Germination and growth *Jatropha curcas* after storage and salt stress

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# Abstract

*Jatropha curcas* L. is a plant that can be used in the production of biofuel with strong resistance to drought. However, the crop presents two important problems: i) rapid loss of viability, resulting from the high respiratory rate of the seeds during the storage period; ii) seed sensitivity when germinated under salinity conditions. To achieve these objectives, two experiments were developed. In the first experiment were verified how the storage of seeds in a drier environment can influence germination, respiration rate and the main biochemical and physiological parameters. In the second experiment, five different genotypes (114, 133, 171, 183 and 218) originating from different producing regions of Brazil were studied under the effect of the addition of NaCl (0, 50, 75, 100 and 150 mM) in water irrigation. In this experiment, we evaluate the germination and initial growth. The results of the first experiment show that the use of desiccant, can stabilized the germinability of the seeds stored, a fact corroborated by the reduction of the water potential of the seeds and reduction of the respiratory rates. On the other hand, we showed that *J. curcas* presents a moderate tolerance to salinity, being able to germinate up to 150 mM NaCl, even though a drastic reduction in the biomass accumulation was observed with the increase of the salt concentration in the irrigation water. The results show that the germination was reduced to values close to 4% in the treatment of 150 mM, while the mean germination time was increased with the increase in the concentration of salts. The biometric and biomass componets were strongly affected by the increase of the salts, while in the biomass allocation parameters, accumulation was observed in the stem of the seedlings. In this sense, genotypes 114, 171 and 183 were shown to be potentially tolerant while genotypes 218 and 133 were sensitive.

**Key words:** salinity tolerance, seed germination, biomass, biofuel, NaCl

# Introduction

*Jatropha curcas* (purgint nut) is a species belonging to the family Euphorbiaceae with multiple uses, abundantly distributed in many tropical and subtropical regions in the Americas, Africa and Asia [[1](#ref-heller1996physic),[2](#ref-takeda1982development)]. Over the last 20 years they have gained a lot of attention as a potential crop for bioenergy production, since their seed oil can easily be converted to good quality biodiesel. In addition, the species does not present as edible and therefore does not compete with the other oilseeds [[3](#ref-Pompelli2011)]. It has a high growth rate, easy propagation, short period until the first fruit harvest, low seed cost [[1](#ref-heller1996physic)], high oil content (40-58%) [[4](#ref-pandey2012jatropha),[5](#ref-pompelli2010environmental)], and good adaptation to different agroclimatic conditions [[6](#ref-Divakara2010)–[8](#ref-gao2008effects)].

Germination is the process that determines when and where the seeds will initiate their growth [[9](#ref-gunster1994seed)], allowing the embryo to germinate and develop as a photosynthetically active organism. It begins with the imbibition of the quiescent seed and ends as the elongation of the embryonic axis, which can be visualized by the emergence of soil surface. At the moment, the reserves contained in the seeds begin to be mobilized by yielding energy to the developing embryo [[10](#ref-bewley2013mobilization),[11](#ref-sanchez2012early)]. In this sense, salinity can affect germination, limiting the absorption of water in the seeds (osmotic effect) [[12](#ref-almansouri2001effect),[13](#ref-hegarty1977seed)] increases the toxicity by ions or the combination of both [[14](#ref-apse1999salt)]. In addition, NaCl may affect the mobilization of reserves [[15](#ref-bouaziz1990consumption)], structural organization and protein synthesis in embryos [[16](#ref-alencar2015ultrastructural)]. In saline environments, plant adaptation during germination are decisive stages for species establishment, and such factors may negatively influence this process [[17](#ref-ungar1995seed)].

If we take into account that *J. curcas* is a potential species for the large-scale generation of biodiesel, it is easy to think that it will be necessary to plant hundreds or thousands of hectares of trees to produce a satisfactory amount for commercial exploitation [[18](#ref-contran2013state),[19](#ref-yang2010selection)]. In this sense, it is also salutary to remember that in times of harvest market prices usually fall a lot [[20](#ref-sumner1989harvest)] and that is where the storage of seeds comes in. However, seed storage is the most factor that negatively affects seed viability, which includes the time elapsed between harvesting and utilization [[21](#ref-marcos1984testes),[22](#ref-marcos1998new)]. Marcos-Filho [[22](#ref-marcos1998new)] describes that seed storage is a major problem for agriculture [[23](#ref-tekrony2006seeds)], since it is responsible for large losses worldwide, especially in the tropics, where high temperatures and high relative humidity prevail during seed maturation and storage [[24](#ref-bilia1994comportamento)]. *J. curcas* does not escape this pattern, since it presents high metabolism, causing its seeds to rapidly lose their viability with storage [[25](#ref-moncaleano2013germination)]. Although deterioration is irreversible and unavoidable, the speed of the process can be controlled by appropriate harvesting, drying and storage techniques [20]. In this sense, the use of drier atmosphere environments could protect seeds [[26](#ref-hay2012evaluation)–[28](#ref-rao2006storability)].

Another factor that negatively influences agriculture is salinity, mainly in irrigated crops [[29](#ref-kumar2008effects)], with NaCl being the predominant salt. Approximately 20% of the world's cultivated land is affected by salts [[30](#ref-sun2009nacl)]. This problem is more relevant in arid and semiarid regions, where the lack of rainfall and the high evaporative demand caused by high temperatures and low relative humidity contribute to soil salinity intensification. In addition, salinity affects plant growth and development [[31](#ref-munns2008mechanisms)], negatively influencing different stages of its development [[12](#ref-almansouri2001effect),[32](#ref-khajeh2003interaction),[33](#ref-khan2003light)]. However, throughout their evolution, plants have developed mechanisms for regulation and tolerance to salts.

Some studies have described the ecophysiological aspects of the tolerance of *J. curcas* to NaCl [[34](#ref-diaz2012tolerance)–[37](#ref-rajaona2012effect)]; However, these studies focused on only one genotype. Although interest in the growth response of *J. curcas* is increasing, there is no known research that has examined the effect of NaCl on different genotypes during germination and early development of seedlings. In this work five distinct genotypes of *J. curcas* exposed to different NaCl treatments were studied to determine tolerance and to understand the morphological and physiological responses of this species under conditions of salinity in their germination and initial development.

Thus, the main hypotheses of this work were (i) to verify if the use of a desiccant agent could help to maintain the viability and germinability of the seeds of *J. curcas* when stored for long periods of time, and (ii) to study the mechanims of the tolerance of *J. curcas* to salinity between the genotypes cultivated in Brazil.

# Materials and methods

**Aging tests.** To test the effect of storage on seed viability, an artificial aging test was used to reduce the water content in the interstices of the seeds with a desiccant material composed of silica gel (Sigma-Aldrish, part number 10087). The genotype used in this experiment was 171 from Maceió, AL, Brazil. In each experimental unit 50 seeds of *J. curcas*, arranged in germination boxes (110 x 110 x 35 mm) were added under a stainless steel mesh suspended 2 cm from the desiccant. Five storage times (i.e., 0, 3, 6, 9, 12 months) were tested and stored under refrigerator at 4 ± 2°C. After each storage period, the seeds were removed to the germination boxes with desiccant and then placed to germinate.

**Germination tests with aged seeds**. After removed from germination boxes with desiccant, the seeds were germinated in germination boxes (110 x 110 x 35 mm) containing two sheets of germination test paper soaked with 2x the weight of the paper in water and were sealed and placed in a growth chamber (mod. NT 708, New Technical Instruments, Piracicaba, SP, Brazil). The incubator was equipped with four cold white fluorescent lamps of 20 W with 40 μmoles m-2 s-1 at the level of the germination boxes. The photoperiod was 12 h and the temperature conditions were 25 ± 0.5°C. Germination were daily evaluated for a period of 25 days. It was considered germinated, the seed whose radicle has emerged from the integument.

