated genetic gain.

201 **Statistical analyses**

202 **Study 1**

203
204 Phenotypic data for grain yield were analyzed separately by season (wet and dry) using the mixed model:

205
$$y_{ijklm} = \mu + g_i + s_j + gs_{ij} + b_k + \varepsilon_{ijk} \quad (1)$$

206 where y_{ijk} is the grain yield measurement for the i th genotype in the j th year and k th complete block; μ
207 is the overall mean; g_i is the fixed effect of genotype; s_j is the random effect of the specific season; gs_{ij} is
208 the random effect of the interaction between the genotype and the specific season; b_k is the random
209 effect of complete block, also referred to as replicate; and ε_{ijklm} is the residual error.

210 All random effects were assumed independently and identically distributed (i.i.d). To correct for days to
211 maturity, days to maturity collected on the same plot as the grain yield data was included in model 1.

212 From model 1, the least-square means and standard errors for grain yield and grain yield corrected for
213 days to maturity were estimated per variety, per season. To obtain a combined mean across seasons,
214 these least-square means were used as the response variable in the linear regression model:

215
$$y_{ij} = \mu + g_i + w_j + \varepsilon_{ij} \quad (2)$$

216 where y_{ij} is the mean yield for genotype i and season j , g_i is the effect of genotype, w_j is the effect of the
217 weather season, and ε_{ij} is the residual error assumed i.i.d.

218 From model 2, the least-square mean for grain yield and grain yield corrected for days to maturity across
219 all seasons were estimated.

220 To assess genetic trends in varieties, means data were used as the response variable in the linear
221 regression model

$$222 \quad y_i = \mu + \alpha d_i + \varepsilon_i \quad (3)$$

223 where y_i is the mean yield of variety i ; d_i is the continuous year date when the variety was released minus
224 1966, which was the earliest variety release year in the dataset; α is the fixed regression coefficient for
225 the genetic trend in yield; and ε_{ijk} is the residual error with $\varepsilon_i \sim N(0, R\sigma_\varepsilon^2)$ where R is a matrix with the
226 diagonal equal to the squared standard error of the least-square means.

227 Model 3 was fitted using the combined means across seasonal data, using wet season data, and using dry
228 season data separately. Model 3 was fitted for both grain yield and grain yield corrected for maturity.

229

230 **Study 2**

231

232 Phenotypic data used for this study originated from both the MET and "Era" trials. For each trial containing
233 varieties sampled for this study that follows the alpha lattice design, phenotypic data for yield and days
234 to maturity were analyzed using the mixed model:

$$235 \quad y_{ijk} = \mu + g_i + r_j + b_{k(j)} + \varepsilon_{ijk} \quad (4)$$

236 where y_{ijk} is the phenotypic measurement for the i th genotype in the j th replicate and k th block; μ is the
237 trial mean; g_i is the effect of genotype; r_j is the random effect of replicate; $b_{k(j)}$ is the random effect of
238 block nested within j th replicate; and ε_{ijk} is the residual error. All random terms were assumed i.i.d. While
239 for MET data that uses the partially balanced incomplete block design, phenotypic data for yield and days
240 to maturity were analyzed using the mixed model:

241 $y_{ij} = \mu + g_i + b_j + \varepsilon_{ij}$ (5)

242 where y_{ij} is the phenotypic measurement for the i th genotype in the j th block; μ is the trial mean; g_i is
243 the effect of genotype; b_j is the random effect of block; and ε_{ij} is the residual error. All random terms
244 were assumed i.i.d. From models 4 and 5, least-square means and standard errors were computed for
245 yield and days to maturity per trial.

246 The trial-specific least-square mean yields for each variety sampled for Study 2 were used as the response
247 variable in the mixed model:

248 $y_{ijk} = \mu + g_i + s_j + t_k + gs_{ij} + \varepsilon_{ijk}$ (6)

249 where y_{ijk} is the least square mean for the i th genotype in the j th season and k th trial; μ is the mean; g_i
250 is the effect of genotype; s_j is the effect of the specific growing season; t_k is the random effect of trial
251 assumed i.i.d.; $gs_{i,j}$ is random effect of the interaction between genotype i and season j assumed i.i.d.;
252 and $\varepsilon_{ijk} \sim N(0, R\sigma_\varepsilon^2)$ where R is a matrix with the diagonal equal to the squared standard error of the
253 least-square means. To correct for maturity, a co-variate for days to maturity was included in model 6.
254 Model 6 was fitted using both wet and dry seasons data, using wet season data alone, and using dry
255 season data alone. Least-square means and standard errors were then computed for grain yield and grain
256 yield corrected for maturity for each variety across seasons and in dry and wet seasons separately.

257 Genetic trends in varieties were assessed using model 3 as in Study 1.

258

259 **Software used**

260 Analyses for the two studies were carried out in the R statistical programming language and environment
261 (R Development Core Team, 2016), using the packages *lme4* and *lsmeans* (Lenth, 2016). Kinship between

262 varieties included in both studies were estimated based on pedigree using the R package *nadiv* (Wolak,
 263 2012).

264 3. RESULTS

265 Results Summary

266 Genetic Trend Model Estimate of variety mean yields in Los Baños, Philippines has been presented in Table
 267 3 There was considerable variation for *per se* yield among varieties in both Study 1 (Supplementary LS
 268 Means Tables 2 and Supplementary trial heritabilities Table 3 respectively) and Study 2 (Supplementary
 269 LS Means Tables 4 and Supplementary trial heritabilities Table 5 respectively).

270 In Study 1, the highest level of genetic gain was in the wet season when data were not corrected for
 271 maturity. It was 0.70% with a yield increase of about 23 kg ha⁻¹year⁻¹. In Study 2, the highest level of
 272 genetic gain was 0.74% with a yield increase of about 19 kg ha⁻¹ year⁻¹ during the wet season when the
 273 data were corrected for maturity. There was a difference of about 4 kg ha⁻¹ year⁻¹ for the highest values
 274 of genetic gain between the studies (Table 3). These were followed by genetic gain of 0.62% (20 kg ha⁻¹
 275 year⁻¹) when corrected for maturity during the wet season in Study 1 and a genetic gain of 0.66% (17 kg
 276 ha⁻¹ year⁻¹) when not corrected for maturity in the wet season in Study 2.

