



HAL
open science

Modeling sorghum-cowpea intercropping for a site in the savannah zone of Mali: Strengths and weaknesses of the Stics model

Amadou Traoré, Gatien Falconnier, Alassane Ba, Fagaye Sissoko, Benjamin Sultan, François Affholder

► To cite this version:

Amadou Traoré, Gatien Falconnier, Alassane Ba, Fagaye Sissoko, Benjamin Sultan, et al.. Modeling sorghum-cowpea intercropping for a site in the savannah zone of Mali: Strengths and weaknesses of the Stics model. *Field Crops Research*, 2022, 285, pp.108581. 10.1016/j.fcr.2022.108581 . hal-04391641

HAL Id: hal-04391641

<https://hal.science/hal-04391641>

Submitted on 22 Jul 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

1 **Modelling sorghum-cowpea intercropping in semi-arid sub-Saharan Africa: strengths and**
2 **weaknesses of the Stics model**

3 Amadou Traoré¹, Gatien N. Falconnier^{2,3}, Alassane Ba¹, Fagaye Sissoko¹, Benjamin Sultan⁴, François
4 Affholder^{2,3}

5 Corresponding author : Gatien.falconnier@cirad.fr

6 ¹IER (Institut d’Economie Rurale, Mali)

7 ²AIDA, Univ Montpellier, CIRAD, Montpellier, France

8 ³CIRAD, UPR AIDA, F-34398 Montpellier, France

9 ⁴ESPACE-DEV, Univ Montpellier, IRD, Univ Guyane, Univ Réunion, Univ Antilles, Univ
10 Avignon

11

12 **Abstract:**

13 Intercropping is a key entry point for sustainable intensification of cropping systems in sub-Saharan
14 Africa where variable rainfall conditions prevail. Crop simulation models can complement field
15 experiments to assess the agronomic and environmental performances of intercropping systems under
16 diverse climatic conditions, including hypothetical future climate. So far, crop models that can handle
17 intercropping, such as STICS, have not often been extensively evaluated for tropical conditions and for
18 species grown by farmers in sub-Saharan Africa. The objective of this study was to evaluate the
19 performance of the calibrated STICS model to simulate sorghum-cowpea intercropping systems in
20 rainfed conditions in West Africa. We used data from field experiments conducted at the N'Tarla
21 Agronomic Station in Mali in 2017 and 2018. Two varieties of sorghum (local and improved) with
22 contrasting photoperiod sensitivities were grown as sole crop and intercropped with cowpea, with
23 additive design. Two sowing dates and two levels of mineral fertilization were also investigated. Model
24 simulations were evaluated with observed data for phenology, leaf area index (LAI), aboveground
25 biomass, grain yield and in-season soil moisture. Large variations in aboveground biomass of sorghum
26 and cowpea was observed in the experiment (i.e. 3.5 – 9.6 t/ha for sorghum and 0.4 - 2.5 t/ha for
27 cowpea), owing to the treatments (i.e. sole vs intercrop, early vs late sowing, no fertiliser input vs
28 fertiliser input). Such variations were satisfactorily reproduced by the model, with EF of 0.81 in

29 calibration and 0.58 in evaluation (with relative RMSE of 23 and 43%). The two main observed features
30 of the intercropping system was well reproduced by the model. Firstly, cowpea and sorghum
31 aboveground biomass decreased with intercropping compared with sole cropping, and the decrease in
32 cowpea biomass was greater than the decrease in sorghum biomass. Secondly, despite a reduction in
33 sorghum and cowpea yield, Land Equivalent Ratio of the intercropping for aboveground biomass was
34 always above one. With regard to grain yield, observed LER was above one only in the non-fertilised
35 treatment. The model failed at reproducing this behaviour, probably because of insufficiently accurate
36 calibration of the process leading to grain yield formation: rRMSE for grain yield was 49% in calibration
37 and 41% in evaluation. Based on these findings, we discuss avenues to improve model calibration and
38 use the model to explore options for sustainable intensification in land constrained sub-Saharan Africa.

