

ISSN 1678-3921

Journal homepage: www.embrapa.br/pab

For manuscript submission and journal contents, access: www.scielo.br/pab

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Received October 14, 2022

Accepted March 06, 2023

How to cite

FIOREZE, S.L.; BORGA, I.; RIBEIRO, E.C. Physiological and yield parameters of wheat as affected by tiller removal and defoliation. **Pesquisa Agropecuária Brasileira**, v.58, e03156, 2023. DOI: https://doi.org/10.1590/ S1678-3921.pab2023.v58.03156. Plant Physiology/ Original Article

Physiological and yield parameters of wheat as affected by tiller removal and defoliation

Abstract – The objective of this work was to evaluate the photosynthetic parameters, yield potential, and response to defoliation of wheat (*Triticum aestivum*) plants subjected to tiller removal. Two experiments were conducted under greenhouse conditions. In the first, the two following cultivars were evaluated for complete tiller removal: TBIO Audaz and BRS 394, with a high and low tillering capacity, respectively. In the second, only 'TBIO Audaz' was subjected to detillering and defoliation at post-anthesis. Tiller removal increased the yield potential of the main stem of both tested cultivars and the CO₂ assimilation potential of the flag leaf, which was possibly a strategy to meet the demands for an increased sink strength, as evidenced by the response curves to irradiance and leaf internal CO₂ concentration. The partial defoliation of 'TBIO Audaz' increased daily CO₂ assimilation, both in intact and detillered plants. Detillered plants show a higher photosynthetic and yield potential of the main stem, but also a greater sensitivity to defoliation in the post-anthesis period.

Index terms: *Triticum aestivum*, photosynthesis, source-sink relationship, tillering.

Parâmetros fisiológicos e produtivos de trigo afetados pela remoção de perfilhos e pela desfolha

Resumo – O objetivo deste trabalho foi avaliar os parâmetros fotossintéticos, o potencial produtivo e as respostas à desfolha de plantas de trigo (Triticum aestivum) submetidas à remoção de perfilhos. Dois experimentos foram conduzidos em cultivo protegido. No primeiro, as duas seguintes cultivares foram avaliadas quanto à remoção total de perfilhos: TBIO Audaz e BRS 394, com alto e baixo potencial de perfilhamento, respectivamente. No segundo, apenas 'TBIO Audaz' foi submetida à remoção de perfilhos e à desfolha no período de pós-antese. A remoção dos perfilhos aumentou o potencial produtivo do colmo principal das duas cultivares testadas e a assimilação de CO₂ da folha bandeira, que, possivelmente, foi uma estratégia para suprir a maior demanda de drenos, como evidenciado pelas curvas de resposta à irradiância e à concentração foliar interna de CO2. A desfolha parcial de 'TBIO Audaz' aumentou a assimilação diária de CO2, tanto em plantas intactas quanto nas com perfilhos removidos. As plantas submetidas à remoção de perfilhos apresentam maior potencial fotossintético e produtivo do colmo principal, mas, também, maior sensibilidade à desfolha no período de pósantese.

Termos para indexação: *Triticum aestivum*, fotossíntese, relação fontedreno, perfilhamento.



Introduction

Wheat (*Triticum aestivum* L.) yield is limited by sink capacity (Lawlor & Paul, 2014; Borrill et al., 2015), being influenced by the ability of plants to allocate resources and by the size and number of sinks (Smith et al., 2018). Therefore, cereal yield can be increased by modifying sink capacity through genetic improvement or management practices (Furbank et al., 2019).

A promising approach to enhance sink strength in wheat plants is the use of signaling molecules, such as trehalose-6-phosphate, to increase sugar allocation in reserve tissues (Griffiths et al., 2016). Tiller inhibition (*tin*) mutants (Kebrom et al., 2012; Kebrom & Richards, 2013; Hendriks et al., 2016) and artificial tiller removal (Guo & Schnurbusch, 2015; Fioreze et al., 2020, 2021) can also be used to increase the yield potential of the ear and provide important information about the effect of inter-stem competition on source-sink relationships.

