

How and why to measure the germination process?

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(received: August 18, 2005; accepted: January 26, 2006)

ABSTRACT – (How and why to measure the germination process?). In the last two centuries, papers have been published including measurements of the germination process. High diversity of mathematical expressions has made comparisons between papers and some times the interpretation of results difficult. Thus, in this paper is included a review about measurements of the germination process, with an analysis of the several mathematical expressions included in the specific literature, recovering the history, sense, and limitations of some germination measurements. Among the measurements included in this paper are the germinability, germination time, coefficient of uniformity of germination (*CUG*), coefficient of variation of the germination time (*CV_t*), germination rate (mean rate, weighted mean rate, coefficient of velocity, germination rate of George, Timson's index, *GV* or Czabator's index; Throneberry and Smith's method and its adaptations, including Maguire's rate; *ERI* or emergence rate index, germination index, and its modifications), uncertainty associated to the distribution of the relative frequency of germination (*U*), and synchronization index (*Z*). The limits of the germination measurements were included to make the interpretation and decisions during comparisons easier. Time, rate, homogeneity, and synchrony are aspects that can be measured, informing the dynamics of the germination process. These characteristics are important not only for physiologists and seed technologists, but also for ecologists because it is possible to predict the degree of successful of a species based on the capacity of their harvest seed to spread the germination through time, permitting the recruitment in the environment of some part of the seedlings formed.

Key words - germinability, germination rate, germination synchrony, germination time, germination uniformity

RESUMO – (Como e por que medir o processo de germinação?). Nos dois últimos séculos, trabalhos têm sido publicados incluindo medidas do processo de germinação. A diversidade das expressões matemáticas existente na literatura tem tornado difícil comparações entre os trabalhos e, algumas vezes, a interpretação dos resultados. Assim, este trabalho apresenta uma revisão sobre medidas de germinação, incluindo a análise de várias expressões matemáticas contidas na literatura específica. Entre as medidas analisadas estão a germinabilidade, tempo de germinação, coeficiente de uniformidade de germinação (*CUG*), coeficiente de variação do tempo de germinação (*CV_t*), velocidade de germinação (velocidade média, velocidade média ponderada, coeficiente de velocidade, velocidade de germinação de George, índice de Timson, *GV* ou índice de Czabator; método de Throneberry e Smith e suas adaptações, incluindo a velocidade de Maguire; *ERI* ou índice de velocidade de emergência, índice de germinação e suas modificações), incerteza associada à distribuição das frequências relativas de germinação (*U*) e índice de sincronia (*Z*). Os limites dessas medidas foram incluídos para facilitar a interpretação e a tomada de decisões durante as comparações. Tempo, velocidade, homogeneidade e sincronia são aspectos que podem ser medidos, e que informam a dinâmica do processo de germinação. Essas características são interessantes não somente para fisiólogos e tecnologistas de sementes, mas também para ecólogos, uma vez que é possível prever o grau de sucesso das espécies, com base na capacidade da safra de sementes em distribuir a germinação através do tempo, permitindo o recrutamento no ambiente de parte das plântulas formadas.

Palavras-chave - germinabilidade, sincronia de germinação, tempo de germinação, uniformidade de germinação, velocidade de germinação

Introduction

During the last two centuries, several methods and mathematical expressions to measure the germination

process have been proposed or discussed (Czabator 1962, Nichols & Heydecker 1968, Goodchild & Walker 1971, Heydecker 1973, Labouriau 1983a, Scott *et al.* 1984, Brown & Mayer 1988, Hilhorst & Karssen 1988, Carneiro *et al.* 1993, Bewley & Black 1994, Santana & Ranal 2004). It is wholesome and welcome, but the diversity of expressions has made comparisons difficult. Several of these expressions have the same name, measuring different characteristics of the germination

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process, some of them have the same meaning, although expressed using different symbols, while different mathematical expressions are used to measure the same characteristic. Besides, some practical attempts were also proposed to analyse the germination process, associating rate and capacity in generic expressions. As pointed out by Bewley & Black (1994), the attempts to incorporate two distinct characteristics of the germination process in the same mathematical expression are not entirely satisfactory.

Despite this diversity, several papers have discussed results based only on germinability and the behaviour of seeds during the germination process has several quantitative aspects that must be considered (Labouriau 1983a, Brown & Mayer 1988, Bewley & Black 1994). Time, rate, homogeneity, and synchrony are important aspects that can be measured, informing the dynamics of this process. These characteristics are important not only for physiologists and seed technologists, but also for ecologists because it is possible to predict the degree of success of a species based on the capacity of their harvest seed to spread the germination through time, permitting the recruitment in the environment of some part of the seedlings formed.

Thus, in this review the history, the sense, and limitations of some germination measurements were recovered and discussed. Mathematical expressions were adequate to the actual notation.