**Biochemical analysis of the seeds used in the aging test.** A portion of 10% of the aged seeds was carefully ground in liquid nitrogen and stored at -20°C until use. For extraction of soluble carbohydrates (TSC), soluble amino acids (TSA) and starch (STR), samples were solubilized in 50% (v/v) ethanol [[38](#ref-trethewey1998combined)], whereas for analysis of total soluble proteins (TSP), the samples were extracted in Stitt buffer [[39](#ref-armengaud2009multilevel)]. The measures of soluble carbohydrates and start, soluble proteins and soluble amino acids the methodologies described by Dubois [[40](#ref-dubois1956colorimetric)], Bradford [[41](#ref-bradford1976rapid)] and Moore and Stein [[42](#ref-moore1954modified)] were used, respectively. For the quantification of the oil content (OIL) the methodology described in detail by Ahmad [[43](#ref-ahmad1981ricinoleic)]. To quantification of glucose (GLC), fructose (FTS) and sucrose (SCR) [[44](#ref-stitt198932)], coupled to the production of 6-phosphogluconate, in the sequential presence of hexokinase, phosphoglucoisomerase, glucose-6-phosphate dehydrogranase and invertase enzymes were used, as described in Stitt [[44](#ref-stitt198932)]. All these analyzes were performed in triplicate.

**Physiological analyzes coupled to the aging test.** The relative water content (RWC) of seeds was calculated as described Moncaleano-Escandon [[25](#ref-moncaleano2013germination)]. The water potential (WTP) of the seeds was quantified with the dewpoint water potential meter (WP4C; Decagon Devices, Pullman, WA, USA), where the seeds were lightly cracked to allow water to pass through the seeds to the internal environment. The values were obtained in MPa. For respiratory rate estimation (RPR), 5% of the seeds used in each storage period were inserted whole in a CO2 flow chamber (6400-09; LiCOR, Lincoln, NE, USA). For each storage time, 10 different samples were used as replicates. In each measurement procedure three cycles of 102 seconds were performed, with a 2 second interval between the readings. During this time the increase in CO2 concentration inside the chamber was monitored. The reference CO2 was calibrated, at each measurement, according to the CO2 concentration of the environment (~ 400 μmol CO2 m-2 s-1). The net respiration rate was expressed in μmol CO2 h-1 g-1 seed.

**Germination tests in the presence of NaCl.** For this experiment seeds of five genotypes (Table 1) of *J. curcas* originating from different producing regions of Brazil, given by EMBRAPA Agroenergia (Brasília, DF – Brazil), where kept at 4°C until their use. The germination was carried out in a greenhouse at the Federal University of Pernambuco, Department of Botany, Recife, PE (8°02"59.0' S; 34°56"54.9' W, 4 masl). The average temperature being recorded during the experiments using a portable mini climatic estation (mod. KR420, Akron Measure Instrument, Leuven, Belgium). In this experiment, five different concentrations of NaCl (0, 50, 75, 100 and 150 mM) were tested in the irrigation water, with zero being destiled water. After the soaking, all seeds were disinfected with NaOCl (2%) for 10 minutes and triply rinsed with distilled water. Thus, the seeds were germinated in polypropylene boxes (20 x 20 x 5 cm) containing 2,500 g of river washed sand, air-dried. Each boxes containing 25 seeds were considered as an experimental unit. The boxes were irrigated daily with 300 mL of water containing Hoagland nutrient solution [[45](#ref-epstein1972mineral)] at the concentration of 25%, this being the volume of irrigation required (i.e., previously tested) to leach excess salts and prevent their accumulation in soil. The germination was evaluated daily for 25 consecutive days until the seedlings were collected. In this experiment, it was considered seed germinated when the aerial part emerged from the soil.

Table 1 Information and location of the five genotypes of Jatropha curcas studied under salinity conditions.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| Genotype | City | State | Location | Altitude (masl) |
| 183 | Jaíba | Minas Gerais | 23°47'55.0'' S 53°18'48'' W | 478 |
| 114 | Umuarama | Paraná | 15°10'27.0'' S 43°53'18'' W | 430 |
| 218 | São Miguel do Araguaia | Goiás | 13°55'57.0'' S 50°09'17'' O | 350 |
| 171 | Maceió | Alagoas | 09°27'60.0'' S 35°49'41'' W | 131 |
| 133 | Santa Inês | Maranhão | 03°39'24.9'' S 45°22'36'' W | 31 |

**Evaluation of germination parameters.** For the two experiments the different germination parameters were calculated: germinability (GRP), mean germination time (MGT), germination uncertainty (GRU) and germination synchrony (GRS). All calculations, graphs and statistics were performed with GerminaR package [[46](#ref-R-GerminaR)].

**Evaluation of biometric parameters and biomass.** The mean seedling height (HGT) and mean stem diameter (STD) were evaluated at the end of the experiment [[25](#ref-moncaleano2013germination)]. For this, the seedlings were collected and separated into three components: leaves, stems and roots. The stem diameter were measured by a digital caliper (Digital Caliper, ROHS, ZAAS Precision, Piracicaba, SP). Leaf area (LFA) was evaluated by plant and for experimental unit (TLFA) [[47](#ref-pompelli2012allometric)]. To estimate dry biomass, all samples were dried by a forced ventilation oven at 70°C for 72 hours. Leaf dry weight (LDW), stem dry weight (STDW) and root dry weight (RDW) were used to calculate several biomass parameters, such as: total dry weight (TDW); shoot dry weight (STDW); Leaf weight ratio (LWR; ratio between LDW and TDW); stem dry weight ratio (SWR; ratio between STDW and TDW); shoot dry weight ratio (STWR, ratio between STDW and TDW) and root dry weight ratio (RWR; ratio between RDW and TDW).

**Experimental design and statistical analysis.** Both germination experiments were conducted in a completely randomized design. For the seed storage experiment, five storage times (0, 3, 6, 9 and 12 months) were used. The salinity experiment was composed of a factorial, where five genotypes of *J. curcas* and five different concentrations of NaCl in irrigation water (0, 50, 75, 100 and 150 mM). Each treatment consisted of four replicates with 25 seeds. Statistical analysis and generation of graphs were performed in the statistical software R [[48](#ref-R-base)]. The analysis of variance (ANOVA) was performed to evaluate the differences between the factors and the comparison of the means with the Student-Newman-Keuls test (p <0.05) [[49](#ref-R-agricolae)]. For the multivariate analysis, correlation analysis was performed [[49](#ref-R-agricolae),[50](#ref-R-corrplot)] and principal components analysis were made [[51](#ref-R-FactoMineR)].

# Results

**Germination of aged seeds.** The germination of the *J. curcas* seeds submitted to storage ranged from 9% to 15%, with values that are statistically similar to each other (Figure 1A). Although germination was not affected by storage, the mean germination time was significantly increased with storage time (Figure 1B). The germination was completely asynchronous for all storage times, with a mean of 0.16 being recorded before storage and close to zero at other times (Figure 1C). As it was observed, the germination in the time was significantly affected by the storage time (Figure 1E). However, the germination uncertainty was not affected by the storage (Figure 1D). It is verified that the seeds before storage and stored for 3 months began their germination on the 3rd and the 4th day after sowing respectively, while in other times the germination of the first seed was only computed from the 6th day. The time for stabilization of the germination was increased as the storage time was increased, being completed at 15 days without storage and at 23 days at 12 months of storage (Figure 1E).