277

278 **TABLE 3** Genetic Trend Model Estimate of variety mean yields for IRRI varieties in Studies 1 and 2

Study	Season	Controlled for maturity	Baseline yield± standard error	Increase of yield in kg ha ⁻¹ year ⁻¹	Yield gain per year	Model R Squared
1	Dry	No	6525.73 ± 214.82 (3.3e-30)	17.32 ± 7.13 (2.0e-02)	0.27%	0.12
1	Wet	No	3305.85 ± 166.73 (6.1e-23)	23.13 ± 5.54 (1.5e-04)	0.70%	0.29
1	Wet and dry	No	4915.87 ± 140.98 (1.2e-32)	20.23 ± 4.68 (9.4e-05)	0.41%	0.31
1	Dry	Yes	6541.79 ± 221.32 (9.8e-30)	16.49 ± 7.3 (2.9e-02)	0.25%	0.11

1	Wet	Yes	3376.64 ± 172.7 (1.0e-22)	20.81 ± 5.63 (6.3e-04)	0.62%	0.25
1	Wet and dry	Yes	4958.17 ± 147.32 (5.2e-32)	18.51 ± 4.89 (4.8e-04)	0.37%	0.25
2	Dry	No	3802.86 ± 327.72 (2.5e-10)	13.02 ± 8.98 (1.6e-01)	0.34%	0.10
2	Wet	No	2729.77 ± 158.09 (1.7e-13)	17.92 ± 4.28 (4.6e-04)	0.66%	0.47
2	Wet and dry	No	3171.47 ± 120.82 (5.7e-17)	17.35 ± 3.32 (4.2e-05)	0.55%	0.58
2	Dry	Yes	3866.37 ± 385.18 (3.0e-09)	14.79 ± 10.59 (1.8e-01)	0.38%	0.09
2	Wet	Yes	2624.13 ± 151.5 (1.7e-13)	19.4 ± 4.09 (1.2e-04)	0.74%	0.53
2	Wet and dry	Yes	3164.63 ± 153.99 (6.4e-15)	19.35 ± 4.26 (2.0e-04)	0.61%	0.51

279 Notes: 1. The values in the column containing baseline yield and standard error are followed by p values
280 in parentheses.

281 Note 2. Results from two independently conducted studies are shown for wet and dry seasons
282 combined, dry season alone, and wet season alone.

283

284 Table 3 shows an increase of yield in kg ha⁻¹ year⁻¹ and yield gain in year⁻¹ ha⁻¹ when controlled for maturity
285 or not controlled for maturity under the dry and wet seasons. Genetic trends in IRRI variety yields in Los
286 Baños, Philippines over a 50-year period, not controlled for maturity and controlled for maturity are
287 shown in Figures 1 and 2, respectively.

288 There were some differences for *per se* yield between the two studies. Study 1, in general, seems to have
289 a higher yield during 2013–2014 compared to Study 2 in 2016–2017 (Supplementary LS Means Tables 2
290 and 3, respectively). In practical terms, yield increases in kg ha⁻¹ year⁻¹ were not very different between
291 the two studies despite differences in the number of entries and locations.

292 During the when the data were not corrected for maturity in Study 1, yield increase was 17.32 kg ha⁻¹ year⁻¹
293 with a standard deviation of 7.13 kg ha⁻¹ and a genetic gain of 0.27% per year. Likewise, in Study 2, the
294 yield increase was 13.02 kg ha⁻¹ year⁻¹ with a standard deviation of 8.98 kg ha⁻¹ and a genetic gain of 0.34%
295 per year.

296 During the wet season when the data were not corrected for maturity in Study 1, the yield increase was
297 $23.13 \text{ ha}^{-1} \text{ year}^{-1} \pm 5.54 \text{ kg ha}^{-1}$ with a percent genetic gain of 0.70% per year. However, for the when data
298 were not corrected for maturity in Study 2, the yield increase was $17.92 \text{ ha}^{-1} \text{ year}^{-1} \pm 4.28 \text{ kg ha}^{-1}$ with a
299 genetic gain of 0.66% per year. The yield increases in Studies 1 and 2 show very similar values and percent
300 genetic gain.

301 In Study 1 when data were not corrected for maturity and wet and dry season data were combined, the
302 yield increase was $20.23 \text{ ha}^{-1} \text{ year}^{-1} \pm 4.68 \text{ kg ha}^{-1}$ with a genetic gain of 0.41% per year. However, in Study
303 2, the yield increase was $17.35 \text{ ha}^{-1} \text{ year}^{-1} \pm 3.32 \text{ kg ha}^{-1}$ with a genetic gain of 0.55% per year. The yield
304 increases in both studies show very similar values and the percent genetic gains were also similar when
305 wet and dry seasons' data were combined.

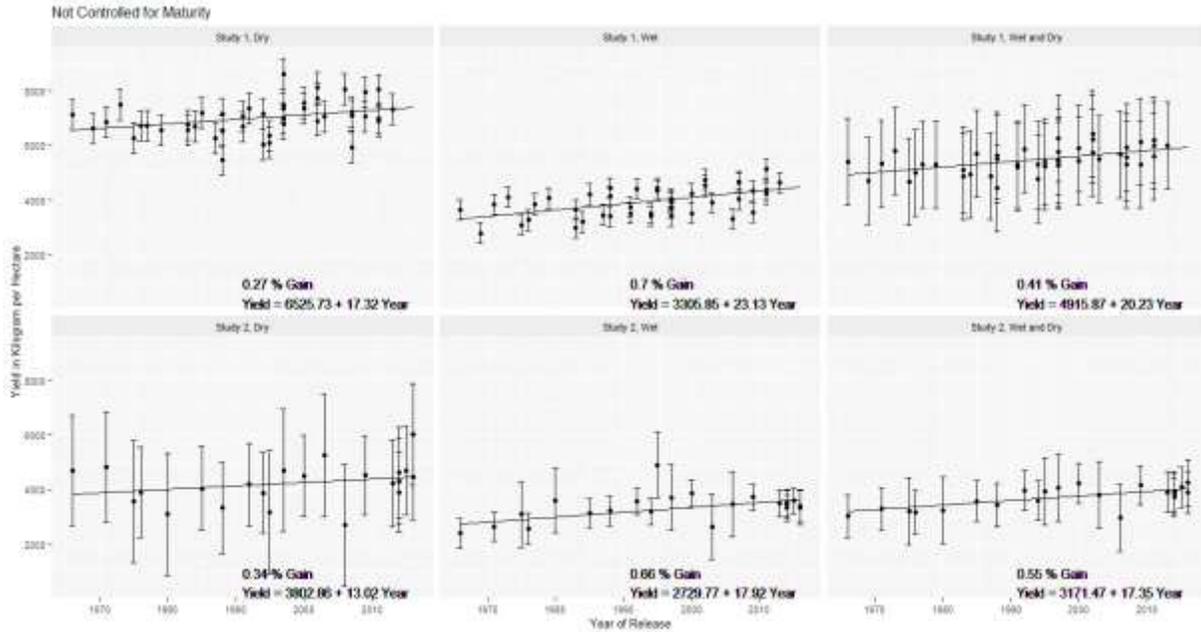
306 In Study 1 during the when data were corrected for maturity, the yield increase was $16.49 \text{ ha}^{-1} \text{ year}^{-1} \pm 7.3$
307 kg ha^{-1} with an annual genetic gain of 0.25%, which was slightly lower than when the data were not
308 corrected for maturity. In Study 2 during when data were corrected for maturity, the yield increase was
309 $14.79 \text{ ha}^{-1} \text{ year}^{-1} \pm 10.59 \text{ kg ha}^{-1}$ with a genetic gain of 0.38%. This was very similar to when the data were
310 not corrected for maturity. In both studies, correcting for maturity did not seem to make much difference
311 in the yield increases or percentage of genetic gain per year during the dry seasons.

