Chapter 6 Constrained Linear Genomic Selection Indices



Abstract The constrained linear genomic selection indices are null restricted and predetermined proportional gain linear genomic selection indices (RLGSI and PPG-LGSI respectively), which are a linear combination of genomic estimated breeding values (GEBVs) to predict the net genetic merit. They are the results of a direct application of the restricted and the predetermined proportional gain linear phenotypic selection index theory to the genomic selection context. The RLGSI can be extended to a combined RLGSI (CRLGSI) and the PPG-LGSI can be extended to a combined PPG-LGSI (CPPG-LGSI); the latter indices use phenotypic and GEBV information jointly in the prediction of net genetic merit. The main difference between the RLGSI and PPG-LGSI with respect to the CRLGSI and the CPPG-LGSI is that although the RLGSI and PPG-LGSI are useful in a testing population where there is only marker information, the CRLGSI and CPPG-LGSI can be used only in training populations when there are joint phenotypic and marker information. The RLGSI and CRLGSI allow restrictions equal to zero to be imposed on the expected genetic advance of some traits, whereas the PPG-LGSI and CPPG-LGSI allow predetermined proportional restriction values to be imposed on the expected trait genetic gains to make some traits change their mean values based on a predetermined level. We describe the foregoing four indices and we validated their theoretical results using real and simulated data.

6.1 The Restricted Linear Genomic Selection Index

Let $H = \mathbf{w'g}$ be the net genetic merit and $I_G = \mathbf{\beta'\gamma}$ the linear genomic selection index (LGSI, see Chap. 5 for details), where $\mathbf{g}, \mathbf{\gamma}, \mathbf{w}$, and $\mathbf{\beta}$ are vectors $t \times 1$ (t= number of traits) of breeding values, genomic breeding values, economic weights, and LGSI coefficients respectively. It can be shown that $Cov(I_G, \mathbf{g}) = \mathbf{\Gamma}\mathbf{\beta}$ is the covariance between \mathbf{g} and $I_G = \mathbf{\beta'\gamma}$, and that $Var(\mathbf{\gamma}) = \mathbf{\Gamma}$ is the genomic covariance matrix of size $t \times t$ (see Chap. 5 for details). The objective of the restricted linear genomic selection index (RLGSI) is to improve only (t - r) of t (r < t) traits (leaving r of them fixed) in a testing population using only genomic estimated breeding values (GEBVs). The RLGSI minimizes the mean squared difference between I_G and H, $E[(H - I_G)^2]$, with respect to $\boldsymbol{\beta}$ under the restriction $Cov(I_G, \mathbf{U}'\mathbf{g}) = \mathbf{U}'\boldsymbol{\Gamma}\boldsymbol{\beta} = \mathbf{0}$, where \mathbf{U}' is a matrix $(t - 1) \times t$ of 1s and 0s, in a similar manner to the restricted linear phenotypic selection index (RLPSI) described in Chap. 3 in the phenotypic selection context.

6.1.1 The Maximized RLGSI Parameters

Let $Var(I_G) = \beta' \Gamma \beta$ be the variance of $I_G = \beta' \gamma$, w'Cw the variance of H = w'g, and $Cov(I_G, H) = w'\Gamma\beta$ the covariance between H = w'g and $I_G = \beta'\gamma$. The mean squared difference between H and I_G can be written as $E[(H - I_G)^2]$, which should be minimized under the restriction $U'\Gamma\beta = 0$ assuming that Γ , C, U', and w are known, i.e., it is necessary to minimize the function

$$f_{R}(\boldsymbol{\beta}, \mathbf{v}) = \mathbf{w}' \mathbf{C} \mathbf{w} + \boldsymbol{\beta}' \boldsymbol{\Gamma} \boldsymbol{\beta} - 2 \mathbf{w}' \boldsymbol{\Gamma} \boldsymbol{\beta} + 2 \mathbf{v}' \mathbf{U}' \boldsymbol{\Gamma} \boldsymbol{\beta}$$
(6.1)

with respect to vectors $\boldsymbol{\beta}$ and $\mathbf{v}' = [v_1 \ v_2 \ \cdots \ v_{r-1}]$, where \mathbf{v} is a vector of Lagrange multipliers. In matrix notation, the derivative results of Eq. (6.1) are

$$\begin{bmatrix} \boldsymbol{\beta} \\ \mathbf{v} \end{bmatrix} = \begin{bmatrix} \boldsymbol{\Gamma} & \boldsymbol{\Gamma} \mathbf{U} \\ \mathbf{U}' \boldsymbol{\Gamma} & \mathbf{0} \end{bmatrix}^{-1} \begin{bmatrix} \boldsymbol{\Gamma} \mathbf{w} \\ \mathbf{0} \end{bmatrix}.$$
(6.2)

Following the procedure described in Chap. 3 (Eqs. 3.2 to 3.5), it can be shown that the RLGSI vector of coefficients that minimizes $E[(H - I_G)^2]$ under the restriction U' $\Gamma\beta = 0$ is

$$\boldsymbol{\beta}_{RG} = \mathbf{K}_G \mathbf{w},\tag{6.3}$$

where $\mathbf{K}_G = [\mathbf{I}_t - \mathbf{Q}_G]$, $\mathbf{Q}_G = \mathbf{U}(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{U}'\Gamma$, w is a vector of economic weights, and \mathbf{I}_t is an identity matrix $t \times t$. When no restrictions are imposed on any of the traits, \mathbf{U}' is a null matrix and $\boldsymbol{\beta}_{RG} = \mathbf{w}$, the optimized LGSI vector of coefficients (see Chap. 5 for details).

By Eq. (6.3), the RLGSI, and the maximized RLGSI selection response and expected genetic gain per trait can be written as

$$I_{\rm RG} = \mathbf{\beta}_{RG}' \mathbf{\gamma},\tag{6.4}$$

$$R_{RG} = \frac{k_I}{L_G} \sqrt{\beta_{RG}' \Gamma \beta_{RG}}$$
(6.5)

and

$$\mathbf{E}_{RG} = \frac{k_I}{L_G} \frac{\Gamma \boldsymbol{\beta}_{RG}}{\sqrt{\boldsymbol{\beta}_{RG}' \Gamma \boldsymbol{\beta}_{RG}}},\tag{6.6}$$

respectively, where k_I is the standardized selection differential (or selection intensity) associated with the RLGSI, and L_G is the interval between selection cycles or the time required to complete a selection cycle using the RLGSI. Equations (6.4) to (6.6) depend only on GEBV information; thus, they are useful in testing populations.

6.1.2 Statistical Properties of RLGSI

Assuming that $H = \mathbf{w}'\mathbf{g}$ and $I_{RG} = \mathbf{\beta}'_{RG}\mathbf{\gamma}$ have bivariate joint normal distribution, $\mathbf{\beta}_{RG} = \mathbf{K}_{G}\mathbf{w}$, and $\mathbf{\Gamma}$, \mathbf{C} , and \mathbf{w} are known, it can be shown that the RLGSI has the following properties:

- 1. Matrices \mathbf{K}_G and \mathbf{Q}_G are idempotent ($\mathbf{K}_G = \mathbf{K}_G^2$ and $\mathbf{Q}_G = \mathbf{Q}_G^2$) and orthogonal ($\mathbf{K}_G \mathbf{Q}_G = \mathbf{Q}_G \mathbf{K}_G = \mathbf{0}$), that is, they are projectors. Matrix \mathbf{Q}_G projects vector $\boldsymbol{\beta} = \mathbf{w}$ into a space generated by the columns of matrix $\mathbf{U}'\Gamma$ due to the restriction $\mathbf{U}'\Gamma\boldsymbol{\beta} = \mathbf{0}$ used when $f_R(\boldsymbol{\beta}, \mathbf{v})$ (Eq. 6.1) is minimized with respect to vectors $\boldsymbol{\beta}$ and \mathbf{v} , whereas matrix \mathbf{K}_G projects \mathbf{w} into a space perpendicular to that generated by the $\mathbf{U}'\Gamma$ matrix columns.
- 2. Because of the restriction $\mathbf{U}' \mathbf{\Gamma} \boldsymbol{\beta} = \mathbf{0}$, matrix \mathbf{K}_G projects vector \mathbf{w} into a space smaller than the original space of \mathbf{w} . The space reduction into which matrix \mathbf{K}_G projects \mathbf{w} is equal to the number of zeros that appears in Eq. (6.6).
- 3. Vector $\beta_{RG} = \mathbf{K}_{G} \mathbf{w}$ minimizes the mean square error under the restriction $\mathbf{U}' \mathbf{\Gamma} \boldsymbol{\beta} = \mathbf{0}$.
- 4. The variance of $I_{RG} = \beta'_{RG} \gamma (\sigma^2_{I_{RG}} = \beta'_{RG} \Gamma \beta_{RG})$ is equal to the covariance between $I_{RG} = \beta'_{RG} \gamma$ and $H = w' g (\sigma_{HI_{RG}} = w' \Gamma \beta_{RG})$.
- 5. The maximized correlation between *H* and *I_{RG}* is equal to $\rho_{HI_{RG}} = \frac{\sigma_{I_{RG}}}{\sigma_H}$, where $\sigma_{I_{RG}} = \sqrt{\beta'_{RG}\Gamma\beta_{RG}}$ and $\sigma_H = \sqrt{\mathbf{w'Cw}}$ are the standard deviations of $I_{RG} = \beta'_{RG}\gamma$ and $H = \mathbf{w'g}$ respectively.
- 6. The variance of the predicted error, $Var(H I_{RG}) = (1 \rho_{HI_{RG}}^2)\sigma_H^2$, is minimal. Note that $Var(H - I_{RG}) = \sigma_{I_{RG}}^2 + \sigma_H^2 - 2\sigma_{HI_{RG}}$, and when $\boldsymbol{\beta}_{RG} = \mathbf{K}_G \mathbf{w}$, $\sigma_{I_{RG}}^2 = \sigma_{HI_{RG}}$, whence $Var(H - I_{RG}) = \sigma_H^2 - \sigma_{I_{RG}}^2 = (1 - \rho_{HI_{RG}}^2)\sigma_H^2$ is minimal.

The statistical RLGSI properties are equal to the statistical RLPSI properties. Thus the RLGSI is an application of the RLPSI to the genomic selection context.

6.1.3 Numerical Examples

To estimate the parameters associated with the RLGSI, we use the real data set described in Chap. 5, Sect. 5.1.8, where we found that, in the testing population, the

estimate of matrix
$$\mathbf{\Gamma}$$
 was $\widehat{\mathbf{\Gamma}} = \begin{bmatrix} 0.21 & 2.95 & 5.00\\ 2.95 & 42.41 & 71.11\\ 5.00 & 71.11 & 121.53 \end{bmatrix}$. We use this matrix and the

GEBVs associated with the traits grain yield (GY, ton ha^{-1}), ear height (EHT, cm), and plant height (PHT, cm) to illustrate the RLGSI theoretical results.