Researches on *tin* lines have shown that tiller suppression improves main stem development and yield (Mitchell et al., 2013; Hendriks et al., 2016), enhancing the strength of spike sinks. Dreccer et al. (2013) added that low-tillering wheat genotypes have high concentrations of water-soluble carbohydrates in the main stem, which may explain the higher yield potential of individual spikes in detillered plants (Guo & Schnurbusch, 2015; Fioreze et al., 2019). According to Furbank et al. (2019), this finding is related to the competition between sinks already in the early growth stages of wheat plants, when tillering represents an intense process of competition between the main stem and tillers for water, nutrients, and light, as well as for photoassimilates.

Despite these important results, there are no known studies on the photosynthetic behavior of detillered plants after anthesis and the impact of detillering on grain filling. Defoliation simulated after anthesis has shown that there may be a long period of limited assimilate availability for grain filling in plants, forcing those with normal tillering to remobilize carbon metabolites from stems or to improve ear photosynthetic activity (Zhang et al., 2020).

The objective of this work was to evaluate the photosynthetic parameters, yield potential, and response to defoliation of wheat plants subjected to tiller removal.

Materials and Methods

Two experiments were conducted under greenhouse conditions during the winter of the 2019 growing season in the municipality of Curitibanos, in the state of Santa Catarina, Brazil (27°16'26.55"S, 50°30'14.41"W, at 988 m altitude). The experimental design was a randomized complete block with four replicates, in a 2x2 (experiment 1) and 2x3 (experiment 2) factorial arrangement.

In experiment 1, the following two wheat cultivars with contrasting tiller capacities were evaluated: TBIO Audaz and BRS 394, with a high and low capacity, respectively. Both cultivars were subjected to two treatments: free tillering (control, with no detillering) and complete detillering of the main stem. For detillered plants, all late-emerging tillers were manually removed. In experiment 2, only 'TBIO Audaz' was evaluated, being subjected to detillering (control and detillering) and defoliation at post-anthesis (control, partial defoliation, and total defoliation). Leaves were removed at the beginning of plant anthesis, and only the flag leaf was maintained for partial defoliation.

The plants used in the study were grown in 3.6 L plastic pots filled with a Cambissolo Háplico (Santos et al., 2018), i.e., a Haplic Inceptisol (Soil Survey Staff, 2014) of clayey texture (550 g kg⁻¹ clay), limed with 1.51 g dm⁻³ limestone at 40 days before sowing. A base fertilizer, consisting of 120 mg dm⁻³ potassium chloride $(60\% \text{ K}_2\text{O})$ and 2.16 g dm⁻³ triple superphosphate (42%) P_2O_5), was mixed with the soil. In each pot, four seeds were sown at a 3.0 cm depth, and, after emergence, seedlings were thinned to two. Side dressing nitrogen fertilization was carried out every 15 days between emergence and anthesis, using urea (45% nitrogen) applied in solution (25 mg dm⁻³ N) to reach 150 mg dm⁻³ N. Soil moisture was maintained close to field capacity throughout the growing period by manual irrigation. The maximum temperature of the greenhouse was set to 30°C with natural irradiation. The propiconazole fungicide was applied between the jointing and flowering stages as a protective management of foliar diseases.

Gas exchange was measured at anthesis using the LI-6400XT portable photosynthesis meter (LICOR Biosciences Lincoln, NE, USA). The curves of the responses to photosynthetic light (A_n versus I) and CO₂ (A_n versus C_i) were plotted in experiment 1, where A_n is the net photosynthetic rate, I is irradiance, and

 C_i is leaf internal CO₂ concentration. Light-response curves were obtained by varying *I* from 1,800 to 0 µmol m⁻² s⁻¹, at 25°C, in a fixed CO₂ concentration of 400 µmol mol⁻¹. The linear portion of the response curve was subjected to the linear regression analysis in order to determine apparent quantum yield and light compensation point according to Habermann et al. (2003).