312 In Study 1 during the wet season, when the data were corrected for maturity, the yield increases was
313 $20.81 \text{ ha}^{-1} \text{ year}^{-1} \pm 5.63 \text{ kg ha}^{-1}$ with a genetic gain of 0.62%. These values were slightly lower compared to
314 when the data were not corrected for maturity. In Study 2 during the wet season, when data were
315 corrected for maturity, the yield increase was $19.4 \text{ ha}^{-1} \text{ year}^{-1} \pm 4.09 \text{ kg ha}^{-1}$ with a genetic gain of 0.74%.
316 These values were similar to when the data were not corrected for maturity. Correcting for maturity once
317 again did not seem to make much difference in either the yield increases or percentage of genetic gain.

318 In Study 1 when data were corrected for maturity and wet and dry season data were combined, the yield
319 increase per year per hectare was $18.51 \text{ ha}^{-1} \text{ year}^{-1} \pm 4.89 \text{ kg ha}^{-1}$ with a genetic gain of 0.37% per year.

320 However, in Study 2 when wet and dry seasons data were combined and the data were corrected for
 321 maturity, the yield increase was $19.35 \text{ ha}^{-1} \text{ year}^{-1} \pm 4.26 \text{ kg ha}^{-1}$ with a genetic gain of 0.61% per
 322 year. Correcting for maturity did not make much difference for genetic gain.

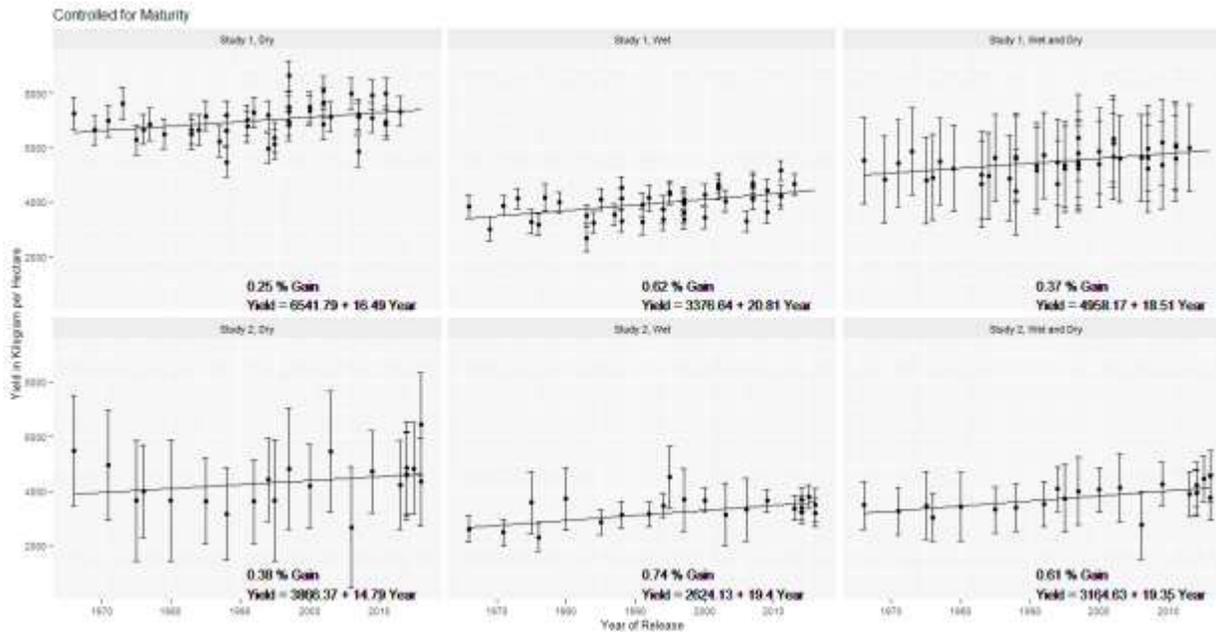
323 **Figure 1. Genetic trends in IRRI variety yields in the Philippines, when not controlled for maturity.**



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326 **Figure 2. Genetic trends in IRRI variety yields in the Philippines when controlled for maturity**



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328 The Figures 1 and 2 show that there is an increasing genetic gain trend among IRRI varieties over a 50 year
329 period. The base line grain yield, as indicated by the regression constant, is higher for Study 1 compared
330 for Study 2, in dry, wet and combined season. In Study 1, only data from one location were used, while in
331 Study 2, data from both the “Era” trials and the MET with seven locations were used.

332 For Study 1, increase in yield as indicated by the slope of the regression lines are higher when grain yield
333 was not corrected for days to maturity while in Study 2 it was observed that the increase in yield was
334 higher when yield was corrected for days to maturity.

335 Likewise, percentages of genetic gain per year in Study 1 are higher when grain yield was not corrected
336 for days to maturity, while in Study 2, it was observed that percentage of genetic gain was higher when
337 grain yield was corrected for days to maturity.

338 In both studies, it was observed that increase in yield and percentage of genetic gain per year is higher in
339 wet season. This is true for grain yield both controlled and not controlled for days to maturity.

340 In Study 1, grain yield corrected and not corrected for days to maturity did not make a significant
341 difference in the analyzed data for base line yield, yield increase, and percentage of genetic gain per year.

342 In Study 2, grain yield corrected and not corrected for days to maturity did make a significant difference
343 in the analyzed data for yield increase and percentage of genetic gain, per year but not for base line yield.

