

Effect of salinity on germination of lettuce cultivars produced in Brazil

ABSTRACT

The challenge of recovering degraded soils due to salinity excess leads to the search for more effective strategies that can overcome this problem. Among these, one of the approaches is the use of resistant plant varieties in affected areas. This study aimed to evaluate the influence of different doses of salt on seed germination and seedling formation of two lettuce cultivars (Hanson and H121) and to verify the existence of tolerance among the cultivars. A completely randomized design was used in a 2x5x4 factorial scheme, where the cultivars were evaluated under five distinct salt doses (0, 25, 50, 75 and 100 mol.m⁻³, conductivities of 0.0, 2.8, 5.4, 8.0 and 10.6 dS.m⁻¹, respectively), with four replicates. In this sense, the following variables were evaluated: germination rate (GR), germination speed index (GSI), seedling height (SH), root length (RL) and percentage of dry matter in relation to fresh matter (DM%). As a result, the Hanson cultivar presented better performance than the H121, under all the different salt doses, in all the studied variables. Also, the EC of 2.8 dS.m⁻¹ did not affect any of the studied variables, including both cultivars. However, EC above 2.8 dS.m⁻¹ had, significantly, reduced the development of the cultivars. The Hanson cultivar was influenced only in the variables SH, RL and DM%, where SH and DM% were influenced by EC values above 8.0 and 10.6 dS.m⁻¹, respectively. The cultivar H121 was significantly influenced in all evaluated variables, which demonstrates its greater susceptibility to salinity.

Keywords: Electrical conductivity, osmotic potential, water consumption, tolerance.

1. INTRODUCTION

Population growth, food security and the scarcity of nonrenewable natural resources are some of the challenging themes that are present in the main discussions about the transformations in contemporary agriculture [1,2]. In a more rational model, priority is given to respect for the environment, where there is a growing concern with the management of natural resources, and at the same time, with the economic viability of the agricultural sector for its professionals, in order to guarantee the sustainability of current and future generations [3,4].

The development of sustainable agriculture is intrinsically related to the possibility of adopting measures that overcome the main challenges of agricultural production, in a rational and environmentally friendly way [5]. Many are the adversities found in agricultural fields throughout the world, but in the past few years, one has been gaining prominence, the soil salinity [6]. Soil salinity can occur naturally due to the characteristics of the source material [7]. However, the most damaging salinity is that induced by modern agricultural systems, where the irrational use of fertilizers and the use of brackish water for irrigation are the main agents that cause this adversity [8,9].

Soil salinity can make it unproductive, indirectly reducing the income of the farmers who cultivate the soil [10]. In Brazil, small farmers are the class that suffers the most from soil salinization, since the low income of these producers does not allow them to adopt

36 technologies to reverse it. Thus, technologies that allow cultivation on lands with an excess
37 of salts are indispensable for the success of these farmers in the field. Among the available
38 technologies, there is the selection of cultivars that tolerate high salinity concentrations.

39 Among the crops produced by small farmers, which suffer from the elevation of salinity
40 levels, there is the lettuce (*Lactuca sativa* L.) [12]. The lettuce stands out because of its low
41 cost of production, market acceptability, and its cultivation requires small areas, which
42 makes it suitable for small farmers [12,13]. Lettuce cultivation is strongly affected by salinity,
43 and about 13% of the yield declines by each unit of electrical conductivity (EC) increased
44 above 1.3 dS m⁻¹ [13]. However, there is variation among cultivars, where some differ from
45 the others regarding their tolerance to high levels of salinity [14,13].

46 The tolerance to salinity is a polygenic characteristic, which makes it difficult to identify
47 tolerant genotypes, due to a large number of genes involved in the trait's control [15,16].
48 Identifying tolerant materials among the cultivars on the market has become an important
49 strategy for expansion of lettuce crops in places where salinity is a limiting factor. Since it
50 reduces the time and cost to obtain superior lineages in breeding programs. Given the
51 above, this study aimed to evaluate the influence of different salt concentrations on
52 germination of two lettuce cultivars, as well as to verify the existence of tolerance to salinity
53 in the cultivars.