Measurement of germination capacity

As pointed out by Bewley & Black (1994), the term germination is often used loosely and sometimes incorrectly, being confounded with seedling growth, which begins when germination finishes. The global germination process is constituted by three partial processes, including the imbibition, the activation process, and intra-seminal growth that is completed with embryo protrusion (Labouriau 1983a). In the practical sense, it is difficult to establish the beginning and the end of each partial process because molecular and cellular events are involved in a multicellular and complex tissue and the reactivation of the process is gradual. Thus, the precision in determining the exact moment of the beginning of the germination process by means of some molecular or cellular criterion is substituted by macroscopic criteria, such as embryo protrusion, that confers some rigor because the same reference point can be used by several people. As was appropriately written by Bewley & Black (1994), the only stage of germination that can time fairly precisely

is its termination. In this manner, the establishment of an easy and common criterion was important because in the absence of this common point, the germination capacity could be under or overestimated, decreasing the chance of comparing results among laboratories and increasing contradictory information about the species.

The germination capacity of one seed, based on a binary answer (germinated/non germinated), is one qualitative attribute of the germination process, generally converted in a quantitative attribute, commonly percentage. According to Labouriau (1983a), germinability of one seed sample is the percentage of seeds in which the germination process reaches the end, in the experimental conditions, by means of the intra-seminal growth that results in the protrusion (or emergence) of one live embryo. In general, it is presented as percentage, accompanied by some dispersion measurement, but it is possible to use proportion rather than percentage, and the proportion values for one or several samples can be submitted to statistical tests (Carvalho *et al.* 2005). As the majority of papers include the germinability through time or at the end of the observations, it should be clear that its importance, meaning, and method of calculation are a consensus among researchers.

Measurements of germination time

The time for the first germination (t_o) that expresses the time of germination of the faster seeds has been used by some authors since 1865 (Edwards 1932, Czabator 1962, Goloff & Bazzaz 1975, Labouriau 1983a, Ranal 1999, Quintanilla *et al.* 2000). According to Labouriau (1983a), when used alone, it is an excessive simplification, since this method disregards the behaviour of the majority of seeds of one sample. The use of the two extreme times, time for the first germination (t_o) and time for the last germination (t_g) was proposed in 1878 (Edwards 1932), perhaps to correct the partiality expressed when only t_o is used, although this method is also limited, concentrating its attention only on the slower and faster seeds (Labouriau 1983a).

A more efficient method to solve this inconvenience is to use measurements of central tendency. Among them is the median, used in the context of germination since 1913 (Czabator 1962, Thompson 1970, Simon *et al.* 1976, Scott *et al.* 1984, Drake 1993, Ranal 1999) to study the time for 50% of germination ($t_{1/2}$). If the frequency distribution of germination is symmetric, the mean time and the median time can be used to express

central tendency of the data. On the other hand, if distribution is asymmetric, mean and median are distant, with the mean located in the tails of distribution where there are smaller frequencies and, in this case, the median is more representative than the mean, as discussed by Costa Neto (1977). The modal time is the point of the distribution in which the highest frequency of germinated seeds is observed. It represents the peak of germination, although sometimes it is not unique. There are seed samples whose distributions present two or more peaks of germination (germination with polymodal time) and, in this case, this measurement is inadequate to synthesize the information.

Labouriau and collaborators used in their papers the mean germination time (\bar{t}), proposed by Haberlandt in 1875 (Labouriau 1983a). It is calculated as the weighted mean of the germination time. The number of seeds germinated in the intervals of time established for data collection is used as weight. In this case, the use of the weighted mean is indispensable, since it considers that a different number of seeds germinate each time. In this manner, $\bar{t} = \frac{\sum_{i=1}^k n_i t_i}{\sum_{i=1}^k n_i}$, where t_i : time

from the start of the experiment to the i^{th} observation (day or hour); n_i : number of seeds germinated in the time i (not the accumulated number, but the number correspondent to the i^{th} observation), and k : last time of germination. This measurement was mentioned by Czabator (1962) as mean length of incubation time. According to the author, it is a measurement of the average length of time required for maximum germination of a seed lot, and is expressed in terms of the same units of time used in making germination counts (hours or days). It was denoted as

$MLIT = (G_1 T_1 + G_2 T_2 + \dots + G_n T_n) / (G_1 + G_2 + \dots + G_n)$, where G : germination count on any counting period; T : time. For Czabator (1962), the expression of $MLIT$ apparently has been used since 1875. Probably it was used by Haberlandt since this year as the mean germination time mentioned by Labouriau (1983a). The variance of germination time is given by

$s_t^2 = \frac{\sum_{i=1}^k n_i (t_i - \bar{t})^2}{(\sum_{i=1}^k n_i - 1)}$, where \bar{t} : mean germination time; t_i : time between the start of the experiment and the i^{th} observation (day or hour); n_i : number of seeds germinated in the time i , and k : last time of germination.