39 *Keywords:* Competition, sowing date, varieties, fertilization, contrasting seasons, aboveground biomass,
40 Mali

41

42 I. Introduction

43 Rainfed agriculture is the most important sector for food security in sub-Saharan Africa (Gowing and
44 Palmer, 2008). However, the region is characterized by low yield due to limited nutrients inputs and
45 weed pressure (Affholder et al., 2013). Rural farmers do not have access to inputs, and low levels of
46 investment in infrastructure generates high transaction costs that lowers the profitability of agricultural
47 activities. These constraints will likely be exacerbated by climate change: the frequency and severity of
48 extreme weather events such as drought and floods is predicted to increase (Connolly-Boutin and Smit,
49 2015; Diedhiou et al., 2018; Sultan and Gaetani, 2016).

50 West Africa must produce more to feed a rapidly growing population. Sorghum is one of the main food
51 crops in West Africa. Sorghum yield is expected to decrease by 13% due to climate change (Sultan et
52 al., 2014). Improving soil fertility and agricultural productivity while adapting to climate change have

53 therefore become the priority objectives of agricultural policies in West African countries
54 (CORAF/WECARD, 2008). In land constrained areas with limited availability of arable land and
55 pastures, intensification of agro-systems will be key. But the experience gained in developed countries
56 indicates that excessive artificialization of the environment, as in “conventional” intensification of
57 agriculture, causes environmental damages. A new form of agriculture to increase agricultural
58 productivity while preserving the ecosystem services that ensure long-term environmental sustainability
59 is needed. At cropping system level, the concept of ecological intensification was coined by to define the
60 set of principles and means necessary to increase primary productivity in the world's major cereal agro-
61 ecosystems (Cassman, 1999). Productivity increase can be achieved by capitalizing on the ecological
62 processes in agro-ecosystems (e.g. biological N₂ fixation), aiming at reducing the use of and need for
63 external inputs (Tilman and Giller, 2013). Innovative cropping systems for ecological intensification
64 need to be assessed with meaningful agronomic, environmental and economic indicators (Affholder et
65 al., 2014). Key ecological services such as carbon storage and biological diversity also need to be
66 addressed (Bonny, 2011).

67 Legumes offer good prospects to increase yield in line with the principles of ecological intensification:
68 legumes can improve soil fertility (Gbakatche et al., 2010), and provide crucial nutrients carry-overs
69 for the subsequent cereal crop (Cassman et al., 2003). Intercropping is the cultivation of two or more
70 species or varieties at the same spatial and temporal resolution (Andrews and Kassam, 1976). In
71 cereal-legume intercropping, inter-species complementarity and facilitation processes can lead to a
72 production benefit of intercropping compared with sole cropping (Li et al., 2003, 1999; Vandermeer,
73 1989). Water use efficiency is greater in intercropping than in sole cropping (Tsubo et al., 2004, Balde et
74 al., 2011). However, the expected benefits of intercropping cereals with legumes are not always
75 obtained in the context of sub-Saharan Africa due to constraints related to cultivar, sowing dates and
76 fertilization levels. Intercropping benefits may be jeopardized by competitions for light, nutrients or water
77 between the intercropped species (e.g. Baldé et al., (2011)). Shifts from competition to facilitation will