54 2. MATERIAL AND METHODS

55 The study was conducted in a completely randomized design in a 2x5x4 factorial scheme.
56 The Hanson (cultivar 1) and H121 (cultivar 2) cultivars were used. To evaluate the tolerance
57 to salinity, five distinct concentrations of sodium chloride (NaCl) were tested: 0, 25, 50, 75
58 and 100 mol.m⁻³. All concentrations of NaCl were diluted in distilled water, whose initial EC
59 was zero. These concentrations corresponded to solutions with the following electrical
60 conductivities: 0.0, 2.8, 5.4, 8.0 and 10.6 dS.m⁻¹, respectively, measured with a conductivity /
61 TDS and salinity meter.

62 To evaluate the germination, four replicates of 25 seeds were made in each treatment,
63 wherein previous tests all the seed lots used in the study presented a germination rate (GR)
64 higher than 90%. The test was performed in a 10 x 1.5 cm petri dish with the use of
65 germitest paper. The paper was accommodated in double sheets at the bottom of the petri
66 dish, and the seeds were evenly distributed over it. The saline solution was added after the
67 seed distribution at 2.5 times the dry paper weight.

68 In order to favor germination, the seeds were kept in a Biochemical Oxygen Demand
69 incubator (BOD) for 7 days, allowing total control of lighting and temperature that was
70 alternated to maintain 12 hours of light and 12 hours of dark at a temperature of 25 ° C.

71 The germination evaluation occurred on the seventh day with the seedling count as
72 proposed by the authors [17]. Also, only seeds with a primary root greater than 2 mm, as
73 established by the authors [18], were considered as germinated. Through the counting data,
74 it was possible to determine the GR in each treatment.
75 Along with the germination test, daily germinated seedlings were counted from the beginning
76 of the sowing until the seventh day, when the germination was stabilized. The germination
77 speed index (GSI) was obtained using the equation (1) proposed by the author [19].

$$78 \text{ GSI} = \sum (n/t)(1)$$

79 Where:

80 n = number of normal seedlings computed on the day of the count;

81 t = number of days from sowing until the day of the count.

82 After the germination stabilization on the seventh day, the variables seedling height (SH),
83 root length (RL) and percentage of dry matter in relation to fresh matter (DM%) were
84 determined. To do so, ten normal seedlings of each replicate were selected. The
85 determination of the SH and RL was performed using a graduated ruler (centimetres). SH
86 was measured from the neck to the apical meristem, while the RL was measured from the
87 neck to the root cap.

88 The DM% was obtained using the equation (2):

89 $DM\% = mf/mi$ (2)

90 Where:

91 DM%: percentage of dry matter in relation to fresh matter;

92 mf: Total dry mass of the seedling;

93 mi: Total fresh mass of the seedling.

94 The seedlings had their fresh and dried masses measured by a precision scale. In order to
95 obtain the dry mass values, the seedlings were left in an oven at 75° C until they reach a
96 constant weight. Lastly, the data were submitted to the Kolmogorov Smirnov normality test,
97 and then the variance analysis was performed. The means between treatments within the
98 same cultivar were compared by the Tukey test, and the means of treatments between the
99 cultivars were compared by the t-test. A simple linear regression analysis was used to infer
100 about the behaviour of the different cultivars under the effect of different salt concentrations
101 in the studied variables. All statistical analyzes were performed at a 5% probability level.

102 3. RESULTS AND DISCUSSION

103 The results presented in table 1 demonstrate that cultivar 1 obtained superior performance in
104 relation to cultivar 2, for all the different EC tested. Cultivar 2 was significantly influenced (p-
105 value <0.005) by the EC in all evaluated variables, while for cultivar 1, only SH, RL and
106 DM% variables were influenced.

107 The EC of 2.8 dS.m⁻¹ did not affect any of the studied variables in both cultivars (Table 1).
108 Similar results were found by the author [14] when studying salinity levels in the germination
109 of the Elba cultivar. Conversely, EC higher than 2.8 dS.m⁻¹ negatively affected the root growth
110 of both cultivars. The authors [20] emphasize the importance of root formation in this
111 phenological phase of the vegetables. The low root development in this phase culminates in
112 fragile seedlings that tend to suffer more from environmental and biological stresses.