The A_n versus C_i curve was determined at 1,600 µmol m⁻² s⁻¹ and 25°C, with measurements starting at 400 µmol mol⁻¹ CO₂. Once the steady state was reached, the CO₂ concentration was gradually decreased to 50 µmol mol⁻¹ and, then, increased stepwise to 1,500 µmol mol⁻¹. CO₂ assimilation and the corresponding internal CO₂ values for the linear portion of the response curve were subjected to the linear regression analysis in order to determine carboxylation efficiency (Farquhar & Sharkey, 1982). In addition, CO₂ assimilation values at a leaf external (AC_e) and internal (AC_i) CO₂ concentration of 400 µmol mol⁻¹ were used to determine the relative effect of stomatal resistance on photosynthesis (S) as described in Farquhar & Sharkey (1982).

Since wheat is markedly known as a sink-limited plant (Lawlor & Paul, 2014; Borrill et al., 2015), in experiment 2, defoliation aimed to simulate a long period of limited assimilate availability for grain filing. A daily curve of net carbon assimilation was plotted in the anthesis stage (except for fully defoliated plants) at ten days after defoliation. During measurements, the CO₂ concentration was fixed at 400 µmol mol⁻¹. Photon influx to the chamber was adjusted according to external *I* values at 6:00 a.m., 9:00 a.m., noon, 3:00 p.m., and 6:00 p.m. The flag leaf of fully defoliated plants was collected and evaluated for length and width.

In the two experiments, plants were harvested at maturity and the main stems were individually evaluated for the following morphological and yield parameters: rachis length, number of spikelets, number of fertile spikelets, number of grains, grain weight, and thousand-grain weight. Basal stem diameter and peduncle diameter were determined only in experiment 2.

Data were subjected to the analysis of variance by the F-test at 5% probability, and means were compared by Tukey's test at 5% probability using the SISVAR software (Ferreira, 2011).

Results and Discussion

In experiment 1, tiller removal increased the photosynthetic response of the two wheat cultivars under high radiation (Figure 1). The absence of intraspecific competition due to detillering resulted in higher maximum A_n values for both cultivars. By contrast, apparent quantum yield was slightly lower in detillered plants, whereas the light compensation point was higher, particularly in 'BRS 394'. Moreover, tiller removal increased carbon assimilation potential under high radiation, and, given that apparent quantum yield was little affected, there also seemed to be an increase in metabolic carbon consumption via dark respiration. According to O'Leary et al. (2017), dark respiration is highly related to plant carbon and nitrogen status.

The increase in the radiation response of detillered plants can be explained by the increase in stomatal conductance (g_s), C_i , and transpiration rate (E) (Figure 2). For both cultivars, g_s , C_i , and E increased with detillering, particularly in BRS 394. Given that the differences between treatments were observed in the region of the curve that represents a limitation of photosynthesis by CO₂ diffusion, g_s acts there as a pathway for increasing C_i values and CO₂ assimilation in detillered plants (Zhao et al., 2020).

Detillering, therefore, caused changes in plant photosynthetic activity in response to the increase in internal CO₂ concentration (A_n versus C_i) (Figure 3). 'TBIO Audaz' showed better responses to tiller removal (Figure 3 A). In the plants of both cultivars, detillering caused a higher catalytic capacity of Rubisco (Figure 3 B and D), but did not affect the AC_e , AC_i , and S parameters (Table 1).

The enhanced photosynthetic potential of the plants might be associated with a high foliar nitrogen content, mainly through the increase in the maximum carboxylation velocity of Rubisco (Cabrera-Bosquet et al., 2009). This can be a key process to explain the highest values of leaf area and photosynthetic activity in detillered wheat plants as a consequence of reduced intraspecific competition. Hendriks et al. (2016) found that an increased root/shoot ratio and foliar nitrogen content in wheat lines expressing the *tin* gene improved water availability, which is another determining factor for maintaining stomatal opening throughout the day.

Considering morphological and yield parameters, tiller removal promoted an increase in the yield

potential of the main stem in both cultivars (Table 1). Detillered plants showed a higher rachis length, grain number per spike, and grain weight per spike, regardless of the cultivar. The interaction between wheat cultivar and tiller removal affected the number of fertile spikelets and thousand-grain weight. In this case, 'TBIO Audaz' plants subjected to tiller removal had a higher number of spikelets, as well as a greater thousand-grain weight (Table 2), although the latter parameter was increased in the two cultivars due to detillering. This finding can be attributed to the higher tillering potential of 'TBIO Audaz' (seven tillers per plant, n = 4). Fioreze et al. (2020, 2021) highlighted

that, the greater the competition between main stem and tillers, the greater the effect of tiller removal.