78 depend on climate, soils and crop management (Cooper et al., 2008). Appropriate and site-specific
79 recommendations for intercropping management requires replicated experiments across contrasting
80 sites. In order to account for climate inter-annual variations, the experiments would need to run over
81 several years, which is challenging in terms of time, cost and technical expertise (Knörzer et al., 2011).
82 To address such limitations, crop simulation models have been used to assess the agronomic and
83 environmental performances of cropping systems under diverse climatic conditions, including
84 hypothetical future climate (Boote et al., 1996). Crop models able to deal with intercropping systems are
85 scant. So far, few studies have been conducted on cereal-legume intercropping. Chimonyo et al. (2016)
86 evaluated the ability of the calibrated APSIM model (The Agricultural Production Systems sIMulator) to
87 reproduce sorghum-cowpea intercropping productivity and water use efficiency in potential (i.e. fully
88 irrigated) and water-stressed (i.e. rainfed) conditions, in South Africa. Despite overall promising
89 simulation accuracy for partitioning of solar radiation between the dominant (sorghum) and understory
90 crop (cowpea), the authors showed that water use efficiency of the intercrop was overestimated by the
91 model in rainfed conditions, probably because of an inadequate simulation of water stress. The latter
92 could be improved with soil calibration based on in-season water measurements. The experiment was
93 set with adequate N fertilizer inputs, so that intercrop response to varying N input was not investigated.
94 Masvaya (2019) also investigated the accuracy of APSIM model in simulating sorghum-cowpea
95 intercropping performance in semi-arid zone in Zimbabwe. The model accurately reproduced grain yield
96 of the two crops for contrasting seasons, soils and nitrogen inputs. However, the model could not be
97 evaluated for intermediary variables (e.g. LAI, soil water), leaving the risk of compensation errors. The
98 STICS model (multidisciplinary simulator for standard crops) was calibrated for wheat-pea intercrop
99 (Corre-Hellou et al., 2009). The model successfully simulated the interactions between the intercropped
100 species for nitrogen uptake but failed to accurately reproduce competition for light. So far, the STICS
101 model has not been tested for intercrops of very contrasting height, as it is the case of sorghum and
102 cowpea for example.

103 This study focuses on sorghum-cowpea intercropping. The objective was to calibrate and evaluate the
104 ability of the STICS soil-crop model to (i) simulate the growth and productivity of a sorghum-cowpea
105 intercropping, (ii) account for possible species competition and/or facilitation for the use of light, water
106 and nitrogen and (iii) test its ability to reproduce the impact of contrasting sorghum varieties, sowing
107 date and fertilization on aboveground biomass and grain yield. Data from two years (2017, 2018) from
108 an experiment conducted at the N'Tarla Agronomic Research Station in Mali were used.

109

110

111 **II. MATERIAL AND METHOD**

112 **2.1 Study Area**

113 This research was conducted at N'Tarla research station (12.6193°N, -5.689°W) in southern Mali. The
114 climate is typical of the semi-arid Sudano-Sahelian zone of West Africa. The rainy season starts in May
115 and ends in October. Average annual rainfall varies between 800 and 1000 mm. Seasonal average
116 temperature is 29 °C (Traore et al., 2013). Farming systems are mixed agro-sylvo-pastoral systems.
117 Cotton (*Gossypium hirsutum* L.) is the main cash crop and is grown in rotation with sorghum (*Sorghum*
118 *bicolor* (L) Moench), millet (*Pennisetum glaucum* (L.) R.Br.), maize (*Zea mays* L.) and legumes such as
119 groundnut (*Arachis hypogaea* L.) and cowpea (*Vigna unguiculata* (L) Walp.). Mineral fertilizer are
120 applied on cotton and maize, with nutrient carry-over benefitting the following cereals (sorghum, millet)
121 (see e.g. Ripoché et al., (2015). Cattle, goats, and sheep are raised by farmers. Large cattle herds
122 move during the dry season to wetter parts of the country (e.g. *Bougouni*, *Kadiolo*) due to local lack of
123 feed. The soils at the research station are classified as Lixisols (FAO, 2006). They have a silty sand
124 texture (10% clay) at the surface (0-35 cm), and clay content increases with soil depth (29.4% between
125 55 and 85 cm) (Table S1). Soil organic carbon is low (<0.1%) (Table 1), pH is about 6 and CEC is less
126 than 3 cmol (+) kg⁻¹ (Traore et al., 2013).