Suppose that on the RLGSI expected genetic gain per trait we impose one and two null restrictions using matrices $\mathbf{U}'_1 = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$ and $\mathbf{U}'_2 = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$ (see Chap. 3, Sect. 3.1.3, for details about matrix \mathbf{U}'). We need to estimate the RLGSI vector of coefficients ($\boldsymbol{\beta}_{RG} = \mathbf{K}_G \mathbf{w}$) as $\hat{\boldsymbol{\beta}}_{RG} = \hat{\mathbf{K}}_G \mathbf{w}$, where $\hat{\mathbf{K}}_G = \begin{bmatrix} \mathbf{I}_3 - \hat{\mathbf{Q}}_G \end{bmatrix}$ and $\hat{\mathbf{Q}}_G = \mathbf{U} (\mathbf{U}' \hat{\Gamma} \mathbf{U})^{-1} \mathbf{U}' \hat{\Gamma}$ are estimates of matrices $\mathbf{K}_G = [\mathbf{I}_3 - \mathbf{Q}_G]$ and $\mathbf{Q}_G = \mathbf{U} (\mathbf{U}' \Gamma \mathbf{U})^{-1} \mathbf{U}' \Gamma$ respectively, and \mathbf{I}_3 is an identity matrix 3 × 3. The estimated \mathbf{Q}_G

matrices for restrictions
$$\mathbf{U}_1' = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$$
 and $\mathbf{U}_2' = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$ were $\widehat{\mathbf{Q}}_{G_1} = \mathbf{U}_1 \left(\mathbf{U}_1' \widehat{\mathbf{\Gamma}} \mathbf{U}_1 \right)^{-1}$

$$\mathbf{U}_{1}'\widehat{\mathbf{\Gamma}} = \begin{bmatrix} 1.0 & 14.05 & 23.81 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \text{ and } \widehat{\mathbf{Q}}_{G_{2}} = \mathbf{U}_{2} (\mathbf{U}_{2}'\widehat{\mathbf{\Gamma}}\mathbf{U}_{2})^{-1} \mathbf{U}_{2}'\widehat{\mathbf{\Gamma}} = \begin{bmatrix} 1.0 & 0 & 11.18 \\ 0 & 1.0 & 0.90 \\ 0 & 0 & 0 \end{bmatrix} \text{ respectively}$$

tively, whereas the estimated \mathbf{K}_{G} matrices for both restrictions were $\widehat{\mathbf{K}}_{G_1} = [\mathbf{I}_3 - \widehat{\mathbf{Q}}_{G_1}]$

$$= \begin{bmatrix} 0 & -14.05 & -23.81 \\ 0 & 1.0 & 0 \\ 0 & 0 & 1.0 \end{bmatrix} \text{ and } \widehat{\mathbf{K}}_{G_2} = \begin{bmatrix} \mathbf{I}_3 - \widehat{\mathbf{Q}}_{G_2} \end{bmatrix} = \begin{bmatrix} 0 & 0 & -11.18 \\ 0 & 0 & -0.90 \\ 0 & 0 & 1.0 \end{bmatrix}$$

Let $\mathbf{w}' = \begin{bmatrix} 5 & -0.1 & -0.1 \end{bmatrix}$ be the vector of economic weights; then the estimated RLGSI vector of coefficients for one and two null restrictions were $\hat{\boldsymbol{\beta}}'_{RG_1} = \mathbf{w}' \hat{\mathbf{K}}'_{G_1} = \begin{bmatrix} 3.78 & -0.1 & -0.1 \end{bmatrix}$ and $\hat{\boldsymbol{\beta}}'_{RG_2} = \mathbf{w}' \hat{\mathbf{K}}'_{G_2} = \begin{bmatrix} 1.12 & 0.09 & -0.1 \end{bmatrix}$ respectively, and the estimated RLGSI for both restrictions can be written as $\hat{\boldsymbol{I}}_{RG_1} = 3.78 \text{ GEBV}_1 - 0.1$ GEBV₂ - 0.1GEBV₃ and $\hat{\boldsymbol{I}}_{RG_2} = 1.12 \text{ GEBV}_1 + 0.09 \text{ GEBV}_2 - 0.1 \text{ GEBV}_3$, where GEBV₁, GEBV₂, and GEBV₃ are the genomic estimated breeding values associated with traits GY, EHT, and PHT respectively in the testing population.

Table 6.1 presents 20 genotypes selected from a population of 380 genotypes and the GEBVs in the testing population ranked according to the estimated RLGSI values for one restriction, where $\mathbf{U}'_1 = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$. The estimated RLGSI values for genotypes 5 and 306 can be obtained as follows: $\hat{I}_{RG_5} = 3.78(-0.6) - 0.1(-8.67) - 0.1(15.97) = 0.196$ and $\hat{I}_{RG_{306}} = 3.78(0.13) - 0.1(1.31) - 0.1(1.66) = 0.194$ respectively. This procedure is valid for any number of genotypes and GEBVs in the testing population.

Assume a selection intensity of 10% ($k_{I_G} = 1.755$); then the estimated RLGSI selection response and expected genetic gain per trait not including the interval length were $\hat{R}_{RG_1} = k_{I_G} \sqrt{\hat{\beta}'_{RG_1} \hat{\Gamma} \hat{\beta}_{RG_1}} = 0.40$ and $\hat{\mathbf{E}}'_{RG_1} = k_I \frac{\hat{\beta}'_{RG_1} \hat{\Gamma}}{\sqrt{\hat{\beta}'_{RG_1} \hat{\Gamma} \hat{\beta}_{RG_1}}}$

Table 6.1 Number of genotypes selected from 380 genotypes of a real testing population; genomic estimated breeding values (GEBVs) associated with three traits: grain yield (GY, ton ha^{-1}), ear height (EHT, cm), and plant height (PHT, cm) in the testing population, and estimated and ranked restricted linear genomic selection index (RLGSI) values obtained in the testing population for one null restriction

	Estimated GE	BVs in the testing	g population	
Number of genotypes	GEBV-GY	GEBV-EHT	GEBV-PHT	Estimated RLGSI
5	-0.6	-8.67	-15.97	0.196
306	0.13	1.31	1.66	0.194
6	0.06	1.83	-1.13	0.157
349	0.37	4.34	8.12	0.153
142	-0.26	-5.47	-5.85	0.149
69	-0.11	-3.43	-2.16	0.143
24	0.03	-0.43	0.19	0.137
192	-0.8	-13.91	-17.7	0.137
33	-0.18	-1.44	-6.71	0.135
18	-0.43	-5.48	-12.08	0.131
21	-1.00	-16.11	-22.96	0.127
41	0.17	1.09	4.08	0.126
351	0.16	2.64	2.15	0.126
323	0.04	-0.79	1.04	0.126
158	-0.49	-8.95	-10.83	0.126
25	-0.24	-3.46	-6.86	0.125
338	0.37	3.88	8.89	0.122
316	-0.01	-0.51	-1.09	0.122
32	-0.19	-3.97	-4.43	0.122
204	-0.46	-7.41	-11.19	0.121

 $= \begin{bmatrix} 0 & -1.42 & -2.58 \end{bmatrix}$ respectively. For two restrictions, with $\mathbf{U}_2' = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$, the estimated RLGSI selection response and expected genetic gains not including the interval length were $\widehat{R}_{RG_2} = k_{I_G} \sqrt{\widehat{\beta}'_{RG_2} \widehat{\Gamma} \widehat{\beta}_{RG_2}} = 0.23$ and $\widehat{\mathbf{E}}'_{RG_2} = k_I \frac{\widehat{\beta}'_{RG_2} \widehat{\Gamma}}{\sqrt{\widehat{\beta}'_{RG_2} \widehat{\Gamma} \widehat{\beta}_{RG_2}}} = \begin{bmatrix} 0 & 0 & -2.29 \end{bmatrix}$ respectively. When the number of

restrictions increases, the estimated RLGSI selection response value decreases, whereas the number of zeros increases in the estimated RLGSI expected genetic gain per trait. The number of zeros in the estimated RLGSI expected genetic gain per trait is equal to the number of restrictions imposed on RLGSI by matrix U', where each restriction appears as 1.

Figure 6.1 presents the frequency distribution of the estimated RLGSI values for one (Fig. 6.1a) and two null restrictions (Fig. 6.1b). For both restrictions the frequency distribution of the estimated RLGSI values approaches the normal distribution.

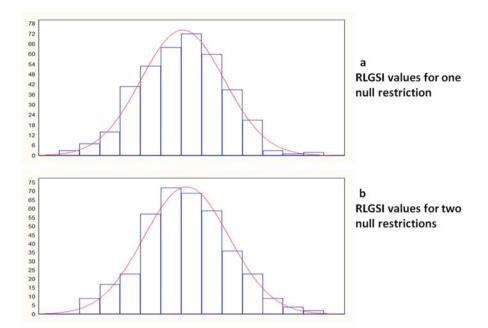


Fig. 6.1 Distribution of 380 estimated restricted linear genomic selection index (RLGSI) values with one (**a**) and two (**b**) null restrictions respectively obtained in a real testing population for one selection cycle in one environment

Now we use the simulated data set described in Chap. 2, Sect. 2.8.1, to compare RLPSI (restricted linear phenotypic selection index, Chap. 3 for details) efficiency versus RLGSI efficiency. Table 6.2 presents the estimated RLPSI and RLGSI selection response for one, two, and three null restrictions imposed by matrices $\begin{bmatrix} 1 & 0 & 0 & 0 \end{bmatrix}$

$$\mathbf{U}_{1}' = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}, \mathbf{U}_{2}' = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}, \text{ and } \mathbf{U}_{3}' = \begin{bmatrix} 1 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 \\ 0 & 0 & 1 & 0 \end{bmatrix}$$
 for five simulated

selection cycles including and not including the interval between selection cycles. In each selection cycle, the sample size was equal to 500 genotypes, each with four repetitions and four traits, whereas the selection intensity was 10% ($k_I = 1.755$); the interval lengths for the RLPSI and RLGSI were 4 and 1.5 years (Beyene et al. 2015) respectively.

Table 6.2 was divided in two parts. The first part presents the estimated RLPSI whereas the second part presents the estimated RLGSI selection responses. Columns 2, 3, and 4 in Table 6.2 present the estimated RLPSI and RLGSI selection responses not including the interval length, whereas columns 5, 6, and 7 present the estimated RLPSI and RLGSI selection response, including the interval length. The averages of the estimated RLPSI selection response not including the interval length for one, two, and three restrictions were 7.04, 5.50, and 3.90, whereas when the interval length was included, the averages were 1.76, 1.38, and 0.98 respectively. The averages of the estimated RLGSI selection response not including the interval length

Table 6.2 Estimated restricted linear phenotypic selection index (RLPSI) and RLGSI selection responses for 1, 2, and 3 null restrictions for 5 simulated selection cycles including and not including the interval between selection cycles. The interval lengths for the RLPSI and the RLGSI were 4 and 1.5 years respectively

	Estimated	RLPSI selection	on response			
	Not inclu	ding interval le	ngth	Including	g interval leng	th ^a
Cycle	1	2	3	1	2	3
1	6.87	5.54	4.13	1.72	1.39	1.03
2	8.45	5.94	4.27	2.11	1.49	1.07
3	7.17	5.79	4.16	1.79	1.45	1.04
4	6.68	5.06	3.72	1.67	1.27	0.93
5	6.02	5.16	3.24	1.51	1.29	0.81
Average	7.04	5.50	3.90	1.76	1.38	0.98
	Estimated	RLGSI selecti	on response	·		
	Not inclu	ding interval ler	ngth	Including	g interval leng	th ^b
Cycle	1	2	3	1	2	3
1	6.41	5.58	4.71	4.28	3.72	3.14
2	5.04	3.47	2.47	3.36	2.32	1.65
3	4.76	3.36	2.22	3.17	2.24	1.48
4	4.51	3.07	2.28	3.01	2.05	1.52
5	4.46	3.10	2.26	2.97	2.07	1.51
Average	5.04	3.72	2.79	3.36	2.48	1.86

^aThe estimated RLPSI selection response was divided by 4

^bThe estimated RLGSI selection response was divided by 1.5

for one, two, and three restrictions were 5.04, 3.72, and 2.79, whereas when the interval length was included the averages were 3.36, 2.48, and 1.86 respectively. These results indicated that when the interval length was included in the estimation of the RLPSI and RLGSI selection response, RLGSI efficiency was greater than RLPSI efficiency, and vice versa, when the interval length was not included the RLPSI efficiency was greater than RLGSI efficiency.