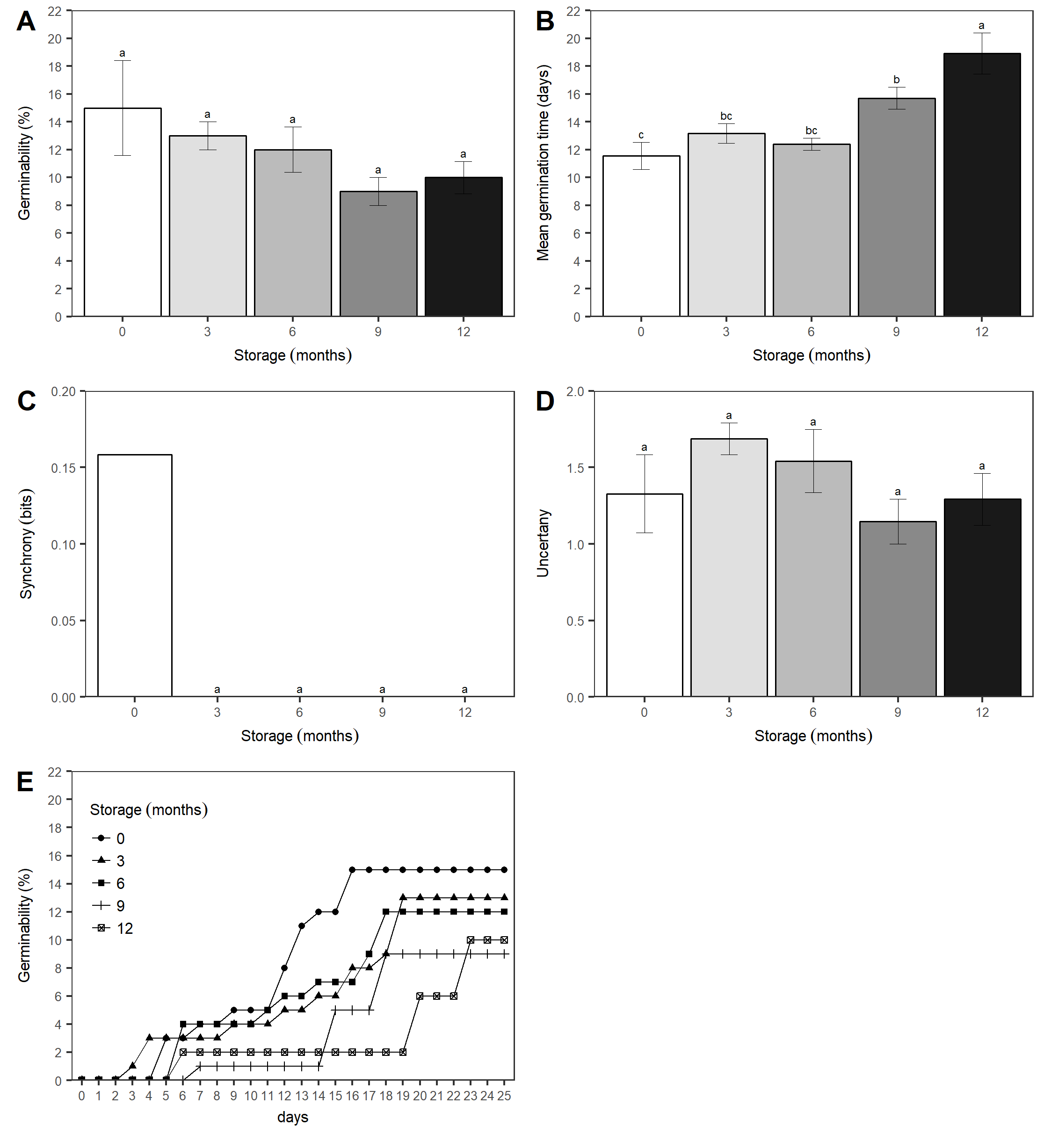


Figure 1 Germinability (A), mean germination time (B), synchrony index (C), germination uncertainty (D) and temporal germination (E) evaluated in Jatropha curcas seeds in genotype 171 stored at 0, 3, 6, 9 and 12 months. The bars represent the mean (± SE). The mean differences between the storage months are represented by the lower case letters (SNK, p = 0.05). n = 4

**Biochemical responses of seeds submitted to aging.** It was verified that the oil content in the seeds remained practically stable until the sixth month of storage at the rate of 35%, with significant reductions from 12th month of storage, when the oil content was approximately 29% (Figure 2A). We verified a strong and negative correlation between oil content in the seeds and storage time (r = -0.91, p ≤ 0.05). On the other hand, the starch was rapidly metabolized with approximate reduction of 33% with time 12 months of storage (Figure 2B). Total soluble protein and total soluble amino acid content was increased by 160% and approximately 67% during storage (Figure 2C-D). A strong and positive correlation (r = 0.92, p ≤ 0.05) is shown between total soluble proteins and amino acid syntheses.

There was a gradual reduction in the total soluble carbohydrate content at the 3rd month of storage compared to the control, without showing differences until the 12th month of storage (Figure 3A). Sucrose levels had a reduction of approximately 49% between 3 and 12 months of storage (Figure 2B), while glucose levels remained stable until the 3rd month of storage when compared to non-stored seeds. From the 6th month, the glucose was rapidly elevated, reaching 71% in the 12th month in relation to the non-stored seeds (Figure 3C). On the other hand, fructose levels did not show a trend during the months evaluated (Figure 3D). We showed that while sucrose levels decreased (r = -0.57, p ≤ 0.05) throughout the storage period, glucose (r = 0.67, p ≤ 0.05) had an opposite behavior, with increase with storage.

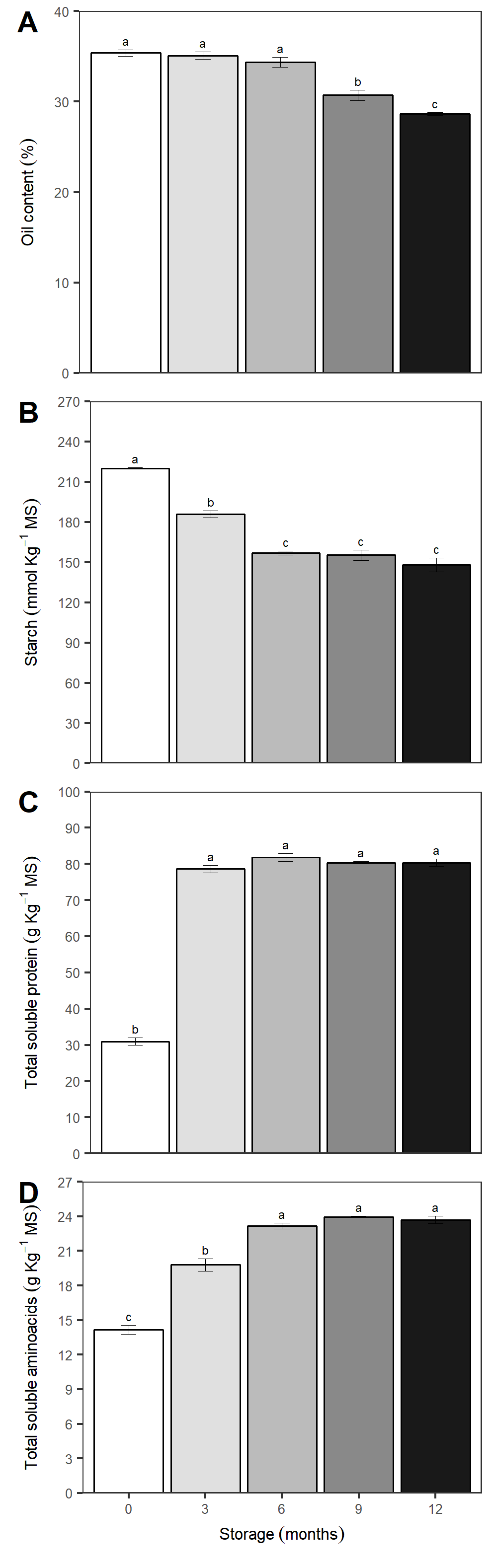


Figure 2 Oil content (A), starch (B), total soluble protein (C), total soluble amino acids (D) evaluated in Jatropha curcas seeds in genotype 171 stored for 0, 3, 6, 9 and 12 months. The bars represent the mean (± SE). The mean differences between the storage months are represented by the lower case letters (SNK, p = 0.05). n = 4