127 2.2 Experimental design

128 The experiment was conducted for two consecutive seasons (2017 and 2018) on the same field.
129 Treatments were kept on the same subplots and not re-allocated from a year to another. Two sorghum
130 varieties, namely Tiemarifing (local variety, V1) and 02-SB-F4DT “Grinka” (improved variety for grain
131 production, V2) were cultivated as sole crops, or intercropped with dual-purpose (i.e. grain and fodder
132 production) cowpea IT89KD-245 “Sangaraka”. V1 is a photoperiod-sensitive variety, and V2 is less
133 sensitive to photoperiod than V1 (Chanterreau, et al., 2013). Sorghum variety and cropping system (sole
134 vs intercrop) were the main treatments. The effect of two sowing dates for sorghum (D1-early sowing,
135 D2-late sowing) was also investigated as a secondary treatment. Sorghum early sowing was done after
136 15 June (to limit the probability of dry spell occurring after sowing) and after a rainfall greater than 20
137 mm. Early sowing (D1) occurred on 24 June in 2017 and 26 June in 2018. Late sowing (D2) of sorghum
138 occurred on 20 July in both years. Cowpea was sown two weeks after early- and late-sown sorghum. In
139 2017, N fertilizer application was 8 kg N/ha. In 2018, two levels of fertilization (F0=0 kgN/ha and
140 F1=38kgN/ha split in two applications) were investigated.

141 Sole and intercropped sorghum was planted at 0.80 m between sowing rows and 0.40 m along the
142 sowing row. Two sorghum plants per hill were kept, leading to a sorghum density of 6.25 plants m⁻² in all
143 treatments. An additive intercropping pattern was chosen where cowpea was sown in continuous line
144 between two sorghum plants (on the sorghum row) leading to a cowpea density of 6.25 plants m².
145 Weeds were controlled using hand hoe three times for D1 and twice for D2, to keep plots free from
146 weed pressure. Intercropped cowpea did not receive any specific fertilization other than that applied on
147 sorghum. Sole cowpea was cultivated without nitrogen application (F0) in 2018. In 2017 there was no
148 sole cowpea plot. Sorghum stem borer (armyworm) and cowpea pests (e.g. *aphids*, *Aphis craccivora*)
149 were controlled with chemicals to prevent damage to crops.

150 The combinations of year, sorghum variety, cropping system, fertilization and sowing date defined 25
151 cropping situations grown on 25 experimental plots where plant growth and soil characteristics were
152 monitored and used for model calibration and evaluation (Table 1).

153 **2.3 Measurements in experimental plots**

154 **2.3.1 Soil measurements**

155 In 2017, a composite sample representing the experimental field was taken at a depth of 0-20 cm before
156 the crops were planted and used to determine soil pH and soil C:N ratio. In 2018, a composite soil
157 sample was taken in each experimental plot with semi-cylindrical augers at a depth of 0-20 cm before
158 crop installation. These samples were sun-dried and sent to the laboratory for determination of total
159 soil nitrogen using the Kjeldahl method (Bremner and Mulvaney, 1982) (Table 1).

160 In 2018, soil volumetric water content was measured with a Diviner 2000 probe placed in vertical access
161 PVC tubes inserted in the soil in each experimental plot. The tubes were installed two weeks before
162 sorghum sowing. The soil-tube interface was carefully filled with fine earth to eliminate gaps at the soil-
163 tube interface, as probe measurements are very sensitive to the quality of the contact between the
164 access tube and the soil (Basinger et al., 2003). The soil was dry at the time of installation, so that the
165 planned maximum depth of 155cm was not reached in some of the experimental plots (the maximum
166 measurement depth in each plot is given in Table 1). Soil water content was measured at 10 cm
167 intervals down to the maximum depth of the tube. Measurements were taken 2-3 times per week from
168 the installation of the tubes to sorghum harvest. Plant-available water across soil profile (i.e. every 10
169 cm to the maximum depth of the tube) was calculated as the difference between soil water content
170 measured at a given time across the profile and soil water content at wilting point. In each experimental
171 plot, plant available water in the 10-cm layers was aggregated to the maximum depth to obtain total
172 plant available water.