The standard deviation is calculated by $s_t = \sqrt{s_t^2}$. The standard error of the mean germination time is another important measurement of variability and is adequate to

measure the accuracy of the calculation of the mean germination time. Thus, $s_{\bar{t}} = s_t / \sqrt{\sum_{i=1}^k n_i}$, where s_t : standard deviation of the germination time; n_i : number of seeds germinated in the time i , and k : last time of germination. Labouriau and collaborators used the values of the confidence intervals to measure the dispersion of mean germination time and to visualize statistical differences between samples or treatments. For the confidence interval the expression is $\bar{t} \pm e$, being $e = t_{\alpha/2, v} s_{\bar{t}}$, where \bar{t} : mean germination time; t : critical value of the "Student" distribution for the significance $\alpha/2$ and $v = \sum_{i=1}^k n_i - 1$ degrees of freedom, and $s_{\bar{t}}$: standard error of the mean germination time.

Edmond & Drapala (1958) used an expression denoted by them as mean days for emergence and expressed by $Mdays = (N_1 G_1 + N_2 G_2 + \dots + N_n G_n) / (G_1 + G_2 + \dots + G_n)$, where G_1, G_2, \dots, G_n : number of seedlings emerged on the day of observation and N_1, N_2, \dots, N_n : number of days counted since the day of sowing until the day of observation. This expression is the mean germination time with different notation. If this expression were to be written in summation, it would be given by $Mdays = \frac{\sum_{i=1}^n N_i G_i}{\sum_{i=1}^n G_i}$, that is the mean germination time, used to evaluate seedling emergence.

Measurements of germination rate

Among the germination measurements, rate has received especial attention of the researchers. According to Labouriau (1983a), the first idea of germination rate was expressed by Kotowski (1926). The author presented the coefficient of velocity for 17 cultivated species, using the expression as coefficient of velocity = $100 (A_1 + A_2 + \dots + A_x) / (A_1 T_1 + A_2 T_2 + \dots + A_x T_x)$, where A_1, A_2, \dots, A_x : number of seedlings counted on the first day, second day, and so on until the last day (x), and T_1, T_2, \dots, T_x : number of days between sowing and the first collection, between the sowing and the second collection, and so on until the last collection (x).

Czabator (1962) mentioned that this coefficient of velocity is the reciprocal of the mean length of incubation time ($MLIT$), whose quotient is multiplied by 100. As Kotowski (1926) wrote that the value of the coefficient of velocity increases when the number of seedlings increases, this information brought several authors to consider this characteristic as a negative point for this

coefficient, limiting its use. However, the number of seeds germinated (or seedlings emerged) is only a weight of this measurement. Thus, the coefficient of velocity is free from the influence of the number of seeds germinated (or seedlings emerged) in the samples or treatments. This mistake was noted by Brown & Mayer (1988) when they analyzed the Kotowski's coefficient of velocity and related methods. According to them, CV is really only the reciprocal of the average germination time, and so is independent of the extent of germination. These authors considered that the usefulness of CV is further limited by its failure to provide any information on the spread of germination.

Later, Nichols & Heydecker (1968) presented the coefficient of velocity of Kotowski as coefficient of velocity of germination, denoting it as

$$CVG = \left(\sum_{i=1}^k f_i / \sum_{i=1}^k f_i x_i \right) 100, \text{ where } f_i: \text{ number of seeds}$$

newly germinating on day i ; x_i : number of days from sowing, and k : last day of germination. Nichols & Heydecker (1968) also observed that the CVG is a reciprocal of the mean germination time. Then, Labouriau (1970) demonstrated that the coefficient of velocity of Kotowski (1926) could be used to calculate the mean germination rate as $\bar{v} = CV/100 = 1/\bar{t}$, where \bar{t} : mean germination time, and CV : coefficient of velocity. In this manner, the mean germination rate (\bar{v}) is defined as reciprocal of the mean germination time, since the mean rate increases and decreases with $1/\bar{t}$, not with \bar{t} (Labouriau 1983b). As the distribution of the reciprocal of time is unknown, the variance of the rate is approximated. Labouriau (1983b) presented in detail the development of the function $\phi(t) = 1/t$, according to the expansion in Taylor series, suggesting a correction to the expression of the variance of the germination rate given by Heydecker (1973). Thus, the variance proposed for rate is $s_v^2 = (\bar{v})^4 s_t^2$, where s_t^2 : variance of germination time and \bar{v} : mean germination rate (Labouriau 1983b).

Sometimes the sense of \bar{v} , a pure rate measurement without the interference of the germinability, is difficult to understand because it is a non-linear measurement of rate. It is similar to that used to measure velocity of chemical reactions (molecules per time unity). It seems compatible with germination process which is a result of a series of chemical reactions inside metabolic complex routes. In physics, frequency is the reciprocal of period. As period is the interval of time to complete one cycle, frequency is the reciprocal of time ($f = 1/T$, where T : period). It is the well known concept of hertz ($1/s = \text{Hz}$, where s : second) that means cycle per second

or the reciprocal of second (s^{-1}). In this sense, \bar{v} is a frequency, since it is the reciprocal of the mean germination time, being expressed in hour^{-1} or day^{-1} .