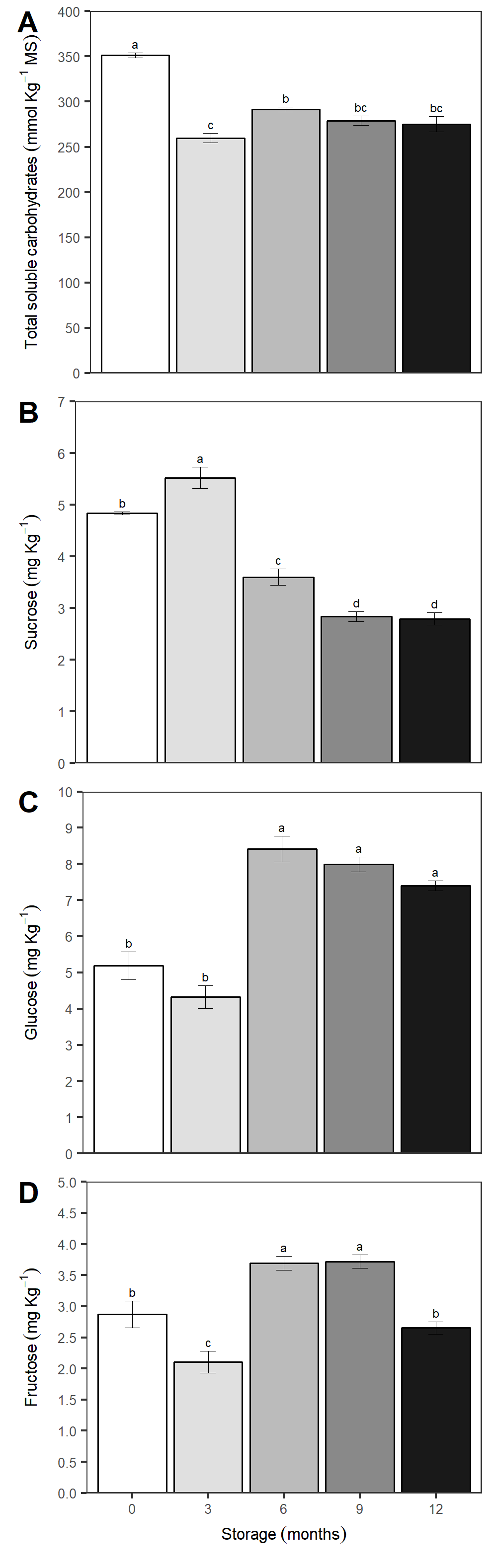


Figure 3 Total soluble carbohydrates (A), sucrose (B), glucose (C) and fructose (D) evaluated in Jatropha curcas seeds in genotype 171 were stored for 0, 3, 6, 9 and 12 months. The bars represent the mean (± SE). The mean differences between the storage months are represented by the lower case letters (SNK, p = 0.05). n = 4

**Physiological responses of seeds submitted to aging.** With the storage time in desiccant agent, it was verified that the water content in the seeds was greatly reduced (Figure 4). Non-stored seeds had 8% of water content, while seeds stored by 12 months had 5.5% (Figure 4A). Concomitantly the water content, the seeds water potential had a strong reduction with the storage time, presenting a strong correlation (r = 0.83, p ≤ 0.05) between these two characteristics. We verified that the water potential of non-stred seeds were -35 MPa, but water potential were reduced to -124 MPa at 12 months of storage (Figure 4B). With the reduction of the relative water content and the water potential there was a strong reduction in the respiratory rate of the seeds (r = 0.88, p ≤ 0.05), ranging 115 mmol CO2 h-1 g-1 MF, in non-stored seeds to 10 mmol CO2 h-1 g-1 MF after 12 months of storage (Figure 4), a reduction of 91% over the seeds storage (Figure 4C).

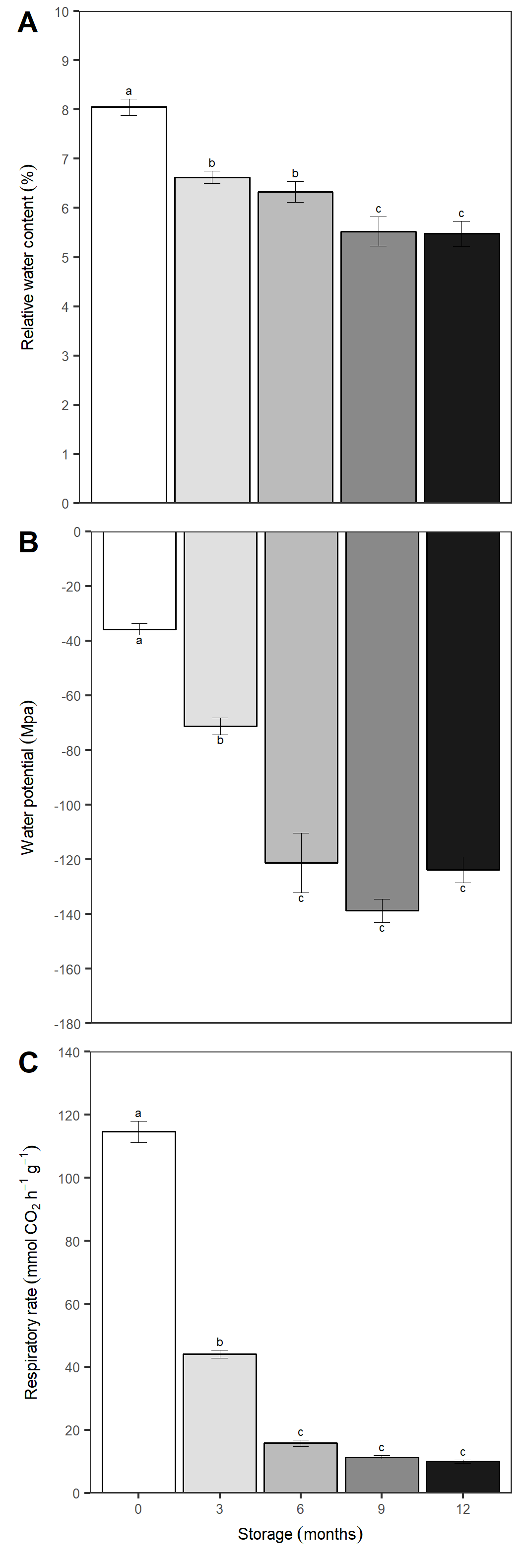


Figure 4 Relative water content (A), water potential (B), seed respiratory rate (C), evaluated in Jatropha curcas seeds in genotype 171 stored at 0, 3, 6, 9 and 12 months. The bars represent the mean (± SE). The mean differences between the storage months are represented by the lower case letters (SNK, p = 0.05). n = 4.