344 Each point in Figures 1 and 2 corresponds to the mean yield of a variety measured in kg ha^{-1} at the Zeigler
345 Experiment Station in IRRI during dry and wet seasons combined (right), during the dry season alone (left),
346 and during the wet season alone (middle) in Study 1 (top) and Study 2 (bottom). The dotted lines show
347 the linear regression between variety mean yield and year of variety release. At the lower right corner of
348 each panel, the percent gain in annual yield and the linear regression equation are displayed.

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353 **Variety relationship**

354 Kinship matrices, which explain the relationship between and among varieties, were calculated to
355 understand if close relationship if any, between and among varieties could be a factor in contributing
356 either positively or negatively to the estimated genetic gain. Kinship heat maps of varieties used in the
357 studies were generated (Figures 3 and 4 for Studies 1 and 2, respectively) to display the relationships
358 visually. Kinship matrices for varieties in the two studies were done using pedigrees recorded for the
359 entries. The relationships among the varieties used were assessed and are shown in the kinship matrices
360 tables for Study 1 (Supplementary Table 6) and Study 2 (Supplementary Table 7). Coefficients of parentage
361 based on pedigrees for Studies 1 and 2 are in Supplementary Tables 8 and 9, respectively. Coefficient of
362 parents is a measure of germplasm diversity used in these studies.

363 In Study 1, the kinship matrix for popular/mega varieties showed that only IR36 seemed to have higher
364 levels of relationship values (>0.4) with 12 out of 44 varieties (27%). IR 64 had higher levels of relationship
365 (>0.4) although there were only with three other varieties. IR72 had higher levels of relationship (>0.4)
366 with 7 of 44 varieties used (Supplementary Table 6).

367 In Study 2, the kinship matrix showed that only IR64 seemed to have higher relationships (>0.4) with 6 of
368 22 varieties. IR36 and IR72 had only three and two varieties, respectively, with higher levels of relationship
369 (>0.4) (Supplementary Table 7). To some extent, this explains that IRRI varieties released so far do not
370 necessarily come from a very narrow genetic base pool of lines

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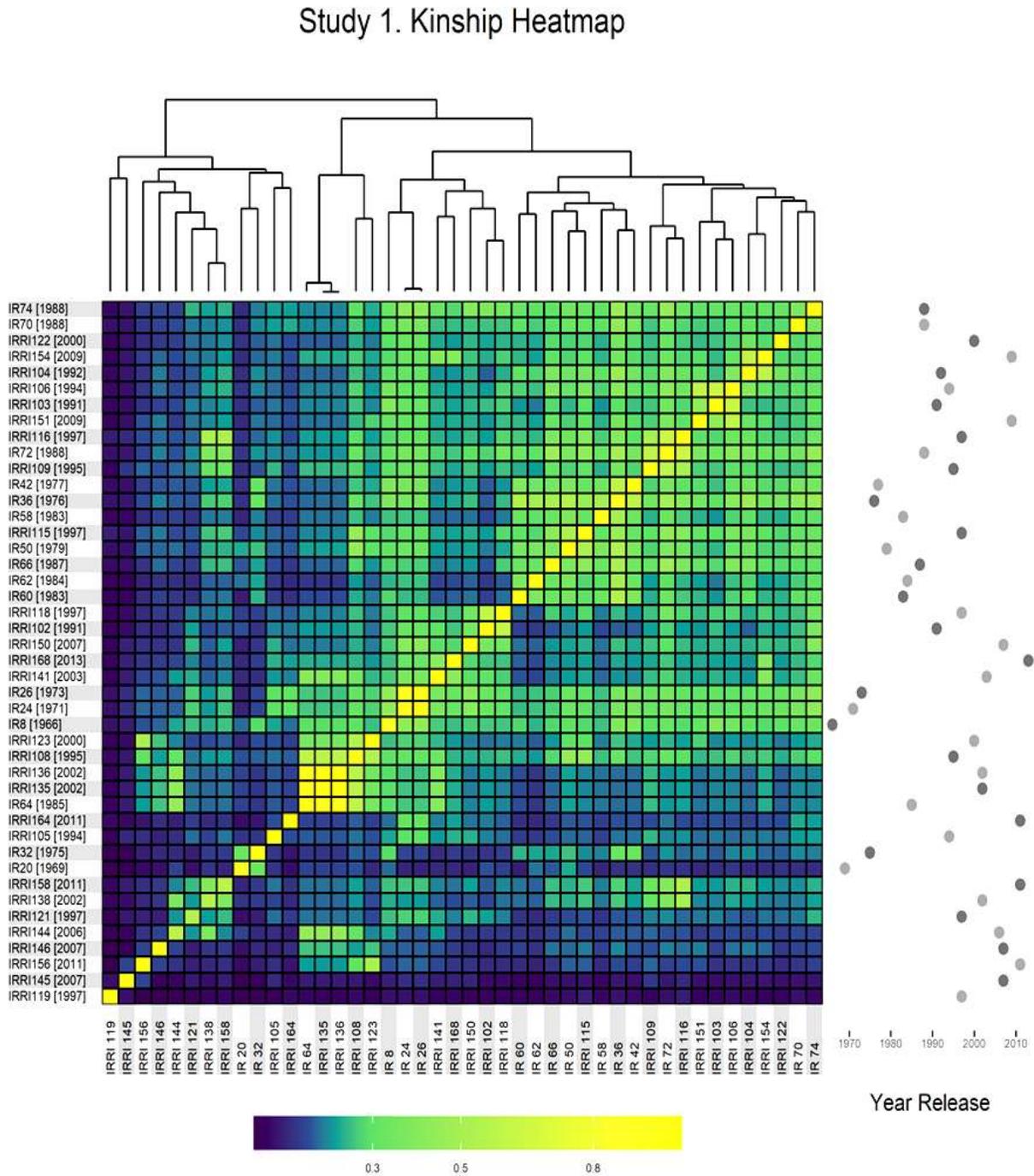
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377 **Figure 3. Kinship heat map of varieties used in Study 1**