In the same way, it is possible to calculate the germination rate as the reciprocal of the median time (Went 1957, *apud* Labouriau 1983a). Thus, $v_{1/2} = 1/t_{1/2}$, where $t_{1/2}$: time for 50% of germination. The use of this expression is not common because its interpretation is more difficult than the mean. Moreover, the variance and the standard deviation calculated in relation to the median present higher values than when they are calculated in relation to the mean. According to Labouriau (1983a), this inconvenience for the use of median was pointed out by Dixon & Massey (1969).

Another alternative to measure the germination rate is to calculate the weighted mean germination rate, using the mean germination rate of each replication of one treatment (or sample) and its variance (Labouriau,

1983c). It can be denoted as $\bar{v} = \frac{\sum_{j=1}^r w_j \bar{v}_j}{\sum_{j=1}^r w_j}$, for

$w_j = n_j / s_j^2$, where w_j : weight of the j^{th} replication; \bar{v}_j : mean germination rate of the j^{th} replication; n_j : number of seeds germinated in the j^{th} replication; s_j^2 : variance of the germination rate of the j^{th} replication, and r : number of replications or sample size (n). This weighted mean germination rate is more representative than the unweighted rate (\bar{v} or \bar{v}_j), since it considers and detaches differences between equal mean rate that present different variances (Labouriau & Osborn 1984). Although \bar{v} is a single mean, unaccompanied by dispersion values, it can be used for comparisons because its mathematical expression includes the variances of the samples or replications involved in its calculation. Thus, its use in the last decades is justifiable, once the statistics programs for computers were scarce. As presently data are more easily processed, the values of \bar{v} for the replications of each treatment can be submitted to the conventional multisample and pairwise comparisons.

Mixed measurements of germination rate

Timson (1965) suggested a measurement named by him as "ten summation" (Σ^{10}). According to the author, this measurement was defined based on the number of days required for seeds of weed species to complete the germination process. In his proposal Timson (1965) presented only a theoretical explanation without any mathematical expression. According to Nichols & Heydecker (1968), this index was originally proposed

by George in 1961 as “germination rate”. Goodchild & Walker (1971) denoted this index as $T = \sum_{i=1}^t g_i(t-j)$, where g_i : germination in time interval i (i varying from 1 to t); t : total number of time intervals, being $j = i - 1$. Goodchild & Walker (1971) emphasized that T cannot be of universal application. As it combines germination rate with the final germination percentage, it is useful for comparisons only when samples have equal germinabilities. Brown & Mayer (1988) wrote clearly that Timson (1965) proposed an index of germination based on the progressive total of daily cumulative germination over 10 days, with supplementary totals over other periods being quoted as Σ^5 for five days, Σ^{20} for 20 days, and so on. The same understanding was given by Baskin & Baskin (1998) who expressed this index as Σ^n , where n is the cumulative daily germination percentage for each day of the study. It means that $\Sigma^n = \sum g_i(t-j)$, i. e., in the first term of this equality the germinability (n) is cumulative as written by Baskin & Baskin (1998) and in the second term the germinability is partial (g_i) as written by Goodchild & Walker (1971). These two expressions, producing the same value, can carry doubts in relation to the use of partial or cumulative germinabilities. According to Tucker & Wright (1965), George proposed a method which involves the sum of the products between the number of seeds germinated and days remaining until completion of the germination test. This method produces a number ranging from NK down to N , where N is the number of seeds included in the test, and K the number of days until the end of the test. Timson (1965) changed the number of seeds germinated proposed by George per percentage of germination. Certainly, the use of percentage of germination is better than number of seeds germinated because the transformation to percentage cancels the effect of the sample size. Allan *et al.* (1962) described a measurement, denoted by them as *ERI*, that is exactly the description of the George method. Labouriau (personal communication) proposed to divide both members of the expression denoted by Goodchild & Walker (1971) as Timson index, by the sum of partial germinabilities (g_i); thus, $T_{\text{mod}} = T / \sum_{i=1}^t g_i$. This proposition minimizes the effect of the final germination percentage and then the Timson index can measure the germination rate with some precision.

Czabator (1962) proposed other index named by him as “germination value” and expressed as $GV = PV \cdot MDG$, where PV : peak value of germination,

and MDG : mean daily germination, which is the mean number of seeds germinating per day. The value of PV represents the maximum cumulative germination percentage divided by the number of days to reach this percentage. In the interpretation of Brown & Mayer (1988), this index is an expression of speed and totality of germination, and their interaction. According to these authors, the values of PV and MDG can be counterbalanced, resulting in equal values for GV for samples or treatments with different behaviour in relation to germination process.

Maguire (1962) presented the germination rate for field or laboratory conditions to evaluate the seedlings vigour. According to the author, the method is an adaptation of the daily counting method used by some researchers, like Throneberry & Smith (1955), for determination of the seedlings vigour. In fact, Maguire (1962) gave a generic mathematical expression for the method presented by Throneberry & Smith (1955) that permits one to obtain the measurement for any intervals of time. The mathematical expression is given by

$$\text{Rate} = \frac{\text{number of normal seedlings}}{\text{days to first count}} + \dots + \frac{\text{number of normal seedlings}}{\text{days to final count}}$$

High values obtained using this expression mean higher seedling vigour of one sample in relation to another. In seed technology this value, named index of velocity of germination or emergence, is used to predict the relative vigour of samples, especially for cultivated species, since samples with the same quantity of seeds germinated can present different values for this index. Although Maguire (1962) had not presented the unit of this measurement, the value calculated using the expression proposed denotes a number of normal seedlings per day.