In experiment 2, the flag leaf length of 'TBIO Audaz', determined in plants subjected to total defoliation, was significantly longer in detillered plants (Figure 4); however, leaf width was not affected. The differences between treatments were observed mainly under high irradiance, similar to the pattern found for the $A_n \times I$ curves (Figure 1). Detillered plants, whether partially defoliated or not, showed the highest daily carbon assimilation (Figure 5), whereas partially defoliated plants exhibited a higher assimilation than those of the control, regardless of the tillering treatment (without and with detillering). This finding is in alignment with that of Macedo et al. (2006), who concluded that the



Figure 1. Irradiance (*I*) versus net photosynthetic rate (A_n) of the TBIO Audaz (A and B) and BRS 394 (C and D) wheat (*Triticum aestivum*) cultivars at anthesis as affected by tiller removal (B and D). Internal figures present the apparent quantum yield (Φ) and light compensation point (Γ). **Significant at 1% probability.

photosynthetic activity of the flag leaf was adjusted to mitigate the effects of defoliation; however, the participation of the ear in CO_2 assimilation must also be considered (Zhang et al., 2020).



Figure 2. Response of stomatal conductance (g_s) , leaf internal CO₂ concentration (C_i) , and transpiration rate (E) to irradiance (I) of the TBIO Audaz (A, C, and E) and BRS 394 (B, D, and F) wheat (*Triticum aestivum*) cultivars at anthesis as affected by tiller removal.



Figure 3. Net carbon assimilation (*An*) versus leaf internal CO_2 concentration (*C*_i) (A and C) and carboxylation efficiency (B and D) of the TBIO Audaz (A and B) and BRS 394 (C and D) wheat (*Triticum aestivum*) cultivars at anthesis as affected by tiller removal. **Significant at 1% probability.

| Table | 1. Ne | et carbon | assimilation | at different | t leaf inte | rnal (AC _i) |) and | external | $(AC_{\rm e})$ | CO_2 c | oncentration | ns, relative | effect |
|--------|--------|-----------|----------------|----------------------|-------------|-------------------------|--------|----------|----------------|----------|---------------|--------------|---------|
| of sto | matal | resistanc | e on photosy | nthesis (S), | and yield | paramete | rs for | the mair | ı stem | of wh | eat (Triticun | n aestivum |) plant |
| cultiv | ars as | affected | by tiller remo | val ⁽¹⁾ . | | | | | | | | | |

| Cultivar (C) | AC_{i} | ACe | S | RL | NS | NFS | NG | GW | TGW |
|--------------------|---|---|-------|-------|-------|-------|-------|-------|-------|
| | (µmol m ⁻² s ⁻¹) | (µmol m ⁻² s ⁻¹) | (%) | (cm) | | | | (g) | (g) |
| TBIO Audaz | 46.89 | 26.15 | 44.21 | 10.6 | 23.3a | 22.1a | 84.8 | 3.4 | 37.9b |
| BRS 394 | 45.61 | 22.88 | 50.29 | 11.0 | 20.5b | 20.1b | 81.1 | 3.6 | 44.1a |
| р | 0.46 | 0.14 | 0.09 | 0.26 | 0.00 | 0.00 | 0.40 | 0.21 | 0.01 |
| Tiller removal (T) | | | | | | | | | |
| No detillering | 44.51 | 23.05 | 48.26 | 9.8b | 21.9 | 20.6b | 67.3b | 2.3b | 34.4b |
| Detillering | 48.00 | 25.97 | 46.23 | 11.8a | 21.9 | 21.6a | 98.6a | 4.7a | 47.6a |
| p | 0.07 | 0.18 | 0.54 | 0.00 | 1.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| $C \times T(p)$ | 0.89 | 0.81 | 0.64 | 0.06 | 1.00 | 0.03 | 0.47 | 0.07 | 0.02 |
| CV (%) | 7.24 | 16.31 | 13.59 | 7.35 | 5.11 | 2.84 | 10.04 | 12.18 | 8.24 |