**Seed germination treated with NaCl.** These experiments were conducted in greenhouse condition. The temperature was 30.6 ± 1.1°C and relative humidity of 70.4 ± 5.8%. The enviromental data were collected every fifteen minutes during all days, 24 hours per day. In this experiment, we verified that the germination was almost zero at 150 mM NaCl for all genotypes. In 0 mM NaCl, seeds of genotype 183 and 114 had 71% and 86% of germination, respectively, with gradual decrease with the increase in NaCl concentration, from 4% to 100 mM and 0% to 150 mM. In another way, genotypes 218 and 133 did not differ by up to 100 mM NaCl, although germination was reduced to approximately 5% at the concentration of 150 mM. Seeds of the genotype 171 showed 65% germination in the control, reducing to 32% and 9% in 50 mM and 150 mM NaCl respectively (Figure 5A). The mean germination time for 0 mM NaCl was 5 to 7 days for all genotypes, while for the 150 mM NaCl concentration the interval was longer, ranging from 7.2 days to 12.3 days. No significant differences were observed in mean germination time from 50 mM to 75 mM, with a general average of 7.5 days for all genotypes (Figure 5B). There was no difference in the germination synchrony for the genotypes up to 75 mM NaCl. However, the synchronization at 100 mM was null at genotype 183, 114 and 218, as well as for genotype 133 in the concentration at 150 mM NaCl. It was observed that the synchrony were always lower than 0.25 for all concentrations and genotypes (Figure 5C), denoting a asynchronous profile. The maximum value for the uncertainty in germination in this experiment was 4.64 bits. Genotypes 183 and 114 showed a tendency to reduce uncertainty with the increase in NaCl concentration. The germination uncertainty was stable up to 75 mM NaCl, but with a significant increase in the concentration of 100 mM, where the uncertainty was 0.5 and 0.3 for the genotype 183 and 114, respectively. Genotypes 218, 171 and 133 showed a trend in increasing uncertainty from 0 mM to 100 mM without showing significant differences. At 150 mM, the uncertainty was 0.3, 0.7 and 0.5 for genotypes 218, 171 and 133 respectively (Figure 5D). The germination in time showed differences for each of the genotypes. The germination in the treatments without salt began between the 3rd and 4th day after sowing; for the treatments with NaCl addition the maximum values of germination were observed between the 9th and 12th day (Figure 6A-E). Regardless of the genotypes, first germination was observed on the 3rd day in 0 mM NaCl, while in the 50 mM and 75 mM NaCl, the germination was generally initiated on the 4th day, but arranged 5th day and 7th day in 100 mM and 150 mM NaCl, respectively. However, regardless of treatments and salinity levels, germination became stable since 13th day (Figure 6).

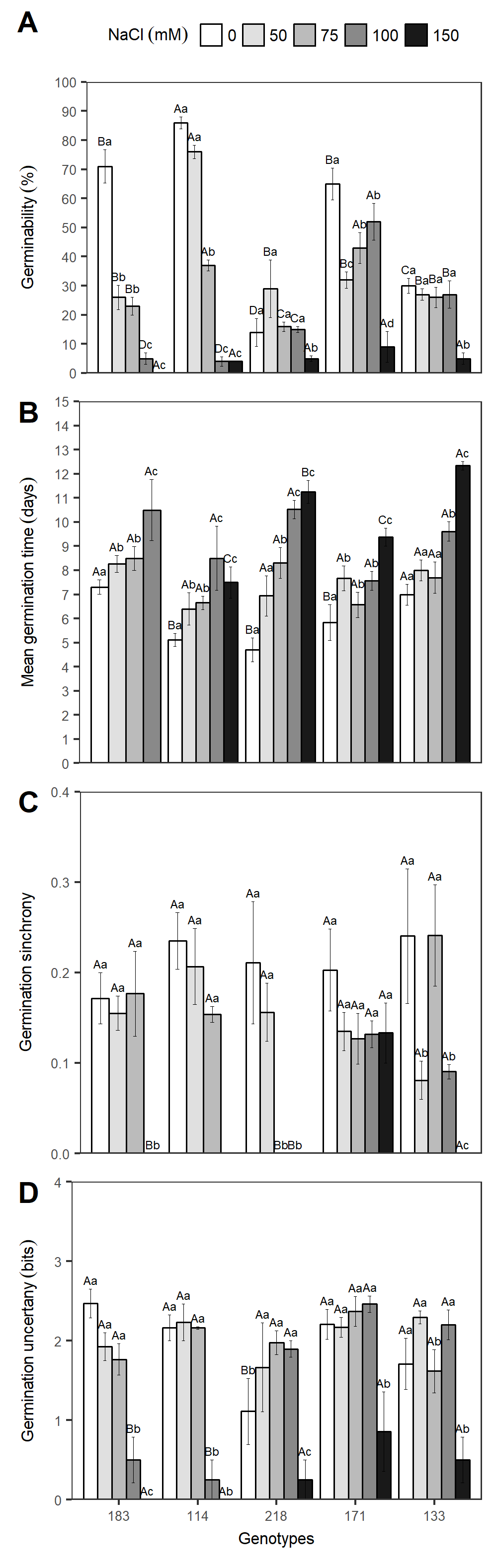


Figure 5 Germinability (A), mean germination time (B), germination synchrony (C) germination uncertainty (D) evaluated in five genotypes of Jatropha curcas L. under different NaCl concentrations (0, 50, 75, 100 and 150 mM). The vertical bars represent the mean (± SE). The mean differences between the accessions are represented by different capital letters and between salt levels by different lowercase letters (SNK, p = 0.05). n = 4