According to Brown & Mayer (1988), the Maguire’s index is a time-weighted cumulative germination that measures the speed of germination and quantifies the seedling vigour, being expressed as

$$T_{\text{mod}} = T / \sum_{i=1}^t g_i$$

where G_i : percentage of seedlings that germinated during the time interval t_i and n : last day of germination. In their comments, the authors inform that there is confusion about what this expression measures, since the expression is not arithmetically equivalent to any accepted measurement of germination rate. The use of percentage of seedlings proposed by Bouton and collaborators in 1976, as commented by Brown & Mayer (1988), is important because it cancels the effect of the sample size. In spite of this modification, the value obtained using this expression is influenced by the total number of seeds that germinated in the experimental

conditions. Thus, it is useful for comparisons only when samples or treatments present the same germinabilities.

Another modification of the method presented by Throneberry & Smith (1955) was presented by Kendrick & Frankland (1969) that proposed a germination index \bar{g} for comparisons of germination times, expressed as $\bar{g} = \Sigma(1/t)$ where t : time for each seed to germinate. Apparently this expression does not have any relation to the method presented by Throneberry & Smith (1955) and Maguire (1962). Nevertheless, when $n_1, n_2, n_3, \dots, n_k$ seeds germinated on the first day, second, third, and k^{th} days, the expression could be written as $\bar{g} = \sum_{i=1}^k n_i / t_i$ that is in itself the Throneberry and Smith's method, adopted by Maguire (1962).

The emergence rate index proposed by Shmueli &

Goldberg (1971) is given by $ERI = \sum_{n=n_o}^{n=c-1} X_n(c-n)$, where

X_n : number of emerged seedlings per linear meter, counted on the day n ; c : number of days counted from sowing until the end of the emergence; n : day of the counting, expressed as number of days after sowing, and n_o : day when the emergence started, expressed as number of days after sowing. Allan *et al.* (1962) used this name and abbreviation to nominate another mathematical expression, as was commented previously. The value of ERI can be divided by total number of emerged seedlings (or germinated seeds), to minimize the effect of germinability (Santana & Ranal 2004).

Then, the new expression is $ERI_{\text{mod}} = ERI / \sum_{n=n_o}^{n=c-1} X_n$.

The germination index proposed by Melville *et al.* (1980) was expressed by $GI = \sum_{i=1}^n |(15 - D_i)G_i| / S$, where n : number of germination counting (days, hours or other type of time unit); 15 : total number of days spent in the germination test plus 1; D_i : number of days until the i^{th} reading; G_i : number of normal seeds germinated in the i^{th} day, and S : total number of seeds used in the test. It means that non germinated seeds are included in the calculus. The S value of the GI expression, index proposed by Melville *et al.* (1980), can be replaced by the number of seeds germinated at the end of the experiment to cancel the effect of the seed sample size (Santana & Ranal 2004).

General considerations about rate germination measurements

One important aspect in relation to these rate measurements is that when the quantity of seeds that

completed the germination process is similar among treatments, all rate measurements have similar accuracy. On the contrary, when germinabilities are different among treatments, high values of G increase values of all mixed rate measurements because they include germination capacity and rate in the same mathematical expression. Negligible to low correlation (*sensu* Davis 1971) between \bar{v} and germinability, obtained by simulations on a set data show that the first is not influenced by the second. Very high correlation *sensu* Davis (1971) between germinability and each one of the mixed rate measurements obtained for the same set data show the influence of the germinability on these measurements. It means that germinability and germination rate are counterbalanced in these indices. These associations were demonstrated for the first time for *Myracrodruon urundeuva* Allemão diaspores (Dorneles *et al.* 2005). As pointed out by Goodchild & Walker (1971) and Brown & Mayer (1988), this inherent ambiguity is the greatest weakness in Timson's method. As the idea of these measurements is the same, this observation is valid for all mixed measurements. Among the rate measurements here discussed, $CV, CVG, v_{1/2}, \bar{v}, \bar{v}, GV, Rate, GR,$ and \bar{g} measure the germination frequency. Timson's index, $T_{\text{mod}}, ERI, ERI_{\text{mod}}, GI,$ and GI_{mod} do not have this characteristic and the germination frequency is substituted by the time in decreasing order. Thus, to the faster seeds a weight corresponding to the greater time is attributed.