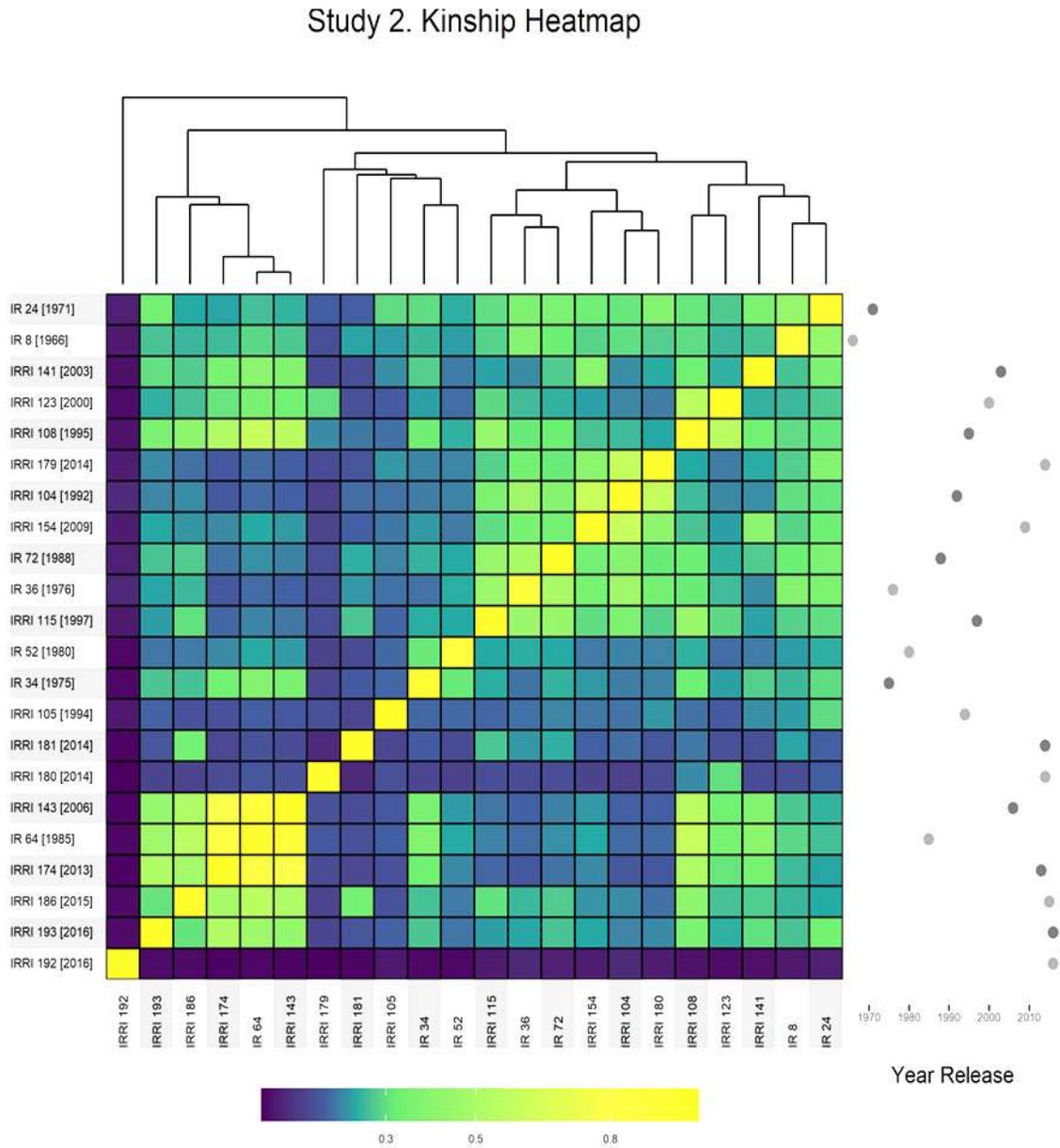
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382 **Figure 4. Kinship heat map of varieties used in Study 2**



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389 4. DISCUSSIONS and CONCLUSIONS

390 The data from the two studies (Table 3) clearly show similar levels of genetic trend among irrigated
391 varieties released at IRRI from 1966 to 2016, suggesting that they achieved greater than a zero rate of
392 genetic gain. In Study 1, the highest level of genetic gain per year was 0.70% in the wet season when not
393 corrected for maturity. In Study 2, the highest level of genetic gain was 0.74% in the wet season when
394 corrected for maturity. Yield increase was 23 kg ha⁻¹ year⁻¹ in Study 1 and 19 kg ha⁻¹ year⁻¹ in Study 2. The
395 lowest level of genetic gain in Study 1 was 0.25% in the dry season when data were corrected for maturity.
396 In Study 2, the lowest level of genetic gain was 0.34% in the dry season when data were not corrected for
397 maturity.

398

399 Comparison of rice with other staple crops, such as wheat, is of interest. The genetic gain from
400 both studies in the wet season was very similar to the 0.7% year⁻¹ gain in advanced spring bread
401 wheat (*Triticum aestivum* L.) lines evaluated in Mexico by CIMMYT from 1977 to 2008 (Lopes et
402 al., 2012) and from 1995 to 2009 with yield gains of 27 kg ha⁻¹ year⁻¹ at 0.65% genetic gain
403 (Sharma et al., 2012). This rate was also similar to yield increases in bread wheat (25 kg ha⁻¹ year⁻¹
404 ¹) in Australia measured from 1958 to 2007 (Sadras & Lawson, 2011) and yield increases of 35.1
405 kg ha⁻¹ year⁻¹ or 0.88% year⁻¹ in Spain from 1930 to 2000 (Sanchez-Garcia et al., 2013). This was
406 also comparable to yield increases in barley grown in the U.K with a genetic gain of 0.39%
407 between 1880 and 1980 (Riggs et al., 1981).

408 Graybosch & Peterson (2010) reported a genetic gain level of 1.1% in winter wheat on the Great Plains of
409 North America from 1959 to 2008, which was similar to that reported by Peng et al. (2000). From 50 years
410 of wheat breeding in Brazil, Rodrigues et al. (2007) assessed that the genetic gain for grain yield was 44.9
411 kg ha⁻¹ year⁻¹. However, in studies conducted by Brennan et al. (2004) to assess annual rate of genetic gain

412 in the wheat breeding program in Wagga Wagga, Australia, they assessed that the annual rate of gain was
413 3.19% while the annual rates of gain in northeastern and southeastern Australia were 2.75 and 3.92%,
414 respectively. In their studies, Brennan & Bialowas (2001) determined that 54% of the gain was due to
415 varieties and 46% to other factors.

416 In both studies, correction for maturity did not appear to make much difference for the analyzed data for
417 yield increases and percentages of annual genetic gain. Additional data from METs (Supplementary Table
418 1), while they added to the data points for increased robustness, they did not make much difference in
419 the genetic trend of one study over the other. Higher *per se* yield in both studies during the dry seasons
420 did not make any difference in the genetic gains seen in these varieties under both trials and years.
421 However, the estimated genetic gains in both studies were lower than the genetic gain of 1% estimated
422 for irrigated rice with yield increases ranging from 75 to 81 kg ha⁻¹ year⁻¹ in trials conducted by Peng et al.
423 (2000).