⁽¹⁾Means followed by equal letters do not differ by Tukey's test, at 5% probability. AC_i , net carbon assimilation when the leaf internal CO₂ concentration is 400 µmol mol⁻¹ (µmol m⁻² s⁻¹); AC_e , net carbon assimilation when the external CO₂ concentration is 400 µmol mol⁻¹ (µmol m⁻² s⁻¹); RL, rachis length; NS, number of spikelets; NFS, number of fertile spikelets; NG, number of grains; GW, grain weight; TGW, thousand-grain weight; p, probability by the F-test; and CV, coefficient of variation.

As in experiment 1, high carbon assimilation rates coincided with high g_s values, which resulted in higher E values (Figure 5 C). Considering that transpiration was markedly higher in plants with an increased photosynthetic activity, water use efficiency was slightly lower in detillered plants (Figure 5 D). In a previous work, Hendriks et al. (2016) found that *tin* lineages have a higher leaf photosynthetic potential

Table 2. Effects of the interaction tiller removal \times cultivar on number of fertile spikelets and thousand-grain weight on the main stem of wheat (*Triticum aestivum*) plants⁽¹⁾.

| Cultivar | Number of fert | ile spikelets | Thousand-grain weight (g) | | | |
|--------------|----------------|---------------|---------------------------|-------------|--|--|
| | No detillering | Detillering | No detillering | Detillering | | |
| 'TBIO Audaz' | 21.3Ba | 23.0Aa | 29.0Bb | 46.8Aa | | |
| 'BRS 394' | 20.0Ab | 20.3Ab | 39.8Ba | 48.4Aa | | |
| LSD | 0.96 | 5 | 5.40 | | | |

⁽¹⁾Means followed by equal letters, uppercase in the lines and lowercase in the columns, do not differ by Tukey's test, at 5% probability. LSD, least significant difference.



Figure 4. Flag leaf length and width of the TBIO Audaz wheat (*Triticum aestivum*) cultivar as affected by tiller removal. **Significant by Tukey's test at 1% probability. ^{ns}Nonsignificant.



Figure 5. Graphs showing: A, daily net carbon assimilation (*An*); B, stomatal conductance (*gs*); C, transpiration rate (*E*); and D, photosynthetic water use efficiency (WUE) of the TBIO Audaz wheat (*Triticum aestivum*) cultivar as affected by tiller removal and partial defoliation (PD).

even under field conditions. Considering that days with *I* values greater than 1,000 μ mol m⁻² s⁻¹ were frequent in the present study, this response is of great importance for carbon accumulation, indicating a lower need for the dissipation of excess-light energy. A similar pattern was observed for daily CO₂ assimilation since detillered plants exhibited a greater daily CO₂ accumulation (Figure 5).

Grain number and weight per spike, as well as thousand-grain weight, were significantly affected by the interaction between tiller removal and defoliation (Table 3). Grain number per spike increased significantly in detillered plants, regardless of defoliation, as observed in experiment 1. Furthermore, this parameter was not influenced by defoliation in plants that were not detillered, but showed the highest values in partially-defoliated detillered plants, which did not differ from those of the control. Similar results were found for grain weight per spike. In detillered

Table 3. Morphological and yield parameters of the main stem of 'TBIO Audaz' wheat (*Triticum aestivum*) plants as affected by tiller removal and defoliation⁽¹⁾.

| Tiller removal (T) | RL (cm) | NG | GW (g) | TGW (g) | |
|---------------------|---------|-----------|--------|---------|--|
| | | | | | |
| No detillering | 12.0 | 78.2 3.02 | | 38.8 | |
| Detillering | 15.1 | 136.5 | 5.31 | 37.8 | |
| р | 0.00 | 0.00 | 0.00 | 0.61 | |
| Defoliation (D) | | | | | |
| Control | 13.1 | 112.4 | 4.63 | 40.9 | |
| Partial defoliation | 14.3 | 119.1 | 5.14 | 42.3 | |
| Total defoliation | 13.2 | 90.5 | 2.73 | 31.5 | |
| р | 0.25 | 0.00 | 0.00 | 0.00 | |
| $T \times D(p)$ | 0.11 | 0.04 | 0.00 | 0.02 | |
| CV (%) | 10.66 | 12.98 | 21.01 | 12.42 | |