Measurements of uniformity of germination

The coefficient of uniformity of germination measures the variability among seeds in relation to the mean germination time of the sample (Heydecker 1973, Bewley & Black 1994) and is expressed by

$$CUG = \frac{\sum_{i=1}^k n_i}{\sum_{i=1}^k (\bar{D} - D_i)^2 n_i};$$

being: $\bar{D} = 100 / CRG; CRG = \left(\frac{\sum_{i=1}^k n_i}{\sum_{i=1}^k D_i n_i} \right) 100$,

where n_i : number of seeds germinated on the i^{th} day and D_i : number of days counted from the day of sowing to the collection of the datum (i). It is important to note that CRG is the Kotowski's coefficient of velocity with a different mathematical notation, and \bar{D} the mean germination time. This mathematical expression is the reciprocal of the variance of the germination time, except the "-1" that appears in the expression of variance as the degrees of freedom of the measurement. If CUG was expressed as variance, high values would indicate germination spread over time. Certainly the author had

the intention that this expression would have a direct interpretation on the sense that high values would be associated with concentrated germination in time. Thus, it is important that it be expressed as a reciprocal measurement of variance, given information about the uniformity of seeds in relation to the germination process. This coefficient can be applied only if the germination frequencies have normal distribution (Heydecker 1973, Bewley & Black 1994). This is a limitation because, in general, seeds in germination process show some deviation from normality in their behaviour in relation to time. Another limitation for the use of this measurement is that the mean germination time of samples or treatments must be the same for comparisons. This is the same limitation associated with variances and standard deviations because the variability measured by them depends on the magnitude of the mean.

To solve this inconveniency, the coefficient of variation of the germination time was proposed to evaluate the germination uniformity or variability in relation to the mean germination time and was applied to seeds of *Myracrodruon urundeuva* Allemão (Dorneles *et al.* 2005) and *Anacardium humile* A. St.-Hil. (Carvalho *et al.* 2005). It is calculated as $CV_t = (s_t / \bar{t})100$, where s_t : standard deviation of the germination time and \bar{t} : mean germination time. The unit of CV_t (%) is easier to interpret than the unit of the coefficient of uniformity of germination (day^{-2}). This inconvenience of the *CUG* unit is the same related to variances, as commented above. Another important point is that the CV_t is a relative dispersion measurement that permits comparisons, independently of the magnitude of the mean germination time.

Measurements of germination synchrony

As seed germination, in general, is assynchronized, it is possible to quantify this characteristic by means of a measurement named synchronization index (\bar{E}), uncertainty associated to the distribution of the relative frequency of germination (U), or informational entropy (H), expressed by Labouriau & Valadares (1976) as

$\bar{E} = -\sum_{i=1}^k f_i \log_2 f_i$, being $f_i = n_i / \sum_{i=1}^k n_i$, where f_i : relative frequency of germination; n_i : number of seeds germinated on the day i , and k : last day of observation. This expression was proposed by Shannon in 1948 to measure the informational entropy or uncertainty associated to the frequency distribution $f_i = \phi(t)$ (Labouriau 1983a). This expression is used by

ecologists to measure the diversity of one environment (Magurran 1988). Thus, high values for this index indicate high diversity and numbers in direction of zero indicate low diversity. When applied to seed germination, the conventional interpretation is in the opposite direction, that is, low values indicate more synchronized germination. The use and sense of this measurement for seed germination were clearly demonstrated by Labouriau & Pacheco (1978) for seeds of *Dolichos biflorus* L., using several frequency polygons with relative frequencies of germination as functions of the time of isothermal incubation of the seeds. The unit of this index is bit, since \log_2 is used (Labouriau & Valadares 1976, Labouriau 1983a, Magurran 1988). Each bit is a binary measurement that counts, in this case, germinated/non germinated. Thus, the transformation of the germination frequencies (f_i) for each time of observation (t_i) in the logarithm of the frequency in base 2 ($\log_2 f_i$) transform a binary measurement (bit) in a weigh. In this manner, $\log_2 f_i$ is the weigh of each frequency registered, being expressed as $f_i \log_2 f_i$. The maximum limit of the Shannon index is the log of the total number of species established in a given area (Hutcheson 1970). For seeds, this limit is the \log_2 of $\sum n_i$, that is, the \log_2 of the total number of seeds germinated in a given condition.

Z index was proposed to evaluate the degree of overlapping in flowering among individuals of one population. Adopting the idea expressed by Primack (1980), the synchrony of one seed with another included in the same replication of one treatment is defined as

$$Z = \sum C_{n_i, 2} / N, \quad \text{being } C_{n_i, 2} = n_i(n_i - 1) / 2 \quad \text{and}$$

$N = \sum n_i(\sum n_i - 1) / 2$, where $C_{n_i, 2}$: combination of the seeds germinated in the time i , two together, and n_i : number of seeds germinated in the time i . Then, $Z = 1$ when the germination of all seeds occur at the same time and $Z = 0$ when at least two seeds could germinate, one at each time.

U is an adaptation of the Shannon index that measures the degree of uncertainty in predicting the informational entropy or the uncertainty associated with the distribution of the relative frequency of germination (Labouriau & Valadares 1976, Labouriau 1983a). In function of this, uncertainty index appears to be better for denomination of this index, as used by Labouriau in his papers published before 1983, than synchronization index. Low values of U indicate frequencies with few

Table 1. Limits of the germination measurements. k : last day of germination; t : total number of time intervals; 1n : total number of seeds in germination condition; this maximum limit means that the maximum rate is reached when all seeds could germinate at the first evaluation; 2n : day of the counting, expressed as number of days after sowing; c : number of days counted from sowing until the end of the emergence; 3n : number of germination counting; 4n : total number of seeds germinated; 5 when time units are included, besides day, hour or other type of time unit could be used.