113 **Table 1. Comparative test of averages for germination rate (GR), germination speed**
114 **index (GSI), seedling height (SH), root length (RL) and percentage of dry matter in**
115 **relation to fresh matter (DM%), for the cultivars 1 and 2 submitted to different**
116 **electrical conductivities.**

EC (dS.m ⁻¹)					
0	2.8	5.4	8	10.6	

Cultivar		GR (%)			
Cultivar 1	99.67±0.33	99.67±0.33 Aa	99.0±1 Aa	98.0±1.15 Aa	98.0±1.15 Aa
	Aa				
Cultivar 2	91.0±1 Ab	89.0±4.43 Aa	74.0±6.63 ABb	58.0±4.76 Bb	37.0±1.91 Cb
		GSI			
Cultivar 1	39.49±0.24	39.34±0.34 Aa	39.54±0.17 Aa	38.41±0.51 Aa	38.0±0.74 Aa
	Aa				
Cultivar 2	24.34±0.94	23.54±1.62	18.29±1.94	13.49±0.71	8.3±0.39 Db
	Ab	ABb	BCb	CDb	
		SH (cm)			
Cultivar 1	2.97±0.08	3.18±0.04 Aa	2.91±0.06 ABa	2.63±0.1 Ba	2.02±0.12 Ca
	Aba				
Cultivar 2	1.62±0.07 Ab	1.46±0.05 Ab	1.42±0.05 Ab	1.06±0.03 Bb	0.79±0.06 Cb
		RL (cm)			
Cultivar 1	1.64±0.15 Aa	1.40±0.08 ABa	1.07±0.06 BCa	0.82±0.02	0.66±0.01 Da
				CDa	
Cultivar 2	0.45±0.07 Ab	0.37±0.01 ABb	0.34±0.02 ABb	0.3±0.03 Bb	0.24±0.01 Bb
		DM (%)			
Cultivar 1	9.93±0.4 Aa	8.28±0.7 Aa	9.05±0.51 Aa	9.39±0.32 Aa	12.91±0.41
					Ba
Cultivar 2	16.37±0.92	14.68±0.96 Ab	14.7±1.04 Ab	20.13±2.70 Ab	30.32±3.48
	Ab				Bb

117 Means followed by the same capital letter, in the same line, do not differ by the Tukey test (P
118 <0.05) and means followed by the same lowercase letter, in the same column, do not differ
119 from each other by the t -test ($P <0.05$).

120 Figure 1A illustrates the influence of EC on the GR of seedlings from both cultivars. Cultivar
121 1 did not suffer influence in the number of germinated seedlings, unlike cultivar 2, which can
122 also be observed in table 1. In cultivar 2, the germination decreased from 91%, in the control
123 group, to 37% for the highest EC tested, demonstrating its sensitivity to high levels of

124 salinity. Studies conducted by the authors [21] demonstrate that may exist different levels of
125 salinity tolerance during the germination process among commercial lettuce cultivars.

126 The absence of influence on the seedlings GR of cultivar 1 may indicate that it tolerates high
127 saline concentrations in this phenological phase. Salinity tolerance in the germination
128 process is associated with the activation of mechanisms that lead to changes in cellular
129 metabolism. Among these mechanisms, there is the ability to accumulate organic ions and
130 low molecular weight solutes in cell vacuoles, to maintain water absorption, even in
131 unfavourable osmotic conditions [22].

132 According to Figure 1B, GSI was different between the evaluated cultivars. For cultivar 1, it
133 was practically constant, not differing between the treatments (Table 1). Conversely, a
134 contrary effect can be observed for cultivar 2, where the decreasing curve in Figure 1B
135 demonstrates how affected it was. These results reinforce the argument previously
136 mentioned, regarding the cultivar 1 tolerance and the cultivar 2 susceptibility when submitted
137 to saline environments. Studies carried out by the authors [23] confirm that the seeds tend to
138 be vulnerable when subjected to the effects of salinity. The GSI of the seeds can be altered
139 by a forced dormancy. The dormancy occurs due to the decrease in the water absorption by
140 the seeds, which negatively affects the imbibition and, consequently, reduces the elongation
141 and divisions of the cells, thus preventing the mobilization of indispensable reserves in the
142 germination process [24,23].