⁽¹⁾RL, rachis length; NG, number of grains; GW, grain weight; TGW, thousand-grain weight; p, probability by the F-test; and CV, coefficient of variation.

plants, thousand-grain weight, grain number per spike, and grain weight per spike reduced drastically as a result of total defoliation (Table 4). However, tiller removal and defoliation did not affect thousand-grain weight and plants that were not detillered, respectively (Table 3).

Yield components were not affected by partial or total defoliation in plants that were not subjected to detillering. Interestingly, in partially defoliated plants, yield potential was maintained due to the increase in the photosynthetic activity of the flag leaf. This increased activity is probably associated with the almost twofold increase in the grain number of detillered plants, which enhanced sink strength.

An increase in yield potential was observed with the increase in the photosynthetic potential of the main stem, combined with the decreased competition for water and nutrients, as evidenced by the higher number of spikelets and initiated flowers. Similar effects were reported for wheat plants subjected to tiller removal, attributed to a greater availability of resources during flower initiation and differentiation (Guo & Schnurbusch, 2015). Furthermore, there was a significant increase in grain weight and thousand-grain weight caused by the increase in the photosynthetic activity of detillered plants at post-anthesis. In this case, the increase in thousand-grain weight can be explained by an increase in nitrogen contents, which is known to occur in the leaves and grains of tin lineages (Hendriks et al., 2016), whereas the increase in the photosynthetic activity of detillered plants can be explained by the increase in sink capacity (grain number and size). Such findings underscore the importance of increasing sink strength as a strategy to enhance source activity in wheat, as shown by Griffiths et al. (2016).

The increase in the daily CO₂ assimilation of partially defoliated plants added to the reduced grain

Table 4. Effects of the interaction tiller removal \times defoliation on number of grains, grain weight, and thousand-grain weight of the main stem of 'TBIO Audaz' wheat (*Triticum aestivum*) plants⁽¹⁾.

| Treatments | Number of | grains | Grain weig | ght (g) | Thousand-grain weight (g) | | |
|---------------------|----------------|-------------|----------------|-------------|---------------------------|-------------|--|
| | No detillering | Detillering | No detillering | Detillering | No detillering | Detillering | |
| Control | 87.0Ba | 137.8Aab | 3.47Ba | 5.80Aa | 39.8Aa | 42.0Aa | |
| Partial defoliation | 78.8Ba | 159.5Aa | 3.14Ba | 7.13Aa | 40.1Aa | 44.6Aa | |
| Total defoliation | 68.8Ba | 112.2Ab | 2.46Aa | 3.00Ab | 36.3Aa | 26.7Bb | |

⁽¹⁾Means followed by equal letters, uppercase in the lines and lowercase in the columns, do not differ by Tukey's test, at 5% probability. LSD, least significant difference.

yield in detillered plants subjected to total defoliation allow of inferring that spike yield potential is more dependent on source capacity in detillered plants. Although the present work was carried out under greenhouse conditions, this result can help to explain why wheat lines expressing the *tin* gene are more susceptible to severe water deficit at post anthesis (Moeller & Rebetzke, 2017; Houshmandfar et al., 2019), which causes a reduction in the photosynthetic metabolism of plants.

Conclusions

1. Tiller removal increases grain number and weight on the main stem of wheat (*Triticum aestivum*) plants.

2. Tiller removal enhances the photosynthetic capacity of the flag leaf.

3. Detillered wheat plants are more sensitive to total defoliation at post-anthesis.

References

BORRILL, P.; FAHY, B.; SMITH, A.M.; UAUY, C. Wheat grain filling is limited by grain filling capacity rather than the duration of flag leaf photosynthesis: a case study using *NAM* RNAi Plants. **PLoS ONE**, v.10, e0134947, 2015. DOI: https://doi.org/10.1371/journal.pone.0134947.