424 There are key differences in the genetic gain study conducted in 1996 by the Peng *et al* (2000) and the
425 two studies as described below.

426 Twice the number of varieties were used in Study 1 with 12 entries common in the two studies. Both
427 studies included all the major varieties released up to the years when they were conducted. However, the
428 later-released entries included in Study 2 did not seem to make any difference in genetic gains. Peng et
429 al. (2000) used nine IRRI-bred varieties and two breeding lines and one external breeding line spanning
430 29 years totaling up to 12 entries ((1966–1995). As previously mentioned, Study 1 included 37 IRRI-
431 developed varieties, two rainfed varieties, and five hybrid varieties spanning 47 years (1966–2013)
432 totaling up to 44 entries. There were eight varieties common to between Study 1 and Peng et al. (2000)
433 study. Study 2 had 21 irrigated varieties and one rainfed variety in the “Era” studies spanning 50 years
434 (1966–2016). There were four entries were common between the Study 2 and Peng et al. (2000) study.

435 Interestingly, the kinship matrices (Figures 3 and 4 and Supplementary Tables 6 and 7) displayed
436 relatedness among the varieties based on their pedigrees. It was important to understand if the lines
437 chosen were highly related to each other and/or to the mega-varieties, such as IR36, IR64, and IR72, based
438 on the pedigree, which may have contributed negatively or positively to the genetic gain observed or if
439 the genetic base was too narrow. Among the 44 varieties in Study 1, only IR36 showed similarity to 12
440 varieties with a kinship value greater than 0.4. In Study 2, among the 22 varieties, only IR36 was similar to
441 three varieties with a value greater than or equal to 0.4. In Study 1, IR64 was only similar to three varieties
442 with a kinship value of 0.4. In Study 2, IR64 was similar to six varieties with kinship values based on
443 pedigrees greater than 0.4. The Peng et al. (2000) study also used IR36 and, although the kinship matrix
444 values could not be developed directly as there are no data on the exact seedlot ID for that year in the
445 IRRI databases. However, based on Study 1 data, only three varieties (IR50, IR60, and IR 72) seem to have
446 a higher level of similarity to IR36 with values greater than 0.4 in the Peng et al. (2000) study. Coefficient
447 of parents based on the pedigrees of the germplasm used in the studies also showed diverse germplasm.
448 This analysis showed that the varieties used in the two studies did not come from a very narrow gene
449 pool. Perhaps for future genetic gain assessment studies, researchers should explore the critical minimum
450 number of varieties that should be included.

451 In general, *per se* yields were higher in both dry and wet seasons for Study 1, compared to Study 2 where
452 yields were lower in general. In Study 1 there was data only from one location. In Study 2, in which data
453 from both “Era” and MET trials in the Philippines were utilized, and due to environmental differences,
454 average yield would perhaps be lower than the single-site data in Study 1. In Peng et al. (2000) study,
455 yields presented in that paper were overall higher than both studies in this paper. These *per se* yield
456 differences perhaps could be attributed to year-to year variation in yield during conduct of the studies. In
457 the future, it would be worth recording weather data at all locations and use as a co-variate to assess
458 year-to-year variation impact on estimating genetic gain, which perhaps would also depend on stability of

459 yield across the years and across environments—especially under climate change scenarios. This will also
460 help to assess the impact of climate change on varietal performance and to develop strategies in breeding
461 programs to mitigate effects of climate change.

462 Interestingly, in both studies, the highest values for annual genetic gain were seen in the wet season
463 rather than the dry season, although *per se* yield in the dry season, as expected, was higher than in the
464 wet seasons. Annual genetic gains were similar in the dry seasons in both studies irrespective of maturity
465 correction. Evaluation during the wet season was not performed in the “Era” experiment conducted in
466 the Peng et al. (2000) although the objective stated in that paper was to determine the yield trend of rice
467 cultivars/lines developed since 1966. In the two studies presented here, evaluations were conducted both
468 in the dry and wet seasons to understand the genetic gain of varieties from a production standpoint as,
469 in Southeast Asia, more rice is grown during the wet seasons.

470 Higher levels of genetic gain for yield seen in the wet season could also be due to enhanced differences
471 among some varieties with improved pest and disease tolerances providing better yields compared to
472 varieties that are not so tolerant of pests and diseases. Therefore, in newly developed and released
473 varieties, genetic gain for yield must be measured in both seasons to demonstrate the yield potential of a
474 variety predominantly during the dry season and to assess genetic gain for yield when varieties have to
475 withstand higher pest and disease pressures during wet seasonal conditions. This will provide information
476 on average performance of a variety under different seasonal conditions that are more relevant to
477 practical variety release purposes. In the future, it would be worth utilizing pest and diseases data as a
478 covariates to assess year-to-year variation impact on estimating genetic gain across the years and across
479 environments, especially under climate change scenarios. Peng et al. (2010) also emphasized maintenance
480 breeding.

481 It should be noted that “Era” trial method has its own limitations. While Study 1 included a larger set of
482 varieties evaluated in one location only, due resource limitations, Study 2 included fewer varieties but

483 with data from more locations albeit in the Philippines only. Maturity was one covariate used in this study
484 however, correcting for maturities did not seem to make significant differences either increase in yield
485 per tons per hectare per year or percent genetic gain. Ideally, such studies need to be conducted across
486 environments and across years with commonality of some varieties used between years to increase
487 connectedness among varieties between the years. For all practical purposes, conducting such “Era” trials
488 across key environments in numerous rice-producing countries would be resource-intensive. Rutkoski
489 (2019), in her studies compared various methods for assessing genetic gain, stated that utilization of
490 estimated breeding values (EBVs) by including long- term check varieties as the method which gave
491 second high level of precision and lowest level of error among all the methods she compared. However,
492 she noted that, for the EBV method to work well, EBV estimation required accurate records of pedigrees
493 from the time that the breeding program began to estimate additive relationships among lines and having
494 good connectivity between years. Rutkoski (2019) also concluded that, when a breeding program does
495 not meet the requisites for EBV estimations, the “Era” method can be used instead because data are not
496 needed from historical datasets. The EBV experiments can be designed to ensure achieving precision,
497 efficiency, correlation and reducing error although they require the availability of varieties released over
498 the years and resources to conduct new trials. She also recommended that, to improve efficiency, several
499 varieties released annually could be used instead of just one variety. In Studies 1 and 2, an effort was
500 made to source more than one variety per year, although for the older varieties, it was difficult to find
501 authentic sources of seed for older varieties in addition to requiring larger and more resource-intensive
502 trials.