173 **2.3.2 Plant measurements**

174 Date of flowering was recorded for sorghum and cowpea in sole and intercropping plots when 50% of
175 the plants reached the stage. Physiological maturity of sorghum was observed when 100% of the plants
176 had completed the stage of dark spots appearing at the edge of the grain attachment on the panicle. For
177 cowpea, physiological maturity was considered when the first pods started to dry. Leaf Area Index (LAI)
178 was measured using a Licor-LAI2000 (Licor, INC.) on four transects (covering three planting rows) with
179 six plants per transect on each measurement date for sole and intercropped sorghum plots (Fig. S1).
180 Measured LAI on the four transects was then averaged to get the final LAI value for the plot at a given
181 date. LAI measurements began on July 15 and continued until maturity with a 15-day interval between
182 measurements. Seven LAI measurements were made for plots with early sowing (D1) and five for plots
183 with late sowing (D2) of sorghum. Only five LAI measurements were taken under cowpea in sole crop
184 condition, because cowpea senescence started before sorghum. For intercropping there were six LAI
185 measurements in plots with D1 and four measurements in plots with D2. For the first three
186 measurements, only sorghum LAI was measured, because cowpea was too small and the LAI meter
187 could not go under its canopy without direct contact of leaves with the lens. For the remaining
188 measurements, LAI was measured (i) under cowpea canopy (to obtain total canopy LAI, i.e. sorghum
189 and cowpea), and (ii) above cowpea canopy (to estimate sorghum LAI). The LAI of intercropped cowpea
190 was estimated as the difference between the two measurements. Sorghum and cowpea aboveground
191 biomass (stem, leaves and parts of the inflorescence at flowering) was sampled in three randomly
192 selected 1 m² quadrats (corresponding to six sorghum stands and six cowpea stands) in each
193 experimental plot every two weeks after crop installation. Fresh samples of sorghum and cowpea were
194 weighted and oven-dried for 48h at 72°C for dry weight estimation. At harvest, yield and yield
195 components were determined: grain yield, grain weight, number of grains, straw weight, thousand seed
196 weight and harvest index for sorghum in 2017 and 2018. For cowpea, grain yield was measured in 2018
197 in 2017, but in 2017 the sample for dry weight estimation was lost before weighing, so that there was no
198 cowpea grain yield data for 2017. In 2018, cowpea number of grains could not be determined due to

199 destruction of the sub-sample by animals during drying on a dedicated but insufficiently protected area
200 in N'Tarla experimental station. Determination of plant nitrogen content was obtained using the Kjeldahl
201 method (Bremner and Mulvaney, 1982) applied to plant samples taken at flowering (aboveground
202 biomass) and at harvest (bulked stems and leaves, and grains) in all experimental plots.

203

204 **2.3.3 Weather data**

205 Daily climate data for the two growing seasons (2017 and 2018) was obtained from observations at the
206 N'Tarla meteorological station located 50 m from the experimental plots. Measured variables included
207 daily rainfall, daily minimum and maximum temperatures, sunshine duration, and wind speed. Global
208 radiation (R_g) was calculated as a function of latitude, maximum possible sunshine duration or daylight
209 hours and solar radiation received at the top of the Earth's atmosphere, using the following equation
210 (usually referred to as Angstrom formula (Allen et al., 1998):

$$211 \quad R_g = \left[a + \left(b \times \frac{n}{N} \right) \right] \times R_a \quad (1)$$

212 Where n is the actual sunshine duration measured in hours, N is the maximum daylight duration (Allen
213 et al., 1998). R_a is the extra-terrestrial radiation, i.e. the solar radiation received at the top of the Earth's
214 atmosphere. N and R_a were computed from latitude and date using astronomic formulas, and the
215 coefficients a and b were set at the values 0.25 and 0.50 respectively, as recommended in Allen et al.,
216 1998.

217

218 **2.4 Assessment of the performance of sorghum-cowpea intercropping**

219 Cropping system performance was evaluated for aboveground biomass and grain yield with sole and
220 intercropping, with the computation of Land Equivalent Ratio (LER). LER is the land required with sole
221 cropping to produce the yield obtained with intercropping with the same management (Willey, 1979). It

222 is a commonly used approach to assess the land use advantage associated with intercropping (Willey
223 and Rao, 1980):

$$224 \text{ LER} = Y_a/S_a + Y_b/S_b \quad (2)$$

225 Where Y_a and Y_b are the yields of intercropped sorghum and intercropped cowpea, respectively, and
226 S_a and S_b are the yields of sole sorghum and sole cowpea, respectively.