Table 6.3 presents the estimated RLPSI (first part) and RLGSI (second part) expected genetic gain per trait not including the interval between selection cycles for one, two, and three null restrictions in five simulated selection cycles. In this case, RLPSI efficiency is greater than RLGSI efficiency because the averages of the estimated RLPSI expected genetic gain per trait were -2.52, 2.26, and 2.26 for one null restriction; 2.84 and 2.65 for two null restrictions; and 3.90 for three null restrictions. For the same set of restrictions, the averages of the estimated RLGSI expected genetic gain per trait were: -1.85, 1.13, and 2.06 for one null restriction; 1.52 and 2.19 for two null restrictions, and 2.79 for three null restrictions. However, divided by the interval length (4 years in the RLPSI), the averages of the estimated RLPSI expected genetic gain per trait were -0.63, 0.57, and 0.57 for one null restrictions. In a similar manner, dividing by the interval length (1.5 years in this case), the averages of the estimated RLGSI expected genetic gain per trait were -1.23, 0.75, null restrictions.

	Estin	nated RLP	SI expe	cted gen	etic ga	ain for	one, tw	o, and th	nree nu	ıll rest	riction	s
	1				2				3			
Cycle	T1	T2	T3	T4	T1	T2	T3	T4	T1	T2	T3	T4
1	0	-2.18	2.03	2.66	0	0	2.77	2.77	0	0	0	4.13
2	0	-3.41	2.33	2.71	0	0	2.87	3.07	0	0	0	4.27
3	0	-2.30	3.12	1.74	0	0	3.11	2.68	0	0	0	4.16
4	0	-2.88	1.42	2.38	0	0	2.35	2.70	0	0	0	3.72
5	0	-1.83	2.38	1.81	0	0	3.12	2.04	0	0	0	3.24
Average	0	-2.52	2.26	2.26	0	0	2.84	2.65	0	0	0	3.90
	Estir	nated RLO	3SI expe	ected gen	netic g	ain foi	: 1, 2, ar	nd 3 null	l restri	ctions		
	1				2				3			
Cycle	T1	T2	T3	T4	T1	T2	T3	T4	T1	T2	T3	T4
1	0	-1.41	1.29	3.72	0	0	1.89	3.70	0	0	0	4.71
2	0	-2.16	1.07	1.81	0	0	1.49	1.98	0	0	0	2.47
3	0	-1.94	1.24	1.57	0	0	1.58	1.78	0	0	0	2.22
4	0	-1.90	1.02	1.60	0	0	1.34	1.73	0	0	0	2.28
5	0	-1.83	1.02	1.61	0	0	1.33	1.77	0	0	0	2.26
Average	0	-1.85	1.13	2.06	0	0	1.52	2.19	0	0	0	2.79

Table 6.3 Estimated RLPSI and RLGSI expected genetic gain per trait for 1, 2, and 3 null restrictions for 5 simulated selection cycles (each with 4 traits) not including the interval length between selection cycles

and 1.37 for one restriction; 1.01 and 1.46 for two restrictions; and 1.86 for three restrictions.

Table 6.4 presents the estimated RLPSI heritability $(\hat{h}_{I_R}^2)$ values, the estimated restricted linear genomic selection index (RLGSI) accuracy $\hat{\varphi}_{HI_{RG}}$) values, the values of $W = \frac{\hat{\rho}_{HI_{RG}}}{\hat{h}_{I_R}} L_{RP} (L_{RP} = 4)$, and the values of $\hat{p} = 100(\hat{\lambda}_R - 1)$, where $\hat{\lambda}_R = \hat{\rho}_{HI_R} / \hat{\rho}_{HI_{RG}}$ and $\hat{\rho}_{HI_R}$ is the estimated RLPSI accuracy, for one, two, and three restrictions for five simulated selection cycles. The RLGSI interval length was $L_{RG} = 1.5$ whereas the averages of the values of $W = \frac{\hat{\rho}_{HI_{RG}}}{\hat{h}_{I_R}} L_{RP}$ for each restriction were 1.22, 0.85, and 0.60; this means that the estimated Technow inequality (Technow et al. 2013), $L_{RG} < \frac{\hat{\rho}_{HI_{RG}}}{\hat{h}_{I_R}} L_{RP}$ (Chap. 5, Eq. 5.18), was not true. Thus, according to the Technow inequality results, for this data set, RLGSI efficiency in terms of time was not greater than RLPSI efficiency. The inequality $L_{RG} < \frac{\hat{\rho}_{HI_G}}{\hat{h}_{I_R}} L_{I_R}$ was not true because the estimated RLGSI accuracy was very low, whereas RLPSI heritability was high. Thus, note that the averages of the estimated RLGSI accuracy for one, two, and

three null restrictions were 0.25, 0.19, and 0.14 respectively, and the averages of the estimated RLPSI heritability values were 0.70, 0.78 and 0.88, respectively. Thus, according to these results, because the estimated RLGSI accuracy is very low and

Table 6.4 Estimated RLPSI heritability $(\widehat{h}_{I_R}^2)$, estimated RLGSI accuracy $(\widehat{\rho}_{HI_{RG}})$, estimated values of $W = \frac{\widehat{\rho}_{HI_{RG}}}{\widehat{h}_{I_R}} L_{RP} (L_{RP} = 4)$, and values of $\widehat{\rho} = 100 (\widehat{\lambda}_R - 1)$, where $\widehat{\lambda}_P = \widehat{\rho}_{HI} / \widehat{\rho}_{HI}$, and $\widehat{\rho}_{HI}$ are the estimated RLPSI accuracy values. for 1. 2. and 3 restrictions for five simulated selection cycles	timated RLF $\frac{1}{n}/\widehat{\partial}_{m}$ at	PSI heritabil $\widehat{\Omega}_{III}$ are	ity $(\widehat{h}_{I_R}^2)$, est the estimate	imated RLC ed RLPSI a	SI accuracy	y $(\widehat{ ho}_{HI_{RG}})$, est ues. for 1. 2	timated value 2. and 3 restr	ss of $W = \frac{\widehat{\rho}_H}{\widehat{h}}$ ictions for fiv	$\frac{I_{RG}}{I_R}L_{RP}$ (L_{RP} = $\frac{I_{RG}}{I_R}$ ve simulated	= 4), and valu selection cvc	es of $\hat{p} = 100$ les	$(\widehat{\lambda}_R-1),$
	RLPSI heri	ritability		RLGSI accuracy	curacy		Estimated	Estimated values of W		Estimated values of \widehat{p}	alues of \hat{p}	
Cycle	1	2	3	1	2	3	1	2	3	1	2	3
-	0.65	0.77	0.89	0.33	0.28	0.24	1.62	1.29	1.02	7.27	-1.40	-12.28
5	0.76	0.80	06.0	0.26	0.18	0.13	1.20	0.80	0.54	84.12	83.76	87.74
ε	0.71	0.80	0.88	0.24	0.17	0.11	1.16	0.77	0.49	80.34	103.03	119.72
4	0.71	0.79	0.89	0.22	0.15	0.11	1.06	0.68	0.48	79.02	97.29	94.65
S	0.67	0.76	0.86	0.22	0.15	0.11	1.07	0.70	0.48	74.31	110.97	80.61
Average	0.70	0.78	0.88	0.25	0.19	0.14	1.22	0.85	0.60	65.01	78.73	74.09

RLPSI heritability is high, RLGSI efficiency was lower than RLPSI efficiency in terms of time.

The last three columns of Table 6.4, from left to right, present the estimated p values, $\hat{p} = 100(\hat{\lambda}_R - 1)$, for one, two, and three null restrictions in five simulated selection cycles. The average of the \hat{p} values indicates that for each of the three restrictions the RLPSI efficiency was 65.05%, 78.73%, and 74.09%, greater than RLGSI efficiency at predicting the net genetic merit. Thus, for this data set, the RLPSI was a better predictor of the net genetic merit than the RLGSI in each cycle.

6.2 The Predetermined Proportional Gain Linear Genomic Selection Index

6.2.1 Objective of the PPG-LGSI

Let $\mathbf{d}' = \begin{bmatrix} d_1 & d_2 & \dots & d_r \end{bmatrix}$ be a vector $1 \times r$ (*r* is the number of predetermined proportional gains) of the predetermined proportional gains imposed by the breeder, and assume that μ_q is the population mean of the *q*th trait before selection. The objective of the predetermined proportional gain linear genomic selection index (PPG-LGSI) is to change μ_q to $\mu_q + d_q$ in the testing population, where d_q is a predetermined change in μ_q . It is possible to solve this problem minimizing the mean squared difference between $I_G = \boldsymbol{\beta}' \boldsymbol{\gamma}$ and $H = \mathbf{w}' \mathbf{g}$, $E[(H - I_G)^2]$, under the restriction $\mathbf{U}' \Gamma \boldsymbol{\beta} = \theta_G \mathbf{d}$, where θ_G is a proportionality constant, or under the

restriction $\mathbf{D}'\mathbf{U}'\mathbf{\Gamma}\boldsymbol{\beta} = \mathbf{0}$, where $\mathbf{D}' = \begin{bmatrix} d_r & 0 & \dots & 0 & -d_1 \\ 0 & d_r & \dots & 0 & -d_2 \\ \vdots & \vdots & \ddots & \vdots & \vdots \\ 0 & 0 & \dots & d_r & -d_{r-1} \end{bmatrix}$ is a matrix $(r-1) \times r$ (see Chap. 3 for details), and d_q ($q = 1, 2, \dots, r$) is the q^{th} element of vector

 $(r-1) \times r$ (see Chap. 3 for details), and d_q (q = 1, 2, ..., r) is the q^{th} element of vector $\mathbf{d}' = [d_1 \quad d_2 \quad ... \quad d_r]$; \mathbf{U}' is a matrix $(t-1) \times t$ of 1s and 0s, and $\mathbf{\Gamma} = \{\sigma_{\gamma_{qq'}}\}$ (q, q' = 1, 2, ..., t, t = number of traits) is a covariance matrix of additive genomic breeding values, $\mathbf{\gamma}' = [\gamma_1 \gamma_2 ... \gamma_t]$.