143 The adjusted curves in Figure 1C show the behaviour of the DM%. This variable is
144 completely associated with the water intake of the seedlings, since the higher dry mass/fresh
145 mass ratio, the lower water content in plant tissues [25]. Results obtained by the authors [26]
146 showed that the water consumption by lettuce plants is reduced linearly by the increase of
147 the EC induced by the addition of salts, and it can be reduced by up to 45% when EC is
148 raised to 10.4 dS m^{-1} . In the present study, from the values of the DM% variable, it can be
149 inferred that the water consumption of the cultivars was differently affected. The adjusted
150 curves for the DM% variable show that the water consumption in cultivar 1 was not
151 influenced by the EC increase, while in cultivar 2 it was reduced when the EC reached 5.0
152 dS m^{-1} .

153 The growth of the seedlings' shoot and the root system was reduced with the progressive
154 increase of the salinity (Figure 1D and 1E). Similar results were found by the authors [21]
155 when studying the influence of salinity on two lettuce cultivars. Seedling development is
156 affected by the decline of phytohormones levels, such as auxins, gibberellins, jasmonic acid
157 and salicylic acid in plant tissues. This decrease is associated with the toxic effect of NaCl
158 excess on plants [27,28]. The reduction of these plant hormones along with a decrease in
159 the osmotic potential, caused by saline stress, results in a smaller number of cell divisions.
160 Consequently, a lower vegetative growth of the seedlings occurs [29,27].