227 $p\text{LER}_a$ and $p\text{LER}_b$ are the partial LER values for each species:

$$228 \text{ pLER}_a = Y_a/S_a \quad (3)$$

$$229 \text{ pLER}_b = Y_b/S_b \quad (4)$$

230 A LER value greater than 1 indicates that there is an advantage for intercropping over sole cropping.
231 Partial land equivalent ratio ($p\text{LER}$) values are used to assess the contribution of each crop to the final
232 LER.

233 **2.5 General description of STICS soil-crop model**

234 STICS soil-crop model (Brisson et al., 2009) simulates carbon, water and nitrogen balances in the soil-
235 crop-atmosphere system. The model simulates the dynamics of agricultural and environmental variables
236 (e.g. crop yield, N in harvested organs, soil moisture and mineral N content, nitrogen leaching and soil
237 organic carbon) over agricultural seasons on a daily time step, taking into account the impact of
238 weather, and soil and crop management practices (e.g. mineral and organic fertilization, irrigation, tillage
239 and residue management). It has been designed as a generic model that can be easily adapted to
240 different crops and environmental conditions. Crops are defined using eco-physiological options (e.g.
241 effect of photoperiod and/or the requirements of cold on crop phenology) and plant parameters. Plant
242 parameters include both crop specific and cultivar-dependent parameters. The version of the model
243 used for this study is the intercropping extension of the STICS sole crop model that can consider two
244 species grown at the same time (Brisson et al., 2004). This model, hereafter referred to as "Stics-

245 intercrop” simulates the competition for light, water and nitrogen between intercropped species at a daily
246 time step (Brisson et al., 2004). In the following section, we describe the general functioning of Stics,
247 and detail the specific features of Stics-Intercrop.

248 **2.6. Description of Stics model and intercrop version specificities**

249 Plant development occurs in several stages associated with LAI and phenological stages (Table 2). The
250 duration between stages is expressed in cumulative thermal time and is specific to species and
251 varieties. Development is controlled by crop temperature (simulated variable). Germination and
252 emergence are controlled by soil temperature and moisture.

253 Crop height varies during growth, using a function relating height to LAI, and is limited to a maximum
254 value for each species (*hautmax* parameter). In the intercropping version, the comparative height of the
255 two crops determines the dominant and the understory (i.e. dominated) plant. The model is thus able to
256 simulate possible inversions in dominance between the two species during crop cycle. The sole-crop
257 model uses Beer's law to compute radiation interception. Stics-intercrop differs on that aspect and uses
258 a "Radiative transfer" formalism to compute radiation interception: the model estimates direct and
259 diffuse radiation received each day for 20 points uniformly distributed along the inter-row, and the
260 fractions intercepted by the foliage of the dominant crop and transmitted to the layer below (Brisson et
261 al., 2009). Each crop has a single extinction coefficient (*ktrou*, the same for direct and diffuse radiation).
262 Row spacing (*interrang* parameter), row orientation (*orientrang* parameter) and canopy volume impact
263 radiation transfers. The canopy volume is defined by (i) basic shapes (*form* parameter, either triangle or
264 rectangle), (ii) a ratio between the height and width of the shape (*rapform* parameter), and (iii) the base
265 height of the shape (*hautbase* parameter), considering absence of leaves between zero and base height
266 (Brisson et al., 2004). The shape and the base height are assumed to be constant throughout crop
267 cycle. Width varies as a function of height, distance between plant rows, LAI, leaf density distribution as
268 a function of height, and the *rapform* parameter. For each intercropped species, daily LAI is calculated