6.2.2 The Maximized PPG-LGSI Parameters

In this subsection, we minimize $E[(H - I_G)^2]$ under the restriction $\mathbf{D}'\mathbf{U}'\Gamma\boldsymbol{\beta} = \mathbf{0}$ and later under the restriction $\mathbf{U}'\Gamma\boldsymbol{\beta} = \theta_G \mathbf{d}$. Under the restriction $\mathbf{D}'\mathbf{U}'\Gamma\boldsymbol{\beta} = \mathbf{0}$, it is necessary to minimize the function

$$f_P(\boldsymbol{\beta}, \mathbf{v}) = \boldsymbol{\beta}' \boldsymbol{\Gamma} \boldsymbol{\beta} + \mathbf{w}' \mathbf{C} \mathbf{w} - 2 \mathbf{w}' \boldsymbol{\Gamma} \boldsymbol{\beta} + 2 \mathbf{v}' \mathbf{D}' \mathbf{U}' \boldsymbol{\Gamma} \boldsymbol{\beta}$$
(6.7)

with respect to $\boldsymbol{\beta}$ and $\mathbf{v}' = \begin{bmatrix} v_1 & v_2 & \dots & v_{r-1} \end{bmatrix}$, where \mathbf{v}' is a vector of Lagrange multipliers. From a mathematical point of view, Eq. (6.7) is equal to Eq. (6.1); thus, the vector of coefficients $\boldsymbol{\beta}$ of the PPG-LGSI should be similar to the vector of coefficients of the RLGSI (Eq. 6.3), i.e., the PPG-LGSI vector of coefficients is equal to

$$\boldsymbol{\beta}_{PG} = \mathbf{K}_P \mathbf{w},\tag{6.8}$$

where now $\mathbf{K}_P = [\mathbf{I}_t - \mathbf{Q}_P]$, $\mathbf{Q}_P = \mathbf{U}\mathbf{D}(\mathbf{D}'\mathbf{U}'\Gamma\mathbf{U}\mathbf{D})^{-1}\mathbf{D}'\mathbf{U}'\Gamma$, w is a vector of economic weights, and \mathbf{I}_t is an identity matrix $t \times t$. When $\mathbf{D}' = \mathbf{U}'$, $\boldsymbol{\beta}_{PG} = \boldsymbol{\beta}_{RG}$ (the RLGSI vector of coefficients), and when \mathbf{U}' is a null matrix, $\boldsymbol{\beta}_{PG} = \mathbf{w}$ (the LGSI vector of coefficients). This means that the PPG-LGSI includes the RLGSI and the LGSI as particular cases.

Under the restriction $U'\Gamma\beta = \theta_G d$ (see Chap. 3 for details) the vector of coefficients of the PPG-LGSI can be written as

$$\boldsymbol{\beta}_{PG} = \boldsymbol{\beta}_{RG} + \boldsymbol{\theta}_{G} \mathbf{U} (\mathbf{U}' \boldsymbol{\Gamma} \mathbf{U})^{-1} \mathbf{d}, \qquad (6.9)$$

where $\boldsymbol{\beta}_{RG} = \mathbf{K}_{G}\mathbf{w}$ (Eq. 6.3), $\mathbf{K}_{G} = [\mathbf{I} - \mathbf{Q}_{G}], \mathbf{Q}_{G} = \mathbf{U}(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{U}'\Gamma$, and $\mathbf{d}' = [d_{1} \quad d_{2} \quad \dots \quad d_{r}]$ is the vector of the predetermined proportional gains imposed by the breeder. It can be shown that θ_{G} , the proportionality constant, can be written as

$$\theta_G = \frac{\mathbf{d}'(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{U}'\Gamma\mathbf{w}}{\mathbf{d}'(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{d}}.$$
(6.10)

When $\theta_G = 0$, $\beta_{PG} = \beta_{RG}$, and when U' is a null matrix, $\beta_{PG} = \mathbf{w}$. Equations (6.8) and (6.9) give the same results, that is, both equations express the same result in a different mathematical way.

The maximized selection response and expected genetic gain per trait of the PPG-LGSI can be written as

$$R_{PG} = \frac{k_I}{L_G} \sqrt{\beta'_{PG} \Gamma \beta_{PG}}$$
(6.11)

and

$$\mathbf{E}_{PG} = \frac{k_I}{L_G} \frac{\mathbf{\Gamma} \boldsymbol{\beta}_{PG}}{\sqrt{\boldsymbol{\beta}_{PG}' \mathbf{\Gamma} \boldsymbol{\beta}_{PG}}},\tag{6.12}$$

respectively, where L_G is the time required to complete a selection cycle using the PPG-LGSI. Equations (6.11) and (6.12) depend only on GEBV information.

6.2.3 Statistical Properties of the PPG-LGSI

Assuming that $H = \mathbf{w}'\mathbf{g}$ and the PPG-LGSI ($I_{PG} = \boldsymbol{\beta}'_{PG}\boldsymbol{\gamma}$) have bivariate joint normal distribution, $\boldsymbol{\beta}_{PG} = \mathbf{K}_P \mathbf{w}$; $\boldsymbol{\Gamma}$, \mathbf{C} , and \mathbf{w} are known, it can be shown that PPG-LGSI has the following statistical properties:

- 1. The vector $\beta_{PG} = \mathbf{K}_P \mathbf{w}$ minimizes the mean square error under the restriction $\mathbf{D}'\mathbf{U}'\mathbf{\Gamma}\boldsymbol{\beta} = \mathbf{0}$.
- 2. The variance of $I_{PG} = \beta'_{PG} \gamma (\sigma_{I_{PG}}^2 = \beta'_{PG} \Gamma \beta_{PG})$ is equal to the covariance between $I_{PG} = \beta'_{PG} \gamma$ and $H = \mathbf{w}' \mathbf{g} (\sigma_{HI_{PG}} = \mathbf{w}' \Gamma \beta_{PG})$.
- 3. The maximized correlation between *H* and I_{PG} (also called PPG-LGSI accuracy) is equal to $\rho_{HI_{PG}} = \frac{\sigma_{I_{PG}}}{\sigma_{H}}$, where $\sigma_{I_{PG}} = \sqrt{\beta'_{PG}\Gamma\beta_{PG}}$ and $\sigma_{H} = \sqrt{\mathbf{w'Cw}}$ are the standard deviations of $I_{PG} = \beta'_{PG}\gamma$ and $H = \mathbf{w'g}$ respectively.

4. The variance of the predicted error,
$$Var(H - I_{PG}) = (1 - \rho_{HI_{PG}}^2)\sigma_H^2$$
, is minimal.

The statistical PPG-LGSI properties are equal to the statistical PPG-LPSI properties, then, the PPG-LGSI is an application of the PPG-LPSI to the genomic selection context.

6.2.4 Numerical Example

To illustrate the PPG-LGSI theory, we use the estimated matrix $\widehat{\Gamma} = \begin{bmatrix} 0.21 & 2.95 & 5.00\\ 2.95 & 42.41 & 71.11\\ 5.00 & 71.11 & 121.53 \end{bmatrix}$ and the GEBVs associated with the traits GY (ton

 ha^{-1}), EHT (cm), and PHT (cm), described in Sect. 6.1.3.

It is necessary to estimate the PPG-LGSI vector of coefficients $\boldsymbol{\beta}_{PG} = \boldsymbol{\beta}_{RG} + \theta_g \mathbf{U} (\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{d}$ (Eqs. 6.9 and 6.10). In Sect. 6.1.3, we showed that the estimated vectors of coefficients of $\boldsymbol{\beta}_{RG} = \mathbf{K}_G \mathbf{w}$ for the null restrictions $\mathbf{U}'_1 = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$ and $\mathbf{U}'_2 = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$ were $\hat{\boldsymbol{\beta}}'_{RG1} = \mathbf{w}'\hat{\mathbf{K}}'_{G1} = \begin{bmatrix} 3.78 & -0.1 & -0.1 \end{bmatrix}$ and $\hat{\boldsymbol{\beta}}'_{RG2} = \mathbf{w}'$ $\hat{\mathbf{K}}'_{G2} = \begin{bmatrix} 1.12 & 0.09 & -0.1 \end{bmatrix}$ respectively, where $\mathbf{w}' = \begin{bmatrix} 5 & -0.1 & -0.1 \end{bmatrix}$. This means that to estimate $\boldsymbol{\beta}_{PG} = \boldsymbol{\beta}_{RG} + \theta_G \mathbf{U}(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{d}$, we need only to estimate $\theta_G \mathbf{U}(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{d}$ for both sets of restrictions.

Consider matrix $\mathbf{U}_{1}' = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$ and let $d_{1} = 7.0$ be the predetermined proportional gain restriction for trait 1. We can estimate θ_{G} and $\mathbf{U}(\mathbf{U}'\Gamma\mathbf{U})^{-1}\mathbf{d}$ as $\widehat{\theta}_{G1} = \frac{7.0(\mathbf{U}_{1}'\widehat{\Gamma}\mathbf{U}_{1})^{-1}\mathbf{U}_{1}'\widehat{\Gamma}\mathbf{w}}{7.0(\mathbf{U}_{1}'\widehat{\Gamma}\mathbf{U}_{1})^{-1}7.0} = \begin{bmatrix} 33.333\\0\\0 \end{bmatrix}$, whence the PPG-LGSI vector of coefficients was $\widehat{\boldsymbol{\beta}}_{PG_{1}} = \widehat{\boldsymbol{\beta}}_{RG_{1}} + \widehat{\theta}_{G_{1}}\mathbf{U}_{1}(\mathbf{U}_{1}'\widehat{\Gamma}\mathbf{U}_{1})^{-1}7.0 = \begin{bmatrix} 5.0\\-0.1\\-0.1 \end{bmatrix}$, and the estimated PPG-LGSI was $\widehat{I}_{PG_1} = 5.0 \text{GEBV}_1 - 0.1 \text{GEBV}_2 - 0.1 \text{GEBV}_3$. In a similar manner, we can estimate the PPG-LGSI vector of coefficients under restrictions $\mathbf{U}_2' = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$ and $\mathbf{d}_2' = \begin{bmatrix} 7 & -3 \end{bmatrix}$. In this case, $\widehat{\boldsymbol{\beta}}_{PG_2} = \widehat{\boldsymbol{\beta}}_{RG_2} + \widehat{\boldsymbol{\theta}}_{G_2} \mathbf{U}_2 (\mathbf{U}_2' \widehat{\boldsymbol{\Gamma}} \mathbf{U}_2)^{-1} \mathbf{d}_2 = \begin{bmatrix} 4.97 \\ -0.18 \\ -0.10 \end{bmatrix}$ and the estimated PPG-LGSI was $\widehat{I}_{PG_2} = 4.97 \text{GEBV}_1 - 0.18 \text{GEBV}_2 - 0.1 \text{GEBV}_3$.

Figure 6.2 presents the frequency distribution of the estimated PPG-LGSI values for one (Fig. 6.2a) and two (Fig. 6.2b) predetermined restrictions, d = 7 and $\mathbf{d}' = \begin{bmatrix} 7 & -3 \end{bmatrix}$ respectively, obtained in a real testing population for one selection cycle in one environment. For both restrictions, the frequency distribution of the estimated PPG-LGSI values approaches the normal distribution.