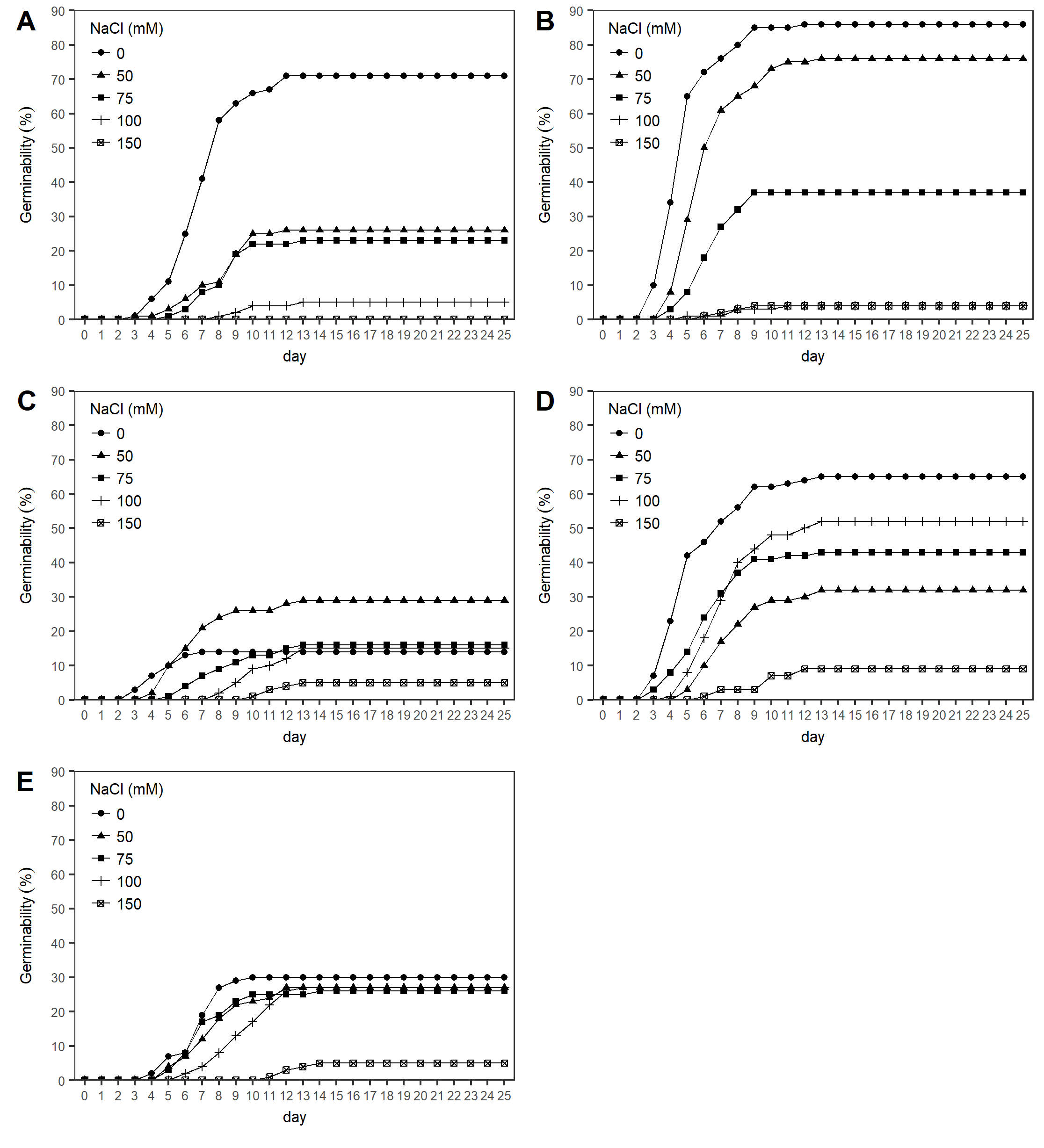


Figure 6 Cumulative germination in the time evaluated in five genotypes of Jatropha curcas L. under different concentrations of NaCl (0, 50, 75, 100 and 150 mM). Genotype 183 (A), 114 (B) 218 (C), 171 (D) and 133 E)

**Biometric and biomass components** Although the germination was evaluated up to the concentration of 150 mM NaCl, the biomass production was only computed up to 100 mM NaCl, and the biomass parameters were strongly affected by the increase in salinity. The leaf area (LFA) was only possible to be evaluated until the concentration of 75 mM NaCl, since these was the parameters most affected by the increase of NaCl in the irrigation solution. The genotypes 218, 171 and 133 presented the highest values in the control concentration, while for 50 mM the genotypes with the highest performance were 133, 114 and 183. The genotypes 133, 114, 171 and 183 presented the highest value in the 0 mM treatment, followed by genotype 183 and 171 that did not show significant differences between them. At the concentration of 50 mM, genotype 114 present larger leaf area, followed by genotype 133, while at the 75 mM concentration there was no difference between the genotypes. There was a tendency of plant height (HGT) reduction with increasing salts. The HGT reached approximately 13 cm in the control, but was strongly reduced until values smaller than 3 cm to 100 Mm of NaCl, fact clearly recorded in genotype 183 (Figure 7A). The stem diameter (STD) presented reduction with the increase of the concentration of NaCl, even without statistical differences. At concentrations of 50 mM and 75 mM, the genotypes 183 and 114 obtained the highest values of stem diameter, while at the concentration of 100 mM they had the lowest values (Figure 7C). For the leaf dry weight ratio, it is possible to observe a tendency in the reduction with the salinity increase, being genotype 218 presenting a ratio of 0.42, followed by genotype 183 with a ratio of 0.39. These two genotypes were the ones that best behaved in this allometric parameter. Genotype 171 showed no significant difference in leaf weight ratio up to 50 mM, while genotype 114 increased its biomass accumulation to 50 mM. All genotypes showed drastic reduction from 75 mM NaCl, with a reduction of biomass accumulation in the leaf with increasing NaCl in the irrigation water (Figure 7F). A distinct profile is verified in the stem dry weight ratio (SWR), which was increased with elevation of salinity, even though no significant effect was observed up to the 50 mM concentration. For the 75 mM concentration, genotype 183 was better than the others, although similar to genotypes 171 and 133 at 100 mM concentration (Figure 7D). The root dry weight ratio (RWR) showed a distinct behavior for each genotype, with genotypes 171, 133 and 218, increasing biomass of the roots up to 75 mM and showed a continuous increment up to 100 mM. On the other hand, the genotype 183 showed increment up to 50 mM and from there had its values reduced. Genotype 114 was apparently unaffected by the increase in NaCl concentration (Figure 7B).