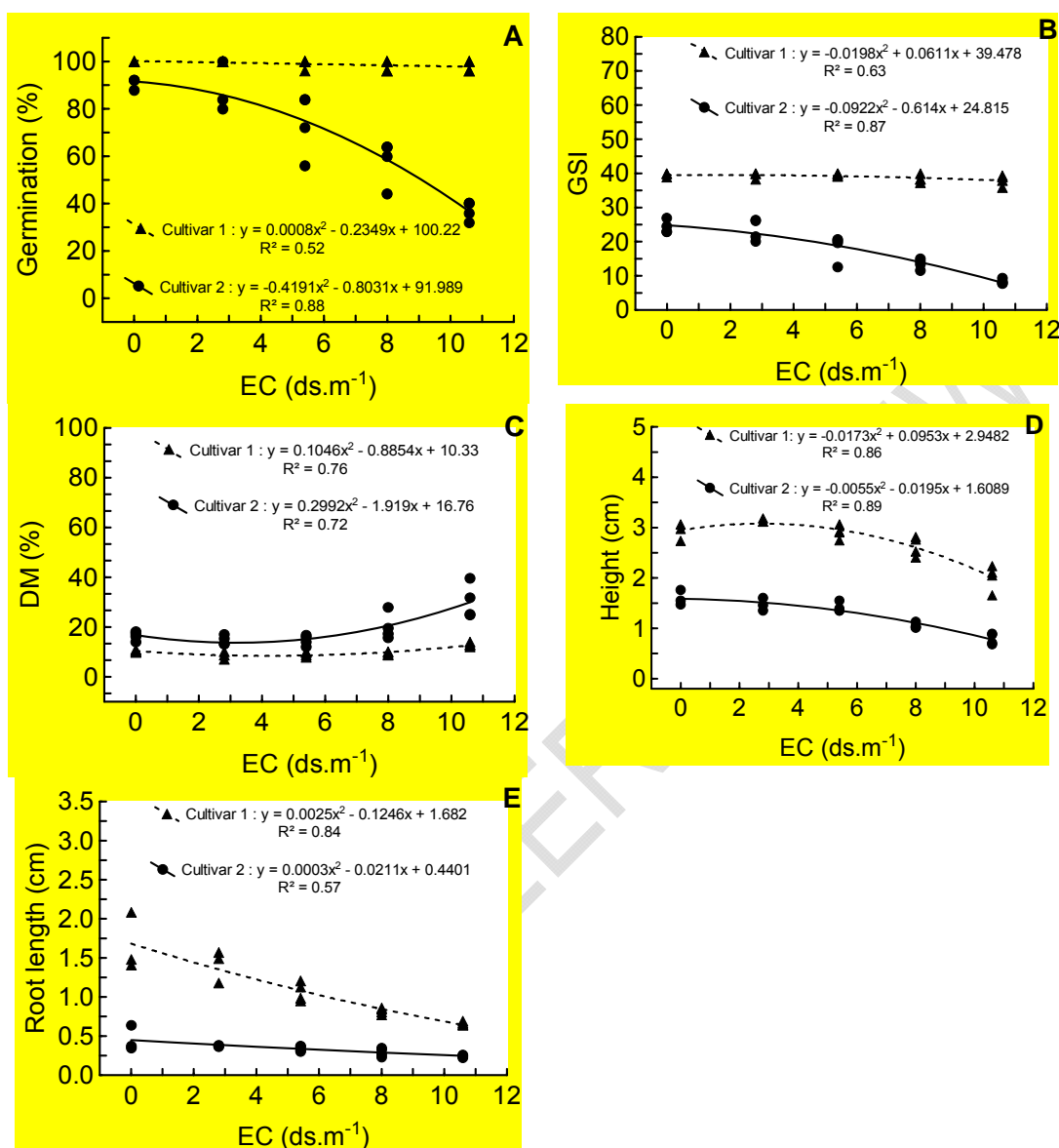


Fig. 1. Germination (A), germination speed index(B), percentage of dry matter in relation to fresh matter (C), seedling height (D) and root length (E) in function of different levels of salinity.

The linear correlation analysis (Figure 2A and 2B) showed that there is a positive correlation between root growth and shoot growth in both cultivars. It shows that, despite the negative influence of salinity on seedling development, the balance between these variables was not affected. Studies carried out by the authors [30,31, 32] in coffee, eucalyptus and potato cultivation, respectively, demonstrate the importance of good aerial and root system relationship for plants. Regarding lettuce cultivation, the relation of plant's root-shoot plays a fundamental role in the seedlings' formation. According to the authors [33], the increase in this ratio produces seedlings with superior quality, due to the better adhesion rates after transplanting.

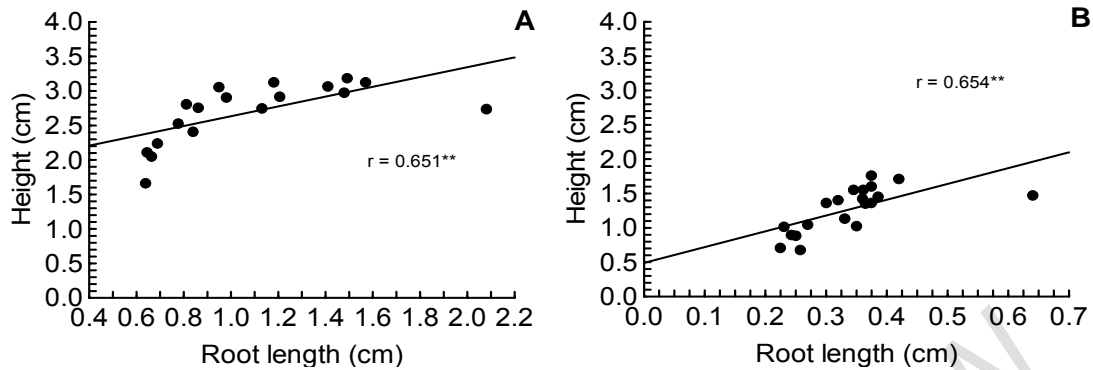


Fig. 2. Pearson's linear correlation between seedling height and root length for cultivar 1 (A) and cultivar 2 (B). * $P < 0.05$; ** $P < 0.001$.

4. CONCLUSION

Salinity affected root growth for both cultivars when EC was greater than 2.8 ds m^{-1} . For all the studied variables, cultivar 1 has shown better performance when compared to cultivar 2, allowing to affirm that it tolerates higher salinity rates in this phenological phase. The cultivar 2 was more susceptible to salinity since all the studied variables were statistically affected by different salt concentrations.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

AUTHORS' CONTRIBUTIONS

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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