Assume a selection intensity of 10% ($k_{I_G} = 1.755$); then, for one predetermined restriction, where $\mathbf{U}'_1 = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$ and $d_1 = 7.0$, the estimated PPG-LGSI selection response and expected genetic gain per trait, not including the interval length, were $\widehat{R}_{PG_1} = k_{I_G} \sqrt{\widehat{\beta}'_{PG_1}\widehat{\Gamma}\widehat{\beta}_{PG_1}} = 1.05$ and $\widehat{E}'_{PG_1} = k_I \frac{\widehat{\beta}'_{PG_1}\widehat{\Gamma}}{\sqrt{\widehat{\beta}'_{PG_1}\widehat{\Gamma}\widehat{\beta}_{PG_1}}} = [0.74 \ 9.92 \ 16.54]$

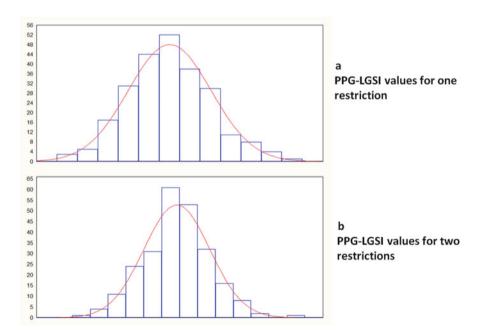


Fig. 6.2 Distribution of 380 estimated predetermined proportional gain linear genomic selection index (PPG-LGSI) values with one (**a**) and two (**b**) predetermined restrictions, d = 7 and $\mathbf{d}' = \begin{bmatrix} 7 & -3 \end{bmatrix}$ respectively, obtained in a real testing population for one selection cycle in one environment

respectively. For two restrictions, with $\mathbf{U}_{2}' = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$ and $\mathbf{d}' = \begin{bmatrix} 7 & -3 \end{bmatrix}$, the estimated RLGSI selection response and expected genetic gains, not including the interval length, were $\widehat{R}_{PG_{2}} = k_{I_{G}} \sqrt{\widehat{\beta}'_{PG_{2}} \widehat{\Gamma} \widehat{\beta}_{G_{2}}} = 0.52$ and $\widehat{\mathbf{E}}'_{PG_{2}} = k_{I} \frac{\widehat{\beta}'_{PG_{2}} \widehat{\Gamma}}{\sqrt{\widehat{\beta}'_{PG_{2}} \widehat{\Gamma} \widehat{\beta}_{PG_{2}}}} = k_{I_{G}} \sqrt{\widehat{\beta}'_{PG_{2}} \widehat{\Gamma} \widehat{\beta}_{PG_{2}}} = 0.52$ and $\widehat{\mathbf{E}}'_{PG_{2}} = k_{I} \frac{\widehat{\beta}'_{PG_{2}} \widehat{\Gamma} \widehat{\beta}_{PG_{2}}}{\sqrt{\widehat{\beta}'_{PG_{2}} \widehat{\Gamma} \widehat{\beta}_{PG_{2}}}} = 0.52$

 $[0.11 - 0.05 \ 0.14]$ respectively.

Now, we use the simulated data set described in Chap. 2, Sect. 2.8.1 to compare PPG-LGSI efficiency versus predetermined proportional gain linear phenotypic selection index (PPG-LPSI) efficiency. Let $\mathbf{U}_1' = \begin{bmatrix} 1 & 0 & 0 \end{bmatrix}$, $\mathbf{U}_2' = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \end{bmatrix}$,

and $\mathbf{U}'_3 = \begin{bmatrix} 1 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 \\ 0 & 0 & 1 & 0 \end{bmatrix}$ be the matrices and $d_1 = 7$, $\mathbf{d}'_2 = \begin{bmatrix} 7 & -3 \end{bmatrix}$, and $\mathbf{d}'_3 = \begin{bmatrix} 7 & -3 \end{bmatrix}$

 $\begin{bmatrix} 7 & -3 & 5 \end{bmatrix}$ the vectors for one, two, and three predetermined restrictions respectively. Table 6.5 presents the estimated PPG-LPSI and PPG-LGSI selection response for each predetermined restriction in five simulated selection cycles including and not including the interval between selection cycles (4 years for the PPG-LPSI and 1.5 years for the PPG-LGSI); estimated PPG-LPSI and PPG-LGSI accuracy; and estimated variance of the predicted error (VPE). In each selection cycle, the sample size was equal to 500 genotypes, each with four repetitions and four traits. The selection intensity was 10% ($k_I = 1.755$).

The averages of the estimated PPG-LPSI selection response not including the interval length were 15.14, 14.87, and 13.30, whereas when the interval length was included, the average selection responses were 3.79, 3.72, and 3.33, for one, two, and three predetermined restrictions respectively (Table 6.5). The averages of the estimated PPG-LGSI selection responses not including the interval length for one, two, and three predetermined restrictions were 14.48, 13.47, and 11.26 respectively, and when the interval length was included, the selection responses were 9.65, 8.98, and 7.51 respectively (Table 6.5). These results indicate that when the interval length was included in the estimation of the PPG-LPSI and PPG-LGSI selection responses, PPG-LGSI efficiency was greater than PPG-LPSI efficiency, and vice versa, when the interval length was not included in the PPG-LPSI and PPG-LGSI selection responses, PPG-LPSI efficiency was higher than PPG-LGSI efficiency.

The averages of the estimated VPE values of the PPG-LPSI for one, two, and three predetermined restrictions were 22.42, 30.56, and 41.17 respectively, whereas the estimated VPE values of the PPG-LGSI (see Sect. 6.2.3 for details) were 59.80, 66.95, and 83.98, respectively, that is, in all selection cycles, the VPE of the PPG-LPSI was lower than that of the PPG-LGSI. This means that for this data set, the PPG-LPSI was a better predictor of the net genetic merit than the PPG-LGSI. These results can be explained by observing that the averages of the estimated PPG-LPSI accuracies were 0.88, 0.86, and 0.77, whereas the estimated PPG-LGSI accuracies were 0.65, 0.68, and 0.57 for each predetermined restriction, that is, the estimated PPG-LGSI accuracies were lower than the estimated PPG-LPSI accuracies for this data set.

	Estimated PP	ISdT-Ddd	G-LPSI selection response	ponse								
	Not including	ding interval length	l length	Including	Including interval length ^a	ngth ^a	Estimated	Estimated PPG-LPSI accuracy	ccuracy	Estimate	Estimated PPG-LPSI VPE	VPE
Cycle	1	2	3		2	Э	1	2	e	1	2	б
	17.81	16.72	15.23	4.45	4.18	3.81	0.91	0.85	0.77	22.53	50.44	57.14
2	15.69	15.59	14.39	3.92	3.90	3.60	0.88	0.88	0.81	22.66	27.88	37.03
3	14.22	14.16	13.18	3.56	3.54	3.30	0.87	0.86	0.80	21.95	26.14	33.55
4	14.34	14.33	11.56	3.59	3.58	2.89	0.86	0.86	0.70	22.84	25.84	45.46
5	13.64	13.56	12.16	3.41	3.39	3.04	0.86	0.85	0.76	22.13	22.49	32.69
Average	15.14	14.87	13.30	3.79	3.72	3.33	0.88	0.86	0.77	22.42	30.56	41.17
ш	Estimated PPG-	PG-LGSI se	-LGSI selection response	onse								
4	Not interval len	l length		Including i	Including interval length	th	Estimated	Estimated PPG-LGSI accuracy	curacy	Estimated	Estimated PPG-LGSI VPE	VPE
Cycle 1		2	3	1	2	3		2	3	1	2	б
1 2	21.22	17.83	16.66	14.15	11.89	11.10	0.65	0.91	0.85	74.75	22.25	35.43
2	13.90	13.52	10.65	9.27	9.01	7.10	0.72	0.70	0.55	58.66	62.06	84.62
3	13.59	13.15	10.70	9.06	8.77	7.13	0.70	0.67	0.55	63.49	67.33	86.30
4	12.30	11.84	9.36	8.20	7.89	6.24	0.61	0.59	0.46	83.26	86.91	103.93
5 1	11.38	11.03	8.96	7.59	7.35	5.97	0.56	0.54	0.44	93.62	96.21	109.63
Average 1	14.48	13.47	11.26	9.65	8.98	7.51	0.65	0.68	0.57	74.75	66.95	83.98

^{ar}The estimated LPSI selection response was divided by 4 and the estimated LGS selection response was divided by 1.5

Table 6.5 Estimated predetermined proportional gain linear phenotypic and genomic selection index (PPG-LPSI and PPG-LGSI respectively) selection responses for 1, 2 and 3 predetermined restrictions for five simulated selection cycles including and not including the interval between selection cycles (4 years

Table 6.6 Estimated PPG-LPSI heritability (\hat{h}_P^2) , values of $W_P = \frac{\hat{\rho}_{HI_G}}{\hat{h}_P} L_P (L_P = 4)$, and the ratio of the estimated PPG-LPSI accuracy $(\hat{\rho}_{HI_P})$ to the estimated PPG-LGSI accuracy $(\hat{\rho}_{HI_{PG}})$: $\hat{\lambda}_P = \hat{\rho}_{HI_P} / \hat{\rho}_{HI_{PG}}$, and values of $\hat{p} = 100(\hat{\lambda}_P - 1)$ for 1, 2 and 3 predetermined restrictions for five simulated selection cycles

	PPG-L	PSI herita	bility	Values	s of W_P		Estimated	ratio value	es (\hat{p})
Cycle	1	2	3	1	2	3	1	2	3
1	0.84	0.77	0.83	4.71	4.13	3.72	-18.62	-6.71	-10.20
2	0.80	0.78	0.83	3.22	3.17	2.42	18.30	20.54	32.04
3	0.77	0.76	0.8	3.18	3.09	2.45	19.89	21.59	31.42
4	0.76	0.75	0.78	2.80	2.71	2.10	29.16	31.84	33.75
5	0.75	0.75	0.79	2.57	2.49	1.97	35.26	36.55	42.35
Average	0.72	0.71	0.76	3.29	3.12	2.53	16.80	20.76	25.87

Table 6.6 presents the estimated predetermined PPG-LPSI heritability (\hat{h}_p^2) values, $W_P = \frac{\hat{\rho}_{HI_G}}{\hat{h}_P} L_P (L_P = 4)$ values, and ratio of the estimated PPG-LPSI accuracy $(\hat{\rho}_{HI_P})$ to the estimated PPG-LGSI accuracy $(\hat{\rho}_{HI_{PG}})$, i.e., $\hat{\lambda}_P = \hat{p}_{HI_P}/\hat{p}_{HI_{PG}}$, and, finally, values of $\hat{p} = 100(\hat{\lambda}_P - 1)$ for one, two, and three null restrictions for five simulated selection cycles.

The averages of the W_P values for one, two, and three null restrictions were 3.29, 3.12, and 2.53, respectively, whereas the PPG-LGSI interval length was 1.5 $(L_G = 1.5)$. This means that the estimated Technow inequality, $L_G < \frac{\hat{\rho}_{HI_G}}{\hat{h}_P} L_P$ (see Chap. 5, Eq. 5.18) was true. Thus, PPG-LGSI efficiency in terms of time was greater

than PPG-LPSI efficiency for this data set. These results coincide with those obtained earlier in this chapter, when we compared PPG-LGSI efficiency versus PPG-LPSI efficiency in terms of interval length. However, the average values of $\hat{p} = 100(\hat{\lambda}_P - 1)$ (see Chap. 5, Eq. 5.15) were, in percentage terms, 16.80%, 20.76%, and 25.85% for each restriction. These latter results indicate that for this data set, the PPG-LPSI was a better predictor of the net genetic merit than the PPG-LGSI. This is because the estimated PPG-LPSI accuracies were higher than the estimated PPG-LPSI accuracies for this data set. We found similar results when we compared the PPG-LPSI VPE versus PPG-LGSI VPE (Table 6.5).