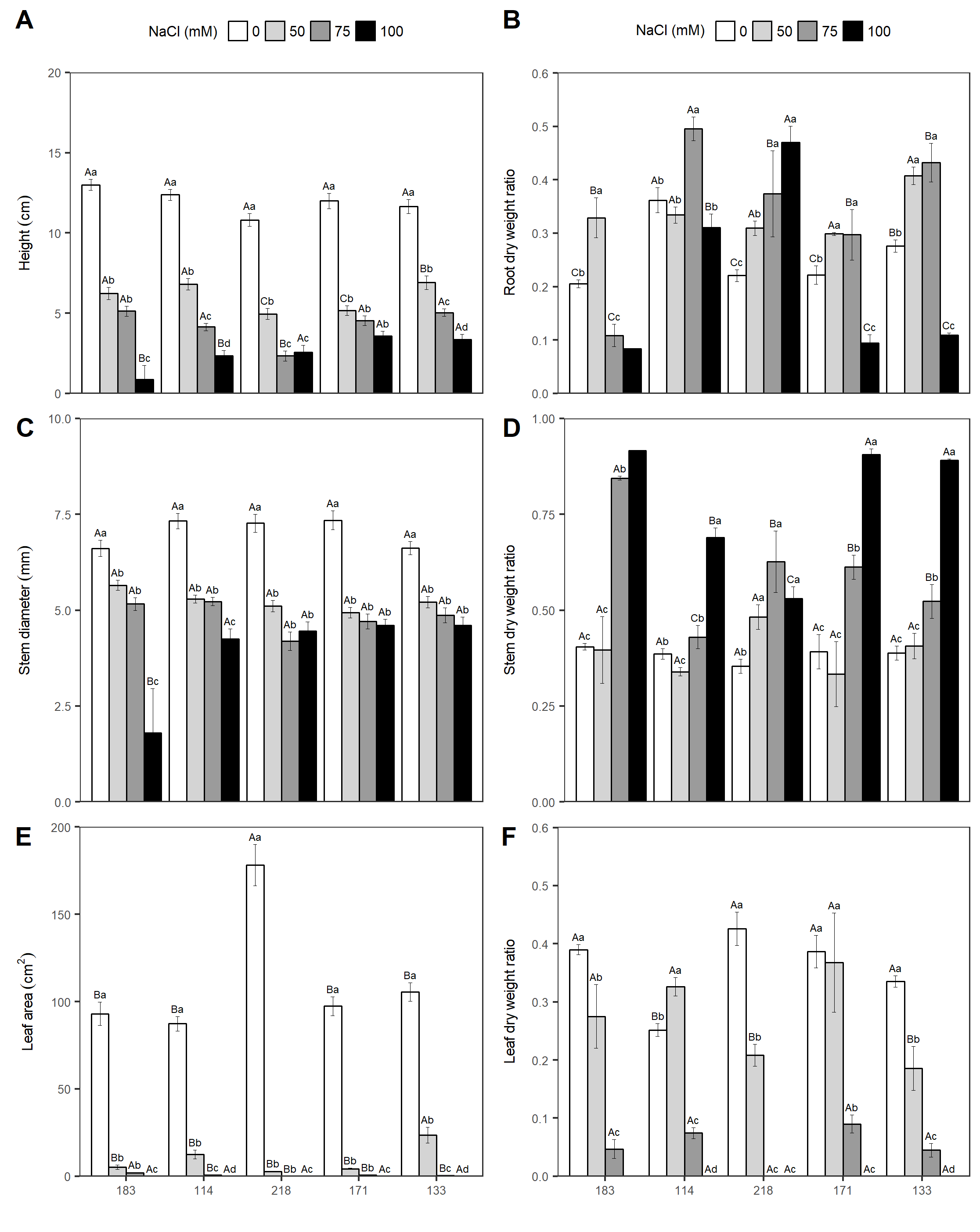


Figure 7 Leaf dry weight ratio (A), stem dry weight ratio (B), root dry weight ratio (C) and shoot dry weight ratio (D) evaluated in five genotypes of Jatropha curcas L. under different concentrations NaCl (0, 50, 75, 100 and 150 mM). The vertical bars represent the mean (± SE). The mean differences between the accessions are represented by different capital letters and between salt levels by different lowercase letters (SNK, p = 0.05). N = 4.

**Multivariate analysis of the salinity experiment.** The germination parameters were negatively affected by the increase in the salt concentration; There was a negative correlation between germination percentage (r = -0.55, p ≤ 0.05) and germination synchrony (r = -0.69, p value ≤ 0.001). On the other hand, the mean germination time was positively significant when correlated with the increase of salts (R = 0.66, p ≤ 0.001). However, there was no significant correlation between germination uncertainty and salt increase (r = -0.23, p = 0.321). The biomass parameters were significantly affected with increasing salts. The correlations between this parameter in relation to the leaf dry weight (r = -0.86, p ≤ 0.001), the leaf area (r = -0.89, p ≤ 0.001), and plant height ( R = -0.89, p ≤ 0.001) and stem diameter (r = -0.91, p ≤ 0.001). Other allometric parameters were also influenced by NaCl addition; the negative relationship between salinity and leaf weight ratio (R = -0.90, p ≤ 0.001) and leaf area ratio (r = -0.94, p ≤ 0.001). On the other hand, were strongly increased with salinity the stem dry weight ratio (R = 0.74, p ≤0.001). The principal components analysis (PCA) shows that approximately 74.4% of the variation can be explained by the parameters studied in this work. In the first component the variables LDW, TLFA, HGT and TDW were the ones that presented the greatest contribution in the variance and are negatively correlated with the increase of NaCl, while the variables with positive correlation were the MGT and SDW. The genotypes that showed the best response were 114, 171 and 183 and the susceptible genotypes 133 and 218 (Figure 8).

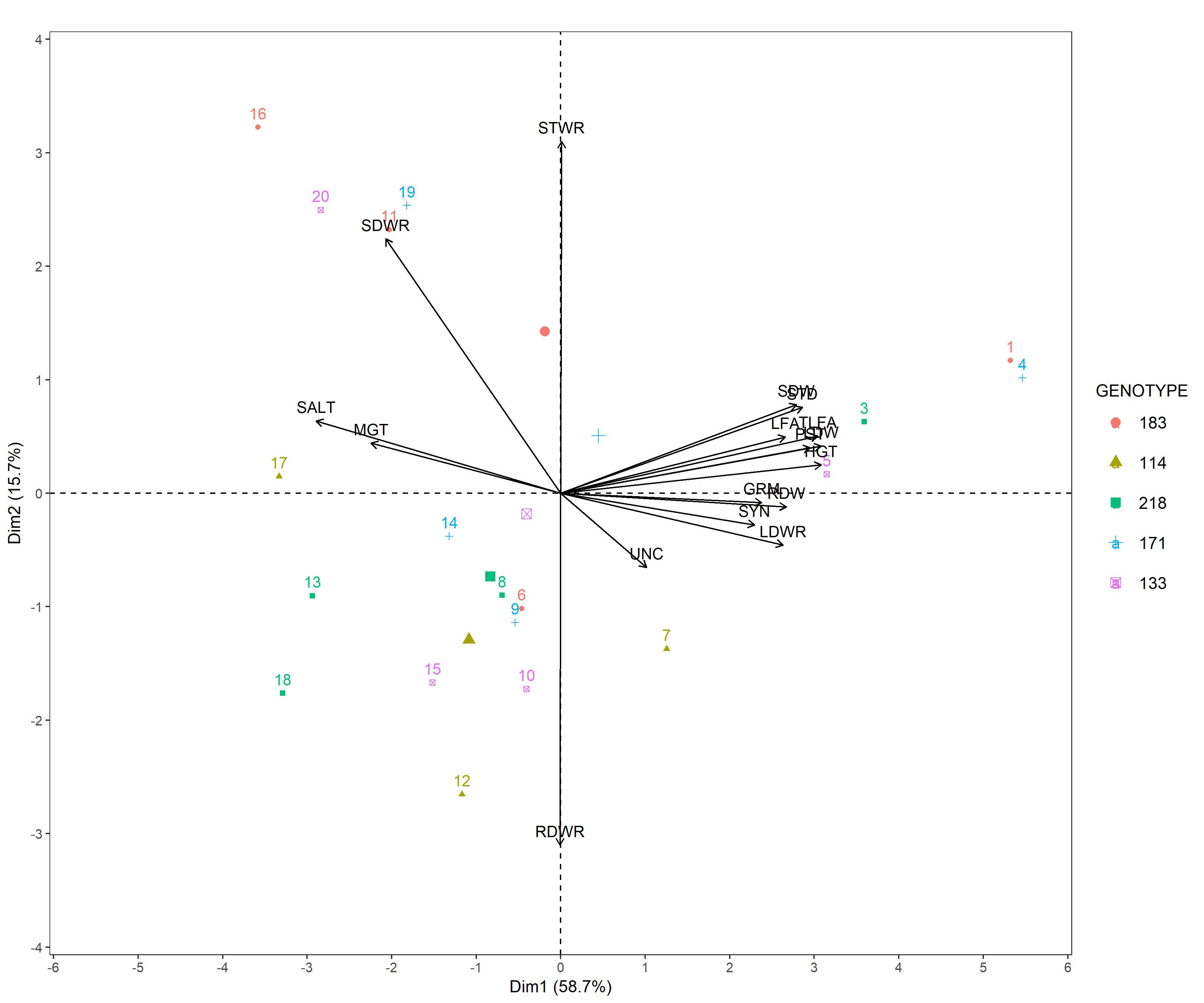


Figure 8 Principal components analysis (PCA) of the variables evaluated in five genotypes of Jatropha curcas L. under different concentrations of NaCl (0, 50, 75, 100 and 150 mM). GRM, Germinability; MGT, mean germination time; UNC, Uncertainty of germination; SYN, Synchronism of germination; LDW, leaf dry weight; RDW, root dry weight; SDW, stem dry weight; TDW, total dry weight; LFA,leaf area; TLFA, total leaf area; LDWR, leaf dry weight ratio; SDWR, stem dry weight ratio; RDWR, root dry weight ratio; STWR, shoot dry weight ratio; HGT, plant length; STD, stem diameter.