Measurement	Limits	Unit ⁵
Germinability	$0 \leq G \leq 100$	%
Mean germination time	$0 < \bar{t} \leq k$	day
Coefficient of variation of the time	$0 < CV_t \leq \infty$	%
Coefficient of uniformity of germination	$0 < CUG < \infty$	day ⁻²
Coefficient of velocity of germination	$0 < CVG \leq 100$	%
Mean germination rate	$0 < \bar{v} \leq 1$	day ⁻¹
Weighted mean rate	$0 < \bar{v} \leq 1$	day ⁻¹
Timson's index	$0 \leq T \leq 100 t$	% day
Timson's index modified	$0 < T_{\text{mod}} \leq t$	day
Czabator's index (germination value)	$0 \leq GV \leq 10,000$	% ² day ⁻²
Rate of Maguire ¹	$0 \leq Rate \leq n$	seed day ⁻¹ or seedling day ⁻¹
Emergence rate index ²	$0 \leq ERI \leq X_n (c - n)$	seed day or seedling day
Emergence rate index modified ²	$0 < ERI_{\text{mod}} \leq (c - n)$	day
Germination index ³	$0 < GI \leq n$	day
Germination index modified ³	$0 < GI_{\text{mod}} \leq n$	day
Germination rate	$0 \leq GR \leq 100$	% day ⁻¹
Germination index	$0 \leq \bar{g} \leq n$	seed day ⁻¹
Uncertainty ⁴	$0 \leq U \leq \log_2 n$	bit
Synchronization index	$0 \leq Z \leq 1$	unit less

peaks, that is, germination more concentrated in time. Only one seed germinating changes the value of U . It means that U measures the degree of spreading of germination through time and can be used, by inference, to measure the synchrony of germination. On the contrary, Z produces a number if and only if there are two seeds finishing the germination process at the same time. Thus, the value of Z measures, in fact, the synchrony of germination, that is, the degree of germination overlapping.

The mathematical limits of each expression here presented represent an important contribution and a good point of reference that make it easier to distinguish between good and bad treatments (table 1).

When zero exists for the germination measurements

Zero and null values are surrounded by doubts that make the statistical analysis difficult. Zero for germinability exists when none seed is able to complete the germination process in a given condition (table 1). The same is valid for Timson's index, GV , $Rate$, ERI , GR , and \bar{g} . Zero for \bar{t} , CV_t , CUG , CVG , \bar{v} , \bar{v} , T_{mod} ,

ERI_{mod} , GI , GI_{mod} , U , and Z do not exist, that is, the values are nulls when germination does not occur. Mathematically all fractional measurements must have as divisor a number different from zero. Thus, when germinability is zero, \bar{t} and \bar{v} must not be calculated. In addition, CV_t and CUG measure variability. It means that both of them exist only when at least two seeds were able to complete the germination process in the experimental condition, at different times. U is zero when one seed was able to complete the germination process. Z is zero when no overlapping will be observed for n germinated seeds and will be null when no seed could complete the germination process.

How to work statistically with germination measurements using samples

When few seeds are produced by one species, the conventional experimental designs are limited. According to the seed analysis rules (ISTA 1985, Brasil 1992), four hundred seeds per treatment are necessary to give representativity for the results. As this number is very high for several native species, Brazilian researchers are preparing specific rules for forest species based on

two hundred seeds as a reliable sample, although not always this minimum could be collected. In this case, all germination measurements with variability associated to them can be calculated because t Student test can be used to compare samples if they have normal distribution. This technique was applied to evaluate the emergence of *Anacardium humile* seedlings (Carvalho *et al.* 2005). The proportion of seeds germinated (or seedlings emerged) is a discrete variable, with binomial distribution and the method of the approximation of the binomial by normal distribution can be used, since $n_1 \hat{p}_1 > 5$, $n_1 \hat{q}_1 > 5$, $n_2 \hat{p}_2 > 5$, and $n_2 \hat{q}_2 > 5$, for $\hat{p}_1 = X_1/n_1$; $\hat{q}_1 = 1 - \hat{p}_1$; $\hat{p}_2 = X_2/n_2$, and $\hat{q}_2 = 1 - \hat{p}_2$, where \hat{p}_1 and \hat{p}_2 are the estimation of the proportion of seeds germinated in the first and second samples; X_1 and X_2 are the number of seeds germinated in the first and second samples; n_1 and n_2 are the initial number of seeds included in the first and in the second samples, respectively. If these conditions are met, comparisons between samples can be processed using the expression of the t Student test for variances heterogeneous to compare proportions,

$$t = (\hat{p}_1 - \hat{p}_2) / \sqrt{\frac{\hat{p}_1 \hat{q}_1}{n_1} + \frac{\hat{p}_2 \hat{q}_2}{n_2}}$$