6.3 The Combined Restricted Linear Genomic Selection Index

The combined restricted linear genomic selection index (CRLGSI) is based on the RLPSI (Chap. 3) and combined linear genomic selection index (CLGSI, Chap. 5) theory. In the RLPSI, the breeder's objective is to improve only (t - r) of t (r < t)

traits, leaving r of them fixed; the same is true for the CRLGSI, but in the latter case, it is necessary to impose 2r restrictions, i.e., we need to fix r traits and their associated r GEBVs to obtain results similar to those obtained with the RLPSI. This is the main difference between the CRLGSI and the RLPSI.

It can be shown that $Cov(I_C, \mathbf{a}_C) = \Psi_C \beta_C$ is the covariance between the breeding value vector $(\mathbf{a}'_C = [\mathbf{g}' \quad \gamma'])$ and the CLGSI, $I_C = \beta'_C \mathbf{t}_C$ (see Chap. 5 for details), where $\mathbf{t}'_C = [\mathbf{y}' \quad \gamma']$. In the CRLGSI, we want some covariances between the linear combinations of \mathbf{a}_C ($\mathbf{U}'_C \mathbf{a}_C$) and CLGSI to be zero, i.e., $Cov(I_C, \mathbf{U}'_C \mathbf{a}_C) = \mathbf{U}'_C \Psi_C \beta_C = \mathbf{0}$, where \mathbf{U}'_C is a matrix $2(t-1) \times 2t$ of 1s and 0s (1 indicates that the trait and its associated GEBV are restricted, and 0 that the trait and its GEBV have no restrictions) and $\Psi_C = \begin{bmatrix} \mathbf{C} & \mathbf{\Gamma} \\ \mathbf{\Gamma} & \mathbf{\Gamma} \end{bmatrix}$ is a block covariance matrix of $\mathbf{a}'_C = [\mathbf{g}' \quad \gamma']$ where \mathbf{C} and $\mathbf{\Gamma}$ are the covariance matrices of breeding (\mathbf{g}) and genomic (γ) values respectively. This problem can be solved by minimizing the mean squared difference between the CLGSI and H ($E[(H - \mathbf{I}_C)^2]$) under the restriction $\mathbf{U}'_C \Psi_C \beta_C = \mathbf{0}$ similar to the RLGSI in Sect. 6.1.

6.3.1 The Maximized CRLGSI Parameters

Let $\mathbf{T}_C = \begin{bmatrix} \mathbf{P} & \mathbf{\Gamma} \\ \mathbf{\Gamma} & \mathbf{\Gamma} \end{bmatrix}$ be the block covariance matrix of $\mathbf{t}'_C = \begin{bmatrix} \mathbf{y}' & \mathbf{\gamma}' \end{bmatrix}$ where \mathbf{P} and $\mathbf{\Gamma}$ are the covariance matrices of phenotypic (\mathbf{y}) and genomic ($\mathbf{\gamma}$) values respectively. Based on the Eq. (6.1) result, it can be shown that the CRLGSI vector of coefficients that minimizes $E[(H - \mathbf{I}_C)^2]$ under the restriction $\mathbf{U}'_C \mathbf{\Psi}_C \mathbf{\beta}_C = \mathbf{0}$ is

$$\boldsymbol{\beta}_{CR} = \mathbf{K}_C \boldsymbol{\beta}_C, \tag{6.13}$$

where $\mathbf{K}_{C} = [\mathbf{I} - \mathbf{Q}_{C}]$, $\mathbf{Q}_{C} = \mathbf{T}_{C}^{-1} \mathbf{\Phi}_{C} (\mathbf{\Phi}_{C}' \mathbf{T}_{C}^{-1} \mathbf{\Phi}_{C})^{-1} \mathbf{\Phi}_{C}'$, $\mathbf{\Phi}_{C} = \mathbf{U}_{C}' \mathbf{\Psi}_{C}$, and $\mathbf{\beta}_{C} = \mathbf{T}_{C}^{-1} \mathbf{\Psi}_{C} \mathbf{a}_{C}$ (the vector of coefficients of the CLGSI, see Chap. 5 for details); \mathbf{T}_{C}^{-1} is the inverse of matrix \mathbf{T}_{C} , and \mathbf{I} is an identity matrix $2t \times 2t$. When no restrictions are imposed on any of the traits, \mathbf{U}_{C}' is a null matrix and $\mathbf{\beta}_{CR} = \mathbf{\beta}_{C}$ (the vector of coefficients of the CLGSI). That is, the CRLGSI is more general than the CLGSI. Similar to the RLPSI and the RLGSI, matrices \mathbf{K}_{C} and \mathbf{Q}_{C} are idempotent ($\mathbf{K}_{C} = \mathbf{K}_{C}^{2}$ and $\mathbf{Q}_{C} = \mathbf{Q}_{C}^{2}$) and orthogonal ($\mathbf{K}_{C}\mathbf{Q}_{C} = \mathbf{Q}_{C}\mathbf{K}_{C} = \mathbf{0}$), that is, \mathbf{K}_{C} and \mathbf{Q}_{C} are projectors. Thus, we can assume that the CRLGSI has similar properties to those described for the RLPSI (see Chap. 3 for details) when matrices $\mathbf{\Psi}_{C} = \begin{bmatrix} \mathbf{C} & \mathbf{\Gamma} \\ \mathbf{\Gamma} & \mathbf{\Gamma} \end{bmatrix}$ and $\mathbf{T}_{C} = \begin{bmatrix} \mathbf{P} & \mathbf{\Gamma} \\ \mathbf{\Gamma} & \mathbf{\Gamma} \end{bmatrix}$ are known.

The maximized selection response and the optimized expected genetic gain per trait of the CRLGSI can be written as

6 Constrained Linear Genomic Selection Indices

$$R_{CR} = \frac{k_I}{L_I} \sqrt{\beta_{CR}' \mathbf{T}_C \beta_{CR}}$$
(6.14)

and

$$\mathbf{E}_{CR} = \frac{k_I}{L_I} \frac{\boldsymbol{\Psi} \boldsymbol{\beta}_{CR}}{\sqrt{\boldsymbol{\beta}_{CR}' \mathbf{T}_C \boldsymbol{\beta}_{CR}}},\tag{6.15}$$

respectively. Although in the RLGSI and the PPG-LGSI the interval between selection cycles is denoted as L_G , in the CRLGSI it is denoted as L_I . This is because the RLPSI and the CRLGSI should have the same interval between selection cycles.

6.3.2 Numerical Examples

To illustrate the CRLGSI theoretical results, we use a real training maize (*Zea mays*) F_2 population with 248 genotypes (each with two repetitions), 233 molecular markers, and three traits: GY (ton ha⁻¹), EHT (cm), and PHT (cm). Matrices **P** and **C** were estimated based on Eqs. (2.22) to (2.24) described in Chap. 2. The

 $\widehat{\mathbf{P}} \text{ and } \widehat{\mathbf{C}} \text{ were estimated based on Eqs. (2.22) to (2.24) described in Chap. 2. The estimated matrices were <math display="block"> \widehat{\mathbf{P}} = \begin{bmatrix} 0.45 & 1.33 & 2.33 \\ 1.33 & 65.07 & 83.71 \\ 2.33 & 83.71 & 165.99 \end{bmatrix}$ and $\widehat{\mathbf{C}} = \begin{bmatrix} 0.07 & 0.61 & 1.06 \\ 0.61 & 17.93 & 22.75 \\ 1.06 & 22.75 & 44.53 \end{bmatrix}.$ In a similar manner, we estimated matrix Γ using Eqs. (5.21) to (5.23) described in Chap. 5. The estimated matrix was $\widehat{\mathbf{\Gamma}} = \begin{bmatrix} 0.07 & 0.65 & 1.05 \\ 0.65 & 10.62 & 14.25 \\ 1.05 & 14.25 & 26.37 \end{bmatrix}.$

To estimate the CRLGSI and its associated parameters (selection response, expected genetic gain per trait, etc.), we need to obtain matrices $\widehat{\mathbf{T}}_{C} = \begin{bmatrix} \widehat{\mathbf{P}} & \widehat{\mathbf{\Gamma}} \\ \widehat{\mathbf{\Gamma}} & \widehat{\mathbf{\Gamma}} \end{bmatrix}$ and $\widehat{\mathbf{\Psi}}_{C} = \begin{bmatrix} \widehat{\mathbf{C}} & \widehat{\mathbf{\Gamma}} \\ \widehat{\mathbf{\Gamma}} & \widehat{\mathbf{\Gamma}} \end{bmatrix}$ using phenotypic and genomic information and the estimated CRLGSI vector of coefficients $\widehat{\boldsymbol{\beta}}_{CR} = \widehat{\mathbf{K}}_{C}\widehat{\boldsymbol{\beta}}_{C}$, where $\widehat{\mathbf{K}}_{C} = [\mathbf{I} - \widehat{\mathbf{Q}}_{C}]$, $\widehat{\mathbf{Q}}_{C} = \widehat{\mathbf{T}}_{C}^{-1}\widehat{\mathbf{\Phi}}_{C}(\widehat{\mathbf{\Phi}}_{C}'\widehat{\mathbf{T}}_{C}^{-1}\widehat{\mathbf{\Phi}}_{C})^{-1}\widehat{\mathbf{\Phi}}_{C}', \ \widehat{\mathbf{\Phi}}_{C} = \mathbf{U}_{C}'\widehat{\mathbf{\Psi}}_{C}$, and $\widehat{\boldsymbol{\beta}}_{C} = \widehat{\mathbf{T}}_{C}^{-1}\widehat{\mathbf{\Psi}}_{C}\mathbf{a}_{C}$.

We have indicated that the main difference between the RLGSI and the CRLGSI is matrix \mathbf{U}_{C}' , on which we now need to impose two restrictions: one for the trait and another for its associated GEBV. Consider the (*Zea mays*) F₂ population described earlier and suppose that we restrict trait GY; then, matrix \mathbf{U}_{C}' should be constructed as $\mathbf{U}_{C_1}' = \begin{bmatrix} 1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 \end{bmatrix}$. If we restrict traits GY and EHT, matrix \mathbf{U}_{C}' should

be constructed as $\mathbf{U}'_{C_2} = \begin{bmatrix} 1 & 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 \\ 0 & 0 & 0 & 0 & 1 & 0 \end{bmatrix}$, etc. The procedure for obtaining matrices $\widehat{\mathbf{K}}_C = [\mathbf{I} - \widehat{\mathbf{Q}}_C], \ \widehat{\mathbf{Q}}_C = \widehat{\mathbf{T}}_C^{-1} \widehat{\mathbf{\Phi}}_C (\widehat{\mathbf{\Phi}}'_C \widehat{\mathbf{T}}_C^{-1} \widehat{\mathbf{\Phi}}_C)^{-1} \widehat{\mathbf{\Phi}}'_C$, and $\widehat{\mathbf{\Phi}}_C = \mathbf{U}'_C \widehat{\mathbf{\Psi}}_C$ is

similar to that described in Chap. 3.

Let $\mathbf{w}' = \begin{bmatrix} 5 & -0.1 & -0.1 & 0 & 0 \end{bmatrix}$ be the vector of economic weights and assume that we restrict trait GY; in this case, according to the estimated matrices $\hat{\mathbf{P}}$, $\hat{\mathbf{C}}$, and $\hat{\mathbf{\Gamma}}$ described earlier, the estimated CRLGSI vector of coefficients was $\hat{\mathbf{\beta}}'_{RG} = [0.076 \ -0.004 \ -0.018 \ 2.353 \ -0.096 \ -0.082]$, whence the estimated CRLGSI can be written as

$$\hat{I}_{CR} = 0.076$$
GY $- 0.004$ EHT $- 0.018$ PHT $+ 2.353$ GEBV_{GY} $- 0.096$ GEBV_{EHT} $- 0.082$ GEBV_{PHT}

where GEBV_{GY}, GEBV_{EHT}, and GEBV_{PHT} are the GEBVs associated with traits GY, EHT, and PHT respectively. The same procedure is valid for two or more restrictions.