503 One limitation in both the Peng et al. (2000) study and Studies 1 and 2 was their preliminary nature since
504 they did not have large sets of historical data like those included in other crops that have utilized multiple
505 locations spanning several decades. Ideally, yield trials should always be conducted in multiple
506 environments and years and have genotype x location (GxL) interactions. Also, annual variations need to

507 be accounted for including the variety's stability of performance. The Peng et al. (2000) study conducted
508 trials in two locations—IRRI, Los Baños and PhilRice, Muñoz). The data were pooled from the trials
509 conducted in Los Baños and Munoz. In Study 1, the "Era" trials were only conducted at IRRI, Los Baños
510 since trials could not be conducted at multiple locations due to insufficient resources. In Study 2, "Era"
511 trials were conducted only in IRRI, Los Baños, but additional data from other MET locations
512 (Supplementary Table 1) in the Philippines for varieties common to "Era" and MET trials were included in
513 the analyses to increase the robustness of analyses. It should also be noted that the genotype x year (GxY)
514 interaction is often as large or larger than the genotype x location (GxL) within a reasonably delineated
515 target population of environments (TPEs) (Cooper et al., 1999; Atlin et al., 2000).

516 Conducting trials across TPEs across years for assessment of genetic gain of varieties, although necessary,
517 is both resource and labor intensive, especially where research funds are limited. To overcome such
518 resource limitations for testing advanced lines under different environments in different rice producing
519 countries and thereby develop data on agronomic adaptation, pest and disease tolerances and selections
520 for locally desired traits, IRRI developed a network system named International Network for Genetic
521 Evaluation of Rice (INGER) for sharing and evaluating advanced lines in among rice-producing countries.
522 INGER has enabled national country breeders to further select the best performers for agronomic
523 adaptation and yield performance over several years and under the national systems' varietal release
524 protocols. This network also allows national country partners to utilize best performers and develop
525 crosses for selections in the local environments. Data from INGER trials could be used for estimations of
526 genetic gains going forward with larger historical datasets to overcome limitations of conducting new
527 large trials spanning multi locations. It should be noted that INGER network system has led to over 1,200
528 varieties being released and that popular mega varieties, such as IR36, IR64, and IR72 (Mackill & Khush,
529 2018; Khush, 1981) have also been released to farmers in the partner countries and high performers
530 among these would also provide connectedness for the data. Evenson and Gollin (1997) conducted

531 economic analyses and calculated that the average value of the varieties released through INGER to be
532 2.5 million USD variety⁻¹ year⁻¹ (in 1997 dollars). In their analysis, Brennan & Malabayabas (2011) reported
533 that such network-based sharing contributed to yield improvement in Southeast Asian countries. This
534 type of sharing and testing of advanced lines has significantly impacted on yield improvements in rice
535 production environments as shown by these economic estimates. Improved yields would also accomplish
536 genetic gain of varieties in the countries and for the CGIAR and IRRI, alignment to food security goals as
537 part of the mission. Sharing germplasm to a wide array of countries with and without CGIAR/IRRI rice
538 breeding activities, also increases diversity of the improved lines in national breeding programs thereby
539 increasing the elite pools of germplasm in partner countries and ensures that no rice-producing country
540 is left behind (Khush & Virk, 2005).

541 A key limitation of such a system of sharing advanced breeding lines or pre-variety lines is that the testing
542 and selections of early-stage IRRI breeding lines would not be done in the countries, thereby missing out
543 on selections for some key traits during early stages of breeding. Therefore, it is desirable that breeding
544 programs develop early-stage breeding regional networks at least in some key countries and select for
545 desired traits suited to rice production areas, achieve a high probability of achieving genetic gain on an
546 annual basis. However, development of an early-stage breeding regional network would require
547 cooperation from all partners in the National Agricultural Research and Extension System (NARES)
548 countries and would be resource-intensive. Several NARES partner countries also have their own national
549 rice breeding programs where IRRI-shared germplasm is utilized both for direct variety releases and
550 crosses therefore, a system of co-operative-early-stage regional breeding by sharing early stage
551 germplasm at least in some regional targeted countries would be desirable to support goals of higher
552 productivity through genetic gain. These early-stage selections from the regional breeding networks could
553 flow into advanced line/pre-variety networks such as INGER and expand the testing in a wider number of

554 countries and provide data for genetic gain assessments using historical sets of data from all INGER
555 partners and measure the effectiveness of breeding.

556 While different methods can be adopted to estimate genetic gain, some key factors that contribute to
557 genetic gain need to be noted. A key factor in yield improvement in crops such as maize has been the
558 coupling of improved management practices with variety improvement. Duvick (2005b), in an assessment
559 of genetic gain in hybrid maize, found that 50–60% of the on-farm yield gains were due to genetic
560 improvement of varieties while about 40–50% were due to changes in management responsible for the
561 improvements in agronomic practices. In rice grown in the United States from 1981 to 2011, yields
562 increased 47% (2.5 t ha^{-1}) at an average of 86 kg ha^{-1} annually. These yield gains have occurred due to
563 improved traits such as a semidwarf habit, disease resistance, specific cultivars, herbicide tolerance, and
564 F_1 hybrids (Moldenhauer et al., 2013). In Australian wheat, Brennan & Bialowas (2001) found that, on
565 average, 54% of the genetic gain was due to yield improvement alone, excluding external factors other
566 than varieties

567 While in countries such as the U.S., Australia, and Brazil, wheat or maize mechanized cropping
568 establishment and harvest systems have contributed to 40% of the yield improvement as mentioned
569 above, in Southeast Asia, application of fertilizers, such as nitrogen for rice crop establishment, and
570 harvesting generally, are not mechanized at the farmer level (Okasha & Hegazy 2020). Even among
571 breeding programs, mechanization is either minimal or lacking completely for crop establishment.
572 Evaluations of breeding lines and varieties under experimental plot trials established using variable,
573 manual crop establishment methods, manual methods of fertilizer applications and harvests conducted
574 under inconsistent moisture conditions could contribute to lower genetic gain in rice.