# Discussion

The seeds of *J. curcas* are classified as orthodox [[27](#ref-hay2013advances)], resistant to desiccation and can present water contents up to 18% [[5](#ref-pompelli2010environmental)] when freshly harvested. Moncaleano-Escandon [[25](#ref-moncaleano2013germination)], showed that the seeds of *J. curcas* can drastically reduce their germinability during storage at temperatures of 4°C or 25°C, accompanied by the reduction of some compounds such as starch and total soluble proteins. This author also showed that at 4°C it was more interesting for the storage of seeds of this species. In this work, we observed that the reduction of the relative humidity in the interstice of the seeds reduced the water content of the seeds and consequently the water potential. Coupled to these facts, the respiration rate were strongly reduced, but not zero, which allowed the viability of the embryo at the expense of the solubilization of reserves, that supported the germinability of the seeds during the storage without significant reduction. However, we shows a strong and positive correlation (r = 0.92, p ≤ 0.05) between total soluble proteins and amino acid syntheses. A possible explanation could be that structural proteins were mobilized to generate carbon skeleton to respiration or amino acids as a compatible solutes that allows maintain the respiration even with reduction of seed moisture. Similarly described previously, we showed a significant correlation between decrease of sucrose and elevation to glucose (r = -0.87, p ≤ 0.05) and fructose (r = -0.56, p ≤ 0.05). The total soluble carbohydrates content must have been reduced by the metabolism of sucrose, which acted as the carbon source for the embryo. Circumstantial evidence of this idea is based on the elevation of glucose and fructose contents while there has been a reduction in the levels of starch and sucrose. Thus, the respiration rate, even low, foresees the live embryo, which allowed the germination, even 12 months after storage, a fact that diverges of Moncaleano-Escandon et al. (2013) [[25](#ref-moncaleano2013germination)] which describes that the germination of *J. curcas* seeds drop near to zero after 12 months of storage, but six months after start of experiments the seeds stored in low temperature drop 27%, while in this experiment the germination remained stable during all storage time. It should be noted that Moncaleano-Escandon et al. (2013) [[25](#ref-moncaleano2013germination)], stored its seeds without any type of desiccant and, in this study, seeds were stored at 4ºC as recommended by Moncaleano-Escandon et al. (2013) [[25](#ref-moncaleano2013germination)], but in boxes with very low RH, promoted by the use of desiccant. Another high evidence of respiration was responsible for the mobilization of reserves can be corroborated by the strong negative correlation between oil, starch and sucrose contents with its degradation products (i.e., total soluble proteins, total soluble amino acids and glucose). In addition, it was reported that seeds containing between 6% and 8% of moisture had a rapid reduction in seed oil content in the first 3 months of storage, together with an increase in the concentration of free fatty acids [[52](#ref-akowuah2012influence),[53](#ref-worang2008quality)] Thus, it is verified that the moisture control in the interstice of the seeds should be taken into account to preserve both the viability and the oil content of the seeds of *J. curcas*.

The negative effects of saline stress were reflected in the delay in the mean germination time, from 3 days in the 0 mM NaCl and up to 13 days in 150 mM NaCl in the irrigation water, which is confirmed by the significant reduction in germination rate. Same patter was previously reported in *J. curcas* by Alencar et al. (2015) [[16](#ref-alencar2015ultrastructural)] which describes a strong and negative correlation between germination rate and mean germination time. Regardless of the storage time, germination was initiated between the third and fifth day, with complete finishing after 23 days of sowing, which occurred mainly with 12 months of storage. Both salinity and storage delayed germination; But among them, storage seems to be the factor that slows down germinability with greater magnitude. It was observed that the germination synchrony was reduced with the increase of the salts concentration, and the salinity stress promoted slow and more disorganized germination, a fact corroborated by the high values of germination uncertainty. That *J. curcas* has an asynchrony in germination, mainly in salt stress, is already very well studied [[5](#ref-pompelli2010environmental),[16](#ref-alencar2015ultrastructural),[25](#ref-moncaleano2013germination),[54](#ref-islam2009effect),[55](#ref-silva2012relationship)]. Therefore, if we analyze the previously published data with those presented of this study we can postulate that synchrony and the uncertainty of the germination can not be considered a good parameters for judgment, at least in *J. curcas*. A possible explanation for this could arise from the fact that *J. curcas* is not yet a domesticated species [[56](#ref-achten2010towards)], which makes it present high levels of uncertainty in germination [[57](#ref-ranal2006and)]; A factor highly related to the survival of the species in its original habitat [[58](#ref-maes2009climatic)] and very unstable from the physiological point of view. Seeds of *J. curcas* can not tolerate up to 150 mM NaCl in the irrigation water for seed germination, and it has very difficulties for seedlings development in concentrations above 75 mM of NaCl. The delay of the germination accompanied with the decrease of the development of the leaves and the reduction of the root growth promotes delay of the autotrophic phase of the plants, in extreme cases, leads to the death of the seedling in the first days after germination. A possible explanation for this is presented by Alencar et al. (2015) [[16](#ref-alencar2015ultrastructural)] since there was a great increase of the Na+ and Cl- contents in the embryonic axes and in the endosperm of the seeds of *J. curcas*. Another possibility is loss of mobilization of cotyledon reserves on germination [[59](#ref-liu2010seed)] affecting seedling establishment [[60](#ref-marques2013increased)].

It is common for plants to increase their stem biomass to the detriment of other organs when subjected to salinity [[61](#ref-dantas2007germination)–[63](#ref-praxedes2010salt)]. Munns [[62](#ref-munns1986whole)] describes this as an indirect effect of decreasing water uptake by roots and lower leaf expansion, while Praxedes [[63](#ref-praxedes2010salt)] describes this effect as lower relative growth rate of the plant as a whole. On the other hand, Bayuelo-Jiménez [[64](#ref-bayuelo2002salinity)] and Debez [[65](#ref-debez2004salinity)] describe that all biometric components are reduced in plants submitted to salinity, whether this species is salinity sensitive [[64](#ref-bayuelo2002salinity)] or a halophyte [[65](#ref-debez2004salinity)]. In addition, Hasewaga [[66](#ref-hasegawa2000plant)] attributed this reduction and/or inhibition of plant growth, when subjected to salinity, to a number of factors such as changes in the water status of the plant caused by the osmotic effect of the salts, increase in the concentration of toxic ions, producing physiological and biochemical variations and alteration in the absorption of essential nutrients.

The principal component analysis showed that the biometric parameters explain better the salinity response than the germination parameters. Genotypes 114, 171 and 183 are possible candidates for salt stress tolerance, whereas genotypes 133 and 218 show sensitivity to NaCl addition in the irrigation water.

In conclusion the viability of germination and oil content in seeds of *Jatropha curcas* can be maintained provided there is control of moisture in the interstices of the seed during storage. Salinity strongly affects the germination and initial development of *Jatropha curcas*, and from the evaluated genotypes, 114, 171 and 183 present as potential candidates for future breeding programs.

# Acknowledgments

The authors thank the National Council for Scientific and Technological Development (CNPq Process nº 404357/2013-0) for the financial support. The first author thanks the Coordination of Improvement of Higher Education Personnel (CAPES) for the scholarship. Special thanks to Dr. Agnaldo Rodrigues de Melo Chaves, Tropical Semiarid Agricultural Research Center, Embrapa Semiárido, Petrolina, PE, Brazil and Embrapa Agroenergy, Brasília, DF, Brazil for giving the seeds used in this study.

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