For \bar{t} , t Student test can be processed by the

expression $t = (\bar{t}_1 - \bar{t}_2) / \sqrt{\frac{s^2_{t_1}}{n_1} + \frac{s^2_{t_2}}{n_2}}$, where \bar{t}_1 and \bar{t}_2 are the mean germination time of the first and second samples; $s^2_{t_1}$ and $s^2_{t_2}$ the variances of the first and second samples; and n_1 and n_2 the number of seeds germinated in the first and in the second samples, respectively. The same is valid for the mean germination rate (\bar{v}), using the expression of variance proposed by Labouriau (1983b),

in the expression $t = (\bar{v}_1 - \bar{v}_2) / \sqrt{\frac{s^2_{v_1}}{n_1} + \frac{s^2_{v_2}}{n_2}}$, where \bar{v}_1 and \bar{v}_2

are the mean germination rate of the first and second samples; $s^2_{v_1}$ and $s^2_{v_2}$ the variances of the first and second samples; and n_1 and n_2 the number of seeds germinated in the first and in the second samples, respectively.

Comparisons between samples for the uncertainty of the seed germination (U) can be made using the method proposed by Hutcheson (1970) and described by Zar (1999).

For the other germination measurements here presented, variability is unknown and its approximation was not proposed by the authors. Then, no statistical test for comparisons can be processed, using samples without replications.

How to work statistically with germination measurements within experimental designs

Within the conventional experimental designs, all germination measurements here discussed can be analysed. In a treatment with five replications, five values of each measurement can be used to calculate the mean and to process the multisample comparisons. In this case, the variability inherent in the germinability, mean germination time and rate, and uncertainty is not considered because only the variability of the experiment can be measured.

Relationships among germination measurements

When statistical analysis are processed, the results are not always the same for \bar{t} and \bar{v} in relation to normality, homogeneity, in the magnitude of the variances, and in relation to the number of categories formed after pairwise comparisons in one experiment. It means that both of them are non linearly associated. Mathematically \bar{t} and \bar{v} are proportionally inverse. Thus, when the first increases, the second obligatorily decreases, but non linearly. These relationships are also valid when the rate of the germination process is measured using mathematical expressions free of the action of the number of seeds germinated as T_{mod} , ERI_{mod} , and GI_{mod} . For mixed rate measurements, increases or decreases as a consequence of the balance between germinability and rate germination can give rise to associations without the same tendency observed for \bar{t} and \bar{v} .

High or low values of \bar{t} are not associated with high or low values of CV_t or CUG , neither with high or low values of U or Z , because seeds could germinate spending a little time (low \bar{t} value) and the germination process could be spread through time or could germinate spending more time (high \bar{t} value), concentrating the process after the first germination.

Another important point is related to the interval between two consecutive evaluations. Depending on this interval, seeds could be considered synchronized and homogeneous or unsynchronized and heterogeneous. Short intervals between two consecutive evaluations makes the determination of \bar{t} more accurate because the history of each seed is registered. On the other hand, short intervals between two consecutive evaluations, show low synchrony, because few seeds are counted at the same evaluation. It means that the mean germination

time and synchrony are overestimated using high intervals between observations. On the contrary, if short intervals are used to count germination, all seeds appear to be without synchrony because it is possible to detect the end of germination (embryo protrusion) for each seed individually. The germination process of *Gossypium hirsutum* L. cv. DeltaOPAL seeds were evaluated in different intervals (Cunha 2002) and confirmed the influence of the interval size on the accuracy of these measurements.

After these considerations, it is clear that the researcher must choose the measurements and the interval of evaluations, according to their objectives. Besides, correlations among the germination measurements prove the nature of each measurement and can help the researchers in their options. It is important to take into account that correlations between variables in completely randomized design models must to be done after the standardization of the effect of treatment (Santana & Ranal 2006). When correlations are applied to the residuals and to the original data, this makes it possible to identify if the association between variables (in this case, germination measurements) is algebraic, physiological, or due to the effect of treatment included in the model. The analysis of the germination can be enriched if in addition to the germinability, \bar{t} and/or \bar{v} , CV_t , Z and/or U values were communicated, because they measure different aspects of the germination process. Finally, all germination measurements could be applied to evaluate seed germination and seedling emergence. Details and intermediate tables to calculate all of them were presented by Santana & Ranal (2004) and the practical and biological sense by Dorneles *et al.* (2005).

Acknowledgments – Important assistance for the understanding and calculus of the Z index was given by Dr. Richard B. Primack and Dr. Fernando Pedroni. Useful discussions about \bar{v} as a frequency were established with Dr. Roberto Hiroki Miwa (Physicist) and Cláudio Franco Muniz (Master Degree Student). The English revision was done by Mr. John David Bagnall. The authors register their sincere thanks, adding a special word of thanks to the Editorial Board of the Revista Brasileira de Botânica for the invitation and opportunity to present and publish this review.

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