Figure 6.3 presents the frequency distribution of the estimated CRLGSI values for one (Fig. 6.3a) and two null restrictions (Fig. 6.3b) using matrices U'_{C_1} and U'_{C_2} , and the real data set of the F₂ population. For both restrictions, the frequency distribution of the estimated CRLGSI values approaches normal distribution.

Suppose selection intensity of 10% ($k_I = 1.755$), matrix $\mathbf{U}_{C_1}' = \begin{bmatrix} 1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 \end{bmatrix} \text{ and that the vector of economic weights is } \mathbf{w}' = \widehat{\mathbf{v}}_{C_1} \widehat{\mathbf{v}}_{C_1} = \widehat{\mathbf{v}}_{C_1} \widehat{\mathbf{v}}_$ $\begin{bmatrix} 5 & -0.1 & -0.1 & 0 & 0 \end{bmatrix}$; then, according to the estimated matrices $\hat{\mathbf{P}}, \hat{\mathbf{C}}$, and $\hat{\mathbf{\Gamma}}$ described earlier, the estimated CRLGSI selection response and the estimated CRLGSI expected genetic gain per trait were $\widehat{R}_{CR} = k_I \sqrt{\widehat{\beta}'_{CR}} \widehat{\mathbf{T}}_C \widehat{\boldsymbol{\beta}}_{CR} = 0.96$ and $\widehat{\mathbf{E}}_{CR}' = k_I \frac{\widehat{\boldsymbol{\beta}}_{CR}' \widehat{\boldsymbol{\Psi}}}{\sqrt{\widehat{\boldsymbol{\beta}}_{CR}' \widehat{\mathbf{T}}_C \widehat{\boldsymbol{\beta}}_{CR}}} = \begin{bmatrix} 0 & -3.53 & -6.03 & 0 & -2.93 & -4.87 \end{bmatrix} \text{ respectively,}$

whereas the estimated CRLGSI accuracy was $\hat{\rho}_{HI_{CR}} = \frac{\hat{\sigma}_{I_{CR}}}{\hat{\sigma}_{II}} = 0.51$ (see Chaps. 3) and 5 for details).

Now, we use the simulated data described in Chap. 2, Sect. 2.8.1 to compare CRLGSI efficiency versus RLGSI efficiency. The criteria for this comparison are the Technow inequality (Eq. 5.18, Chap. 5) and the ratio of the estimated CRLGSI accuracy $(\hat{\rho}_{HI_{CR}})$ to the estimated RLGSI accuracy $(\hat{\rho}_{HI_R})$ expressed as percentages (Eq. 5.17, Chap. 5), i.e., $\hat{p} = 100(\hat{\lambda}_{CR} - 1)$, where $\hat{\lambda}_P = \hat{\rho}_{HI_{CR}}/\hat{\rho}_{HI_R}$, for one, two, and three null restrictions for five simulated selection cycles.

Table 6.7 presents the estimated CRLGSI heritability (h_C^2) , the estimated RLGSI accuracy $(\hat{\rho}_{HI_R})$, the values of $W_C = \frac{\hat{\rho}_{HI_R}}{\hat{h}_I} L_I$ $(L_I = 4)$, and the values of

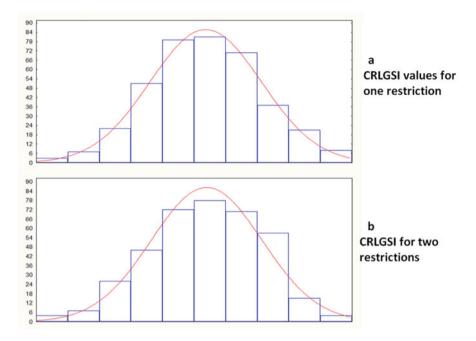


Fig. 6.3 Distribution of 244 estimated combined restricted linear genomic selection index (CRLGSI) values with one (**a**) and two (**b**) null restrictions respectively obtained in a real training population for one selection cycle in one environment

 $\hat{p} = 100(\hat{\lambda}_{CR} - 1)$, where $\hat{\lambda}_{CR} = \hat{\rho}_{HI_{CR}}/\hat{\rho}_{HI_R}$ and $\hat{\rho}_{HI_{CR}}$ is the estimated CRLGSI accuracy, for one, two, and three null restrictions for five simulated selection cycles. The averages of the $W_C = \frac{\hat{\rho}_{HI_R}}{\hat{h}_C} L_I$ values for one, two, and three null restrictions were 1.26, 0.92, and 0.59 respectively, whereas the RLGSI interval length was 1.5 $(L_G = 1.5)$. This means that the estimated Technow inequality $(L_G < \frac{\hat{\rho}_{HI_G}}{\hat{h}_I} L_I)$ was not true. Thus, for this data set, RLGSI efficiency in terms of time is not greater than CRLGSI efficiency. The inequality $L_G < \frac{\hat{\rho}_{HI_G}}{\hat{h}_I} L_I$ was not true because the estimated RLGSI accuracy was very low, whereas CRLGSI heritability was high. Thus, note that the averages of the estimated RLGSI accuracy for one, two, and three null restrictions were 0.25, 0.19, and 0.14 respectively, whereas the averages of the estimated CRLGSI heritability values were 0.72, 0.75, and 0.89 respectively. Thus, according to these results, when the estimated RLGSI accuracy is very low and the estimated CRLGSI heritability is high, RLGSI efficiency will be lower than CRLGSI efficiency in terms of time.

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$(L_I = 4)$, and values of \widehat{p}		$= 100 \left(\widehat{\lambda}_{CR} - \right)$	1), where $\widehat{\lambda}_i$	$c_R = \widehat{ ho}_{Hl_{CR}}/2$	$\widehat{ ho}_{HI_R}$ and $\widehat{ ho}_I$	Huck is the e	stimated CR	tLGSI accu	racy, for 1,	2, and 3 null	= $100(\hat{\lambda}_{CR} - 1)$, where $\hat{\lambda}_{CR} = \hat{\rho}_{HI_{CR}}/\hat{\rho}_{HI_R}$ and $\hat{\rho}_{HI_{CR}}$ is the estimated CRLGSI accuracy, for 1, 2, and 3 null restrictions for five	n or five
simulated selection cycles	tion cycles											
	CRLGSI he	neritability		RLGSI accuracy	uracy		Values of W_C	N_C		Values of \hat{p}		
Cycle	1	2	3	1	2	3	1	2	3	1	2	3
1	0.39	0.41	0.90	0.33	0.28	0.24	2.11	1.75	1.01	15.15	50.00	8.33
2	0.82	0.84	0.89	0.26	0.18	0.13	1.15	0.79	0.55	46.15	72.22	61.54
ю	0.80	0.84	0.88	0.24	0.17	0.11	1.07	0.74	0.47	66.67	82.35	81.82
4	0.82	0.85	0.89	0.22	0.15	0.11	0.97	0.65	0.47	68.18	93.33	72.73
5	0.77	0.82	0.91	0.22	0.15	0.11	1.00	0.66	0.46	72.73	93.33	81.82



61.25

78.25

53.78

0.59

0.92

1.26

0.14

0.19

0.25

0.89

0.75

0.72

Average

The last three columns of Table 6.7, from left to right, present the average of the values of $\hat{p} = 100(\hat{\lambda}_{CR} - 1)$, for one, two, and three null restrictions of five simulated selection cycles. According to these results, CRLGSI efficiency was 53.78%, 78.25%, and 61.25% higher than RLGSI efficiency. Thus, for this data set, the CRLGSI was a better predictor of the net genetic merit than the RLGSI.

6.4 The Combined Predetermined Proportional Gains Linear Genomic Selection Index

In the PPG-LPSI described in Chap. 3, the vector of the PPG (predetermined proportional gains) was $\mathbf{d}' = \begin{bmatrix} d_1 & d_2 & \dots & d_r \end{bmatrix}$. However, because the combined predetermined proportional gains LGSI (CPPG-LGSI) uses phenotypic and GEBV information jointly to predict the net genetic merit, the vector of the PPG (\mathbf{d}_{C}) should twice standard be the vector ď. that is, $\mathbf{d}'_C = \begin{bmatrix} d_1 & d_2 & \cdots & d_r & d_{r+1} & d_{r+2} & \cdots & d_{2r} \end{bmatrix}$, where we would expect that if d_1 is the PPG imposed on trait 1, then d_{r+1} should be the PPG imposed on the GEBV associated with trait 1, etc. In addition, in the CPPG-LGSI, we have three possible options for determining (for each trait and GEBV) the PPG, e.g., for trait 1, $d_1 = d_{r+1}$, $d_1 > d_{r+1}$, or $d_1 < d_{r+1}$. This is the main difference between the standard PPG-LPSI described in Chap. 3 and the CPPG-LGSI.

6.4.1 The Maximized CPPG-LGSI Parameters

It can be shown that the vector of coefficients of the CPPG-LGSI can be written as

$$\boldsymbol{\beta}_{CP} = \boldsymbol{\beta}_{CR} + \theta_{CP} \boldsymbol{\delta}_{CP}, \tag{6.16}$$

where

$$\theta_{\rm CP} = \frac{\beta_C' \Phi_C (\Phi_C' \widehat{\mathbf{T}}_C^{-1} \Phi_C)^{-1} \mathbf{d}_C}{\mathbf{d}_C' (\Phi_C' \widehat{\mathbf{T}}_C^{-1} \Phi_C)^{-1} \mathbf{d}_C}$$
(6.17)

is a proportionality constant. In addition, in Eq. (6.16), $\boldsymbol{\beta}_{CR} = \mathbf{K}_C \boldsymbol{\beta}_C$ is the vector of coefficients of the CRLGSI (Eq. 6.13), $\boldsymbol{\delta}_{CP} = \mathbf{T}_C^{-1} \boldsymbol{\Phi}_C (\boldsymbol{\Phi}_C' \widehat{\mathbf{T}}_C^{-1} \boldsymbol{\Phi}_C)^{-1} \mathbf{d}_C$, $\boldsymbol{\Phi}_C' = \mathbf{U}_C' \boldsymbol{\Psi}_C$, and $\boldsymbol{\beta}_C = \mathbf{T}_C^{-1} \boldsymbol{\Psi}_C \mathbf{a}_C$ (the vector of coefficients of the CLGSI). When $\boldsymbol{\theta}_{CP} = 0$, $\boldsymbol{\beta}_{CP} = \boldsymbol{\beta}_{CR}$, and if $\boldsymbol{\theta} = 0$ and \mathbf{U}_C' is the null matrix, then $\boldsymbol{\beta}_{CR} = \boldsymbol{\beta}_C$.

Thus, the CPPG-LGSI is more general than the CRLGSI and the CLGSI, and includes the latter two indices as particular cases. In addition, it can be shown that the CPPG-LGSI has the same properties as the PPG-LPSI described in Chap. 3.