575 These studies, while recognizing that there has been genetic gain in IRRI's irrigated rice breeding program,
576 have clearly demonstrated that there is a need for further improvement of genetic gain for yield in rice
577 because OECD/FAO (2019) has estimated a need for a 40 million ton increase in rice required by 2028. A

578 higher level of genetic gain for yield is critical along with traits for mitigating effects of climate change
579 (e.g., flood and drought tolerances) and managing abiotic stress in soils. Studies have shown that, for
580 every 1-degree Celsius increase in temperature, there is a yield reduction of up to 10% with a reduction
581 in economic growth by 1.3% (Wassmann & Jagadish, 2014). Interestingly, the Peng study et al. (2000)
582 recorded much higher yields in 1998 compared to Study 1 (2013–2014) and Study 2 (2016–2017). While
583 there is also a case for improvements in agronomic practices without much need for additional inputs
584 from farmers, there is an important need for breeding programs to develop breeding strategies with
585 available resources to improve yield and other desired traits to increase realized genetic gains for yield
586 and other consumer-desired traits.

587 A major difficulty for rice is that, unlike other crops such as maize and wheat, rice is consumed directly as
588 a grain and so consumer and value chain stakeholder preferences and market demand must also have
589 consideration beyond genetic gain for yield when selecting new varieties. In this regard, studies such as
590 those by Laborte et al. (2015) would assist in assessment of consumer needs in rice producing countries.
591 Therefore, while it is important to have genetic gain for yield in rice varieties, it is also important to select
592 for a combination of multiple traits required for not only yield, but climate resilience and quality traits as
593 well. This simultaneous trait selections would help in improving the performance of varieties on the farm,
594 increasing adoption by farmers of the improved varieties, and ensuring marketability due to alignment
595 with consumer desires.

596 The studies in this paper do not focus on recommending an ideal breeding method to increase yield
597 genetic gain in rice as comparison was not done using varieties developed by different breeding methods.
598 The varieties used in these studies were developed using different methods over a long period defined as
599 an “Era”. Studies comparing different breeding method generated varieties would require using rice
600 varieties in multi environments across multiple years in addition to being resource-intensive.

601 Utilization of available molecular tools for selection of diverse improved parents and development of a
602 new generation of crosses will lead to a better certainty of the presence of desired genes and prediction
603 of performance of those crosses. Implementation of single-seed decent methods, either in the field or in
604 the glasshouse, will reduce costs of a breeding program while also improving the program's speed and
605 agility. Heffner et al. (2010), in their studies comparing marker-assisted selections (MAS) and genomic
606 selections (GS) in wheat and maize, indicated that, even at lower genomic estimated breeding values
607 (GEBVs) for complex traits, the expected annual gain from genomic selections in maize was three times
608 that from MAS and in winter wheat twice that from MAS therefore genomic selections can accelerate
609 genetic gain due to shorter breeding cycles. Voss-Fels et al. (2019) in their paper on accelerating genetic
610 gain using genomic selection stated that, genomic selection in crops offers a way to reduce breeding
611 cycles, especially when combined with molecular tools, and rapid generation advancement. This is quite
612 specific to crops and breeding programs and needs continued research. Apart from reduction in breeding
613 cycles, plant breeding programs can fast-track the outputs of the varieties by utilizing methods such as
614 selection index and utilize economic weights for selections of parents (Batista et al., 2018). Details of
615 various aspects of developing a selection index for plant breeding have been described by Céron-Rojas &
616 Crossa (2018).

617 For all practical purposes, studies such as 'Era' studies or analysis of historical data sets need to be
618 conducted at regular intervals utilizing best practices to assess genetic gain of the new varieties using in
619 multi-location trials under climate change scenarios in the major rice-producing environments. This will
620 enable breeding programs to develop strategies for release of enriched varieties that are used by the
621 farmers for better adoption and increased profitability. Assessments need to be done to estimate the
622 required target population of environments for early breeding line selections for yield, pest and disease
623 resistances and tolerances. Consumer-desired grain quality traits under multi environment trials of
624 advanced lines must also be assessed. Simulated studies may provide genetic gain to be targeted by

625 breeding programs to develop crossing plans and strategies to ensure target is reached with necessary
626 population structures in the breeding programs. Assessment of realized genetic gain at the farm level will
627 also provide a critical feedback loop that breeders will require to make necessary improvements for faster
628 breeding cycles and catering to changing markets and demands. To that end, it is critical that necessary
629 investments be made for both early breeding line evaluations in Target Population of Environments (TPEs)
630 as well as advanced breeding lines in the TPEs. Since IRRI itself does not release varieties directly but
631 instead nominates varieties to the NARES partners in the national variety-release systems, it should be
632 emphasized that research collaboration on a sustained basis will require continual funding and longer-
633 term engagement to meaningfully achieve genetic gain in farmers' fields.

634 **Abbreviations in alphabetical order**

635 CIMMYT: International Maize and Wheat Improvement Center

636 DS: Dry seasons

637 EBVs: Estimated breeding values

638 FAO: Food and Agriculture Organization

639 FLW: Flowering time

640 GEBV: Genomic estimated breeding value

641 GS: Genomic selections

642 IRRI: International Rice Research Institute

643 INGER: International Network for Genetic Evaluation of Rice

644 MAS: Marker-assisted selections

645 MET: Multi Environment Trials

646 OECD: Organization for Economic Cooperation and Development

647 PH: plant height

648 SES: Standard evaluation system

649 TN: Number of productive tillers
650 TPE: Target Population of Environments
651 WS: Wet seasons
652 RCBD: Randomized Complete Block Design

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812 **Declarations**

813 **Ethics approval and consent to participate**

814 Not Applicable

815 **Consent for Publication**

816 Not applicable

817 **Availability of Data and Material**

818 All datasets supporting the conclusions of this article are included in this article (and its additional
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821 **Competing interests**

822 The authors declare that they have no competing interests

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831 This study was undertaken as a research study designed and executed by the authors.

832 **Authors’ contribution**

833 SV is the corresponding author. SV conceived and designed the experiments and organized all analysis for
834 Studies 1 and 2. BCYC conceived and designed the experiments and provided the data for analyses for
835 Study 1. SV contributed to majority of the writing of the article, submission for internal reviews, revisions

836 and interpretation of the data and preparation of tables and figures for manuscripts. SV was assisted by
837 AG for all the statistical analyses narrative writing and development of figures and tables. BCYC assisted
838 with the writing, revisions and provided all data for the study 1. AG conducted all data analyses for studies
839 1 and 2. SV also organized and edited and revised the manuscript for submission. SV for study 2 fully
840 organized the conduct of the work in the field, collection, collation of data and assisted by AP. BCYC,
841 designed the study 1, contributed to the work in the field, organized the collection, collation of data for
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Supplementary Files

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- [SupplementaryTable4.Study2LSmeansSEvarieties.xlsx](#)
- [SupplementaryTable5.Study2trialHeritabilities.xlsx](#)
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