CABRERA-BOSQUET, L.; ALBRIZIO, R.; ARAUS, J.L.; NOGUÉS, S. Photosynthetic capacity of field-grown durum wheat under different N availabilities: a comparative study from leaf to canopy. **Environmental and Experimental Botany**, v.67, p.145-152, 2009. DOI: https://doi.org/10.1016/j.envexpbot.2009.06.004.

DRECCER, M.F.; CHAPMAN, S.C.; RATTEY, A.R.; NEAL, J.; SONG, Y.; CHRISTOPHER, J.J.T.; REYNOLDS, M. Developmental and growth controls of tillering and watersoluble carbohydrate accumulation in contrasting wheat (*Triticum aestivum* L.) genotypes: can we dissect them? **Journal of Experimental Botany**, v.64, p.143-160, 2013. DOI: https://doi.org/10.1093/jxb/ers317.

FARQUHAR, G.D.; SHARKEY, T.D. Stomatal conductance and photosynthesis. **Annual Review of Plant Physiology**, v.33, p.317-345, 1982. DOI: https://doi.org/10.1146/annurev. pp.33.060182.001533.

FERREIRA, D.F. Sisvar: a computer statistical analysis system. **Ciência e Agrotecnologia**, v.35, p.1039-1042, 2011. DOI: https://doi.org/10.1590/S1413-70542011000600001.

FIOREZE, S.L.; DRUN, R.P.; WUADEN, A.F.; MAZZUCO, V.; OLIVEIRA, J.C. de. Source-sink relationships of wheat plants obtained by the application of systemic herbicides. **Pesquisa Agropecuária Brasileira**, v.56, e01600, 2021. DOI: https://doi.org/10.1590/S1678-3921.pab2021.v56.01600.

FIOREZE, S.L.; MICHELON, L.H.; TUREK, T.L.; DRU, R.P.; DALORSALETA, J.C.S. Role of nonproductive tillers as transient sinks of assimilates in wheat. **Bragantia**, v.79, p.180-191, 2020. DOI: https://doi.org/10.1590/1678-4499.20190365.

FIOREZE, S.L.; VACARI, J.; TUREK, T.L.; MICHELON, L.H.; DRUN, R.P. Componentes produtivos do trigo em função da temperatura no período de diferenciação de espiguetas. **Revista de Ciências Agroveterinárias**, v.18, p.24-32, 2019. DOI: https://doi.org/10.5965/223811711812019024.

FURBANK, R.T.; JIMENEZ-BERNI, J.A.; GEORGE-JAEGGLI, B.; POTGIETER, A.B.; DEERY, D.M. Field crop phenomics: enabling breeding for radiation use efficiency and biomass in cereal crops. **New Phytologist**, v.223, p.1714-1727, 2019. DOI: https://doi.org/10.1111/nph.15817.

GRIFFITHS, C.A.; SAGAR, R.; GENG, Y.; PRIMAVESI, L.F.; PATEL, M.K.; PASSARELLI, M.K.; GILMORE, I.S.; STEVEN, R.T.; BUNCH, J.; PAUL, M.J.; DAVIS, B.G. Chemical intervention in plant sugar signalling increases yield and resilience. **Nature**, v.540, p.22-29, 2016. DOI: https://doi.org/10.1038/nature20591.

GUO, Z.; SCHNURBUSCH, T. Variation of floret fertility in hexaploid wheat revealed by tiller removal. Journal of Experimental Botany, v.66, p.5945-5958, 2015. DOI: https://doi.org/10.1093/jxb/erv303.

HABERMANN, G.; MACHADO, E.C.; RODRIGUES, J.D.; MEDINA, C.L. CO₂ assimilation, photosynthetic light response curves, and water relations of 'Pêra' sweet orange plants infected with *Xylella fastidiosa*. **Brazilian Journal of Plant Physiology**, v.15, p.79-87, 2003. DOI: https://doi.org/10.1590/S1677-04202003000200003.

HENDRIKS, P.W.; KIRKEGAARD, J.A.; LILLEY, J.M.; GREGORY, P.J.; REBETZKE, G.J. A tillering inhibition gene influences root-shoot carbon partitioning and pattern of water use to improve wheat productivity in rainfed environments. **Journal of Experimental Botany**, v.67, p.327-340, 2016. DOI: https://doi.org/10.1093/jxb/erv457.