The maximized selection response and the expected genetic gain per trait of the CPPG-LGSI can be written as

$$R_{CP} = \frac{k_I}{L_I} \sqrt{\beta_{CP}' \mathbf{T}_C \beta_{CP}}$$
(6.18)

and

$$\mathbf{E}_{CP} = \frac{k_I}{L_I} \frac{\boldsymbol{\Psi} \boldsymbol{\beta}_{CP}}{\sqrt{\boldsymbol{\beta}_{CP}' \mathbf{T}_C \boldsymbol{\beta}_{CP}}},\tag{6.19}$$

respectively. Although in the RLGSI and the PPG-LGSI the interval between selection cycles is denoted as L_G , in the CPPG-LGSI it is denoted as L_I . This is because the RLPSI and the CPPG-LGSI should have the same interval between selection cycles because they use phenotypic information to predict the net genetic merit.

6.4.2 Numerical Examples

Similar to the CRLGSI, to illustrate the CPPG-LGSI results we use the real training maize (*Zea mays*) F_2 population with 248 genotypes, 233 molecular markers, and three traits—GY (ton ha⁻¹), EHT (cm), and PHT (cm)—where $\widehat{\mathbf{P}} = \begin{bmatrix} 0.45 & 1.33 & 2.33 \\ 1.33 & 65.07 & 83.71 \\ 2.33 & 83.71 & 165.99 \end{bmatrix}$, $\widehat{\mathbf{C}} = \begin{bmatrix} 0.07 & 0.61 & 1.06 \\ 0.61 & 17.93 & 22.75 \\ 1.06 & 22.75 & 44.53 \end{bmatrix}$, and $\widehat{\mathbf{\Gamma}} = \begin{bmatrix} 0.07 & 0.65 & 1.05 \\ 0.65 & 10.62 & 14.25 \\ 1.05 & 14.25 & 26.37 \end{bmatrix}$ were the estimated matrices of \mathbf{P} , \mathbf{C} , and $\mathbf{\Gamma}$

respectively.

We can obtain the estimated CPPG-LGSI vector of coefficients as $\hat{\boldsymbol{\beta}}_{CP} = \hat{\boldsymbol{\beta}}_{CR} + \hat{\boldsymbol{\theta}}_{CP}\hat{\boldsymbol{\delta}}_{CP}$ (Eq. 6.16). Suppose that we restrict trait GY and its associated GEBV with matrix $\mathbf{U}_{C_1}' = \begin{bmatrix} 1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 \end{bmatrix}$ and the vector of predetermined restriction $\mathbf{d}_C' = \begin{bmatrix} 7 & 3.5 \end{bmatrix}$. In Sect. 6.3.2, we showed that the estimated CRLGSI vector of coefficients was $\hat{\boldsymbol{\beta}}_{CR}' = \begin{bmatrix} 0.076 & -0.004 & -0.018 & 2.353 & -0.096 & -0.082 \end{bmatrix}$; then, we only need to calculate $\hat{\boldsymbol{\theta}}_{CP}$ and $\hat{\boldsymbol{\delta}}_{CP}$ to obtain the vector of coefficients $\hat{\boldsymbol{\beta}}_{CP}$.

Let $\mathbf{w}' = \begin{bmatrix} 5 & -0.1 & -0.1 & 0 & 0 \end{bmatrix}$ be the vector of economic weights. It can be shown that $\hat{\theta}_{CP} = 0.00030$ is the estimated value of the proportionality constant and $\mathbf{\delta}'_{CP} = \begin{bmatrix} 0.56 & -77.28 & 40.89 & 49.44 & 77.28 & -40.89 \end{bmatrix}$. Thus, the estimated CPPG-LGSI vector of coefficients was $\hat{\boldsymbol{\beta}}'_{CR} = \begin{bmatrix} 0.76 & -0.030 & -0.004 & 2.369 & -0.070 & -0.096 \end{bmatrix}$, whence the estimated CPPG-LGSI can be written as

$$\widehat{I}_{CP} = 0.076 \text{GY} - 0.03 \text{EHT} - 0.004 \text{PHT} + 2.369 \text{GEBV}_{\text{GY}} - 0.070 \text{GEBV}_{\text{EHT}} - 0.096 \text{GEBV}_{\text{PHT}},$$

where GEBV_{GY}, GEBV_{EHT}, and GEBV_{PHT} are the GEBVs associated with traits GY, EHT, and PHT respectively. The same procedure is valid for two or more restrictions. Note that because $\hat{\theta}_{CP} = 0.0003$ is very small, the estimated CPPG-LGSI and CRLGSI values were very similar.

Figure 6.4 presents the frequency distribution of the estimated CPPG-LGSI values for one (Fig. 6.4a) and two predetermined restrictions (Fig. 6.4b) using $\begin{bmatrix} 1 & 0 & 0 & 0 & 0 \end{bmatrix}$

matrices
$$\mathbf{U}_{C_1}'$$
 and $\mathbf{U}_{C_2}' = \begin{bmatrix} 1 & 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 \\ 0 & 0 & 0 & 0 & 1 & 0 \end{bmatrix}$, the vectors of the PPG

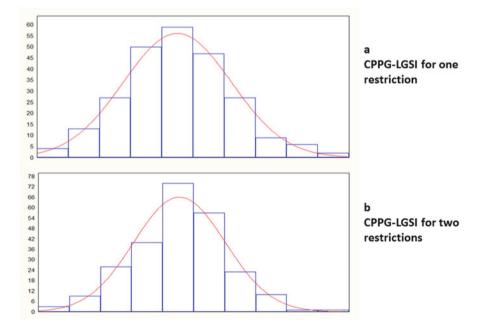


Fig. 6.4 Distribution of 244 estimated combined predetermined proportional gain linear genomic selection index (CPPG-LGSI) values with one (a) and two (b) predetermined restrictions, d = 7 and $\mathbf{d}' = \begin{bmatrix} 7 & -3 \end{bmatrix}$ respectively, obtained in a real training population for one selection cycle in one environment

 $\mathbf{d}_{C1}' = \begin{bmatrix} 7 & 3.5 \end{bmatrix}$ and $\mathbf{d}_{C2}' = \begin{bmatrix} 7 & -3 & 3.5 & -1.5 \end{bmatrix}$, and the real data set F₂. For both restrictions, the frequency distribution of the estimated CPPG-LGSI values approaches normal distribution.

Suppose a selection intensity of 10% ($k_I = 1.755$) and that we restrict trait GY and its associated GEBV. The estimated CPPG-LGSI selection response and expected genetic gain per trait were $\hat{R}_{CP} = k_I \sqrt{\hat{\beta}'_{CP} \hat{T}_C \hat{\beta}_{CP}} = 0.98$ and $\hat{E}'_{CP} = k_I \frac{\hat{\beta}'_{CP} \hat{\Psi}}{\sqrt{\hat{\beta}'_{CP} \hat{T} \hat{\beta}_{CP}}}$ = $[0.007 - 3.647 - 5.760 \ 0.004 - 2.829 - 4.711]$ respectively, whereas

= $\begin{bmatrix} 0.007 & -3.647 & -5.760 & 0.004 & -2.829 & -4.711 \end{bmatrix}$ respectively, whereas the estimated CPPG-LGSI accuracy was $\hat{\rho}_{HI_{CP}} = \frac{\hat{\sigma}_{I_{CP}}}{\hat{\sigma}_{H}} = 0.52$. Once again, because $\hat{\theta}_{CP} = 0.0003$, the latter results are very similar to the CRLGSI results.

Now, we use the simulated data described in Chap. 2, Sect. 2.8.1, to compare CPPG-LGSI efficiency versus PPG-LGSI efficiency. The criteria for this comparison are the Technow inequality (Chap. 5, Eq. 5.18) and the ratio of CPPG-LGSI accuracy ($\rho_{HI_{CP}}$) to PPG-LGSI accuracy (ρ_{HI_P}) expressed as percentages (Chap. 5, Eq. 5.17), $\hat{p} = 100(\hat{\lambda}_{CP} - 1)$, where $\hat{\lambda}_{CP} = \hat{\rho}_{HI_{CP}}/\hat{\rho}_{HI_P}$ for one, two, and three null restrictions in five simulated selection cycles.

Table 6.8 presents the estimated CPPG-LGSI heritability (\hat{h}_{I}^{2}), the estimated PPG-LGSI accuracy ($\hat{\rho}_{HI_{CP}}$), values of $W_{CP} = \frac{\hat{\rho}_{HI_G}}{\hat{h}_{I}}L_{I}$ ($L_{I} = 4$) and $\hat{p} = 100(\hat{\lambda}_{CP} - 1)$, where $\hat{\lambda}_{P} = \hat{\rho}_{HI_{CP}}/\hat{\rho}_{HI_{P}}$ and $\hat{\rho}_{HI_{P}}$ is the estimated CPPG-LGSI accuracy, for one, two, and three null restrictions in five simulated selection cycles. The averages of the estimated W_{CP} values for one, two, and three predetermined restrictions were 3.60, 3.31, and 2.50 respectively, whereas the PPG-LGSI interval length was 1.5 ($L_{G} = 1.5$). This means that the estimated Technow inequality, $L_{G} < \frac{\hat{\rho}_{HI_{G}}}{\hat{h}_{I}}L_{I}$, was true. Thus, for this data set, PPG-LGSI efficiency is greater than CPPG-LGSI efficiency in terms of time.

The last three columns of Table 6.8, from left to right, present the values of $\hat{p} = 100(\hat{\lambda}_{CP} - 1)$, for one, two, and three null restrictions in five simulated selection cycles. The average values of $\hat{p} = 100(\hat{\lambda}_{CP} - 1)$ for each of the three restrictions, in percentage terms, were 37.19%, 32.82%, and 37.08% respectively. This means that the CPPG-LGSI efficiency was greater than PPG-LGSI efficiency at predicting the net genetic merit.

three null restrictions for fiv	ictions for fiv	re simulated s	ve simulated selection cycles	es								
	CPPG-LGSI	I heritability		PPG-LGSI accuracy	accuracy		Values of W _{CP}	W _{CP}		Values of \hat{p}		
Cycle	-	2	3	1	2	3	1	2	3	1	2	3
-	0.41	0.41	0.85	0.65	0.91	0.85	6.25	5.68	3.69	24.62	-3.30	-12.94
2	0.75	0.86	0.84	0.72	0.70	0.55	3.33	3.02	2.40	25.00	24.29	38.18
n	0.78	0.85	0.81	0.70	0.67	0.55	3.17	2.91	2.44	30.00	32.84	40.00
4	0.78	0.84	0.82	0.61	0.59	0.46	2.76	2.57	2.03	49.18	49.15	56.52
S	0.80	0.83	0.82	0.56	0.54	0.44	2.50	2.37	1.94	57.14	61.11	63.64
Average	0.70	0.76	0.83	0.65	0.68	0.57	3.60	3.31	2.50	37.19	32.82	37.08

 $(\widehat{\rho}_{H_{Cp}})$, values of $W_{Cp} = \frac{\widehat{\rho}_{H_C}}{\widehat{h}_I} L_I (L_I = 4)$, and $\widehat{p} = 100(\widehat{\lambda}_{Cp} - 1)$, where $\widehat{\lambda}_P = \widehat{\rho}_{H_{Cp}}/\rho_{H_P}$ and $\widehat{\rho}_{H_P}$ is the estimated CPPG-LGSI accuracy, for one, two, and **Table 6.8** Estimated combined predetermined proportional gain linear genomic selection index (CPPG-LGSI) heritability (\hat{h}_1^2) , estimated PPG-LGSI accuracy

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