HOUSHMANDFAR, A.; REBETZKE, G.J.; LAWES, R.; TAUSZ, M. Grain yield responsiveness to water supply in near-isogenic reduced-tillering wheat lines – an engineered crop trait near its upper limit. **European Journal of Agronomy**, v.102, p.33-38, 2019. DOI: https://doi.org/10.1016/j.eja.2018.11.003.

KEBROM, T.H.; CHANDLER, P.M.; SWAIN, S.M.; KING, R.W.; RICHARDS, R.A.; SPIELMEYER, W. Inhibition of tiller bud outgrowth in the *tin* mutant of wheat is associated with precocious internode development. **Plant Physiology**, v.160, p.308-318, 2012. DOI: https://doi.org/10.1104/pp.112.197954.

KEBROM, T.H.; RICHARDS, R.A. Physiological perspectives of reduced tillering and stunting in the tiller inhibition (*tin*) mutant of wheat. **Functional Plant Biology**, v.40, p.977-985, 2013. DOI: https://doi.org/10.1071/FP13034.

LAWLOR, D.W.; PAUL, M.J. Source/sink interactions underpin crop yield: the case for trehalose 6-phosphate/SnRK1 in improvement of wheat. **Frontiers in Plant Science**, v.5, art. 418, 2014. DOI: https://doi.org/10.3389/fpls.2014.00418.

MACEDO, T.B.; PETERSON, R.K.D.; WEAVER, D.K. Photosynthetic responses of wheat, *Triticum aestivum* L., plants

to simulated insect defoliation during vegetative growth and at grain fill. **Environmental Entomology**, v.35, p.1702-1709, 2006. DOI: https://doi.org/10.1093/ee/35.6.1702.

MITCHELL, J.H.; REBETZKE, G.J.; CHAPMAN, S.C.; FUKAI, S. Evaluation of reduced-tillering *(tin)* wheat lines in managed, terminal water deficit environments. **Journal of Experimental Botany**, v.64, p.3439-3451, 2013. DOI: https://doi.org/10.1093/jxb/ ert181.

MOELLER, C.; REBETZKE, G. Performance of spring wheat lines near-isogenic for the reduced-tillering '*tin*' trait across a wide range of water-stress environment-types. Field Crops **Research**, v.200, p.98-113, 2017. DOI: https://doi.org/10.1016/j. fcr.2016.10.010.

O'LEARY, B.M.; LEE, C.P.; ATKIN, O.K.; CHENG, R.; BROWN, T.B.; MILLAR, A.H. Variation in leaf respiration rates at night correlates with carbohydrate and amino acid supply. **Plant Physiology**, v.174, p.2261-2273, 2017. DOI: https://doi.org/10.1104/pp.17.00610.

SANTOS, H.G. dos; JACOMINE, P.K.T.; ANJOS, L.H.C. dos; OLIVEIRA, V.Á. de; LUMBRERAS, J.F.; COELHO, M.R.; ALMEIDA, J.A. de; ARAÚJO FILHO, J.C. de; OLIVEIRA, J.B. de; CUNHA, T.J.F. **Sistema brasileiro de classificação de solos**. 5.ed. rev. e ampl. Brasília: Embrapa, 2018. 356p.

SMITH, M.R.; RAO, I.M.; MERCHANT, A. Source-sink relationships in crop plants and their influence on yield development and nutritional quality. **Frontiers in Plant Science**, v.9, art. 1889, 2018. DOI: https://doi.org/10.3389/fpls.2018.01889.

SOIL SURVEY STAFF. **Keys to soil taxonomy**. 12th ed. Washington: USDA, 2014. 360p.

ZHANG, M.; GAO, Y.; ZHANG, Y.; FISCHER, T.; ZHAO, Z.; ZHOU, X.; WANG, Z.; WANG, E. The contribution of spike photosynthesis to wheat yield needs to be considered in processbased crop models. **Field Crops Research**, v.257, art.107931, 2020. DOI: https://doi.org/10.1016/j.fcr.2020.107931.