269 as a function of crop temperature, phenological stage, plant density, and water and nitrogen stress. The
270 LAI growth phase follows a logistic curve as a function of thermal time. Aboveground biomass daily
271 growth is computed by multiplying the intercepted radiation by a potential radiation use efficiency factor
272 that depends on the species and crop development phase (i.e. juvenile, vegetative and reproductive
273 stages). Root growth is decoupled from aboveground biomass growth. Root front growth and root
274 density are computed as a function of soil temperature, soil moisture and bulk density. Roots absorb
275 water and mineral nitrogen but allocation of biomass to root is not explicitly simulated. Water and
276 nitrogen uptake in intercropping is calculated in STICS according to root density of each crop. Root
277 densities above 0.5 cm/cm^3 of soil are not taken into account for water and nitrogen uptake since above
278 this threshold, water and nitrogen uptake are assumed not to be limited by root density (Corre-Hellou et
279 al., 2009). The simulated descent of root front is driven by thermal time using soil temperature with a
280 species-specific rate. Soil moisture content below wilting point or above field capacity can reduce or
281 stop root front growth. Root length growth is calculated by a logistic function similar to that of leaf
282 growth. Roots are distributed in the profile as a function of (i) the amount of roots already present and(ii)
283 soil constraints (drought, anoxia, penetrability). Differing with the sole crop version that uses a single
284 crop coefficient, Stics-intercrop computes the water requirements of the two species with a resistive
285 schemes applied at a daily time step to account for microclimatic effect (e.g. transfer of heat below the
286 dominant plant, see Brisson et al. (2004) for more details). The soil environment is assumed to be the
287 same for both crops, i.e. the horizontal heterogeneity of soil is ignored (but vertical heterogeneity is
288 accounted for). The interactions between the two root systems result from the influence of the soil on
289 the root profile of each crop through its penetrability and water dynamics. Soil supply is determined by
290 the balance between inputs (including possible capillary rise) and losses (soil evaporation, runoff and
291 drainage).

292 Water stress and nitrogen stress that affect plant growth and grain yield are taken into account in the
293 model using indices of reduced leaf growth and biomass accumulation under water or nutrient limiting

294 conditions (Brisson et al., 2009). Potential transpiration is a function of Leaf Area Index (LAI) and daily
295 potential evapotranspiration. The latter is as calculated using the approach of Shuttleworth & Wallace
296 (Brisson et al., 1998). Actual transpiration is calculated as the minimum between soil water supply and
297 maximal crop demand. The water stress factor that affects radiation use efficiency and plant
298 transpiration is calculated as the root transpirable fraction of soil water, which reduces growth and
299 transpiration (Brisson et al., 2009). Daily plant nitrogen accumulation depends on biomass accumulation
300 and soil nitrogen availability. Nitrogen uptake for each species depends on root depth, root distribution
301 in soil layers and crop nitrogen demand. Daily nitrogen demand is the product of plant growth rate by
302 the N amount corresponding to maximum N dilution in the plant. Soil supply is calculated per 1 cm
303 elemental layer up to maximum root system depth. The nitrogen stress factor affects (i) LAI, (ii) radiation
304 use efficiency and (iii) senescence. The N stress factor is calculated as the ratio between actual
305 nitrogen concentration and critical nitrogen concentration in the crop if the former is lower than the latter,
306 and is set to one otherwise. Stress factors are calculated daily and vary between zero (complete stress)
307 and one (no stress). A complete description of the equations and parameters governing stresses
308 definition can be found in Brisson et al. (2008). STICS also simulates N acquisition by N₂ fixation in the
309 case of legume crops. Potential symbiotic N₂ fixation by legumes is a function of nodule life cycle and
310 plant growth rate. Actual fixation depends on inhibitory effect related to sub-optimal temperature, soil
311 moisture, and/or excess of nitrate in soil (Brisson et al., 2008).

312

313 **2.7. Model parameters, model calibration and evaluation**

314 Growing seasons and the experimental treatments (cropping system, sowing date, sorghum variety and
315 fertilizer input) defined 25 cropping situations (8 in 2017 and 17 in 2018) corresponding to the
316 experimental plots listed in Table 1. Each experimental plot corresponds in this study to a "simulation

317 unit". The 17 simulation units of 2018 were used for model calibration. The remaining 8 simulations units
318 of 2017 were used for independent model evaluation.

319 **2.7.1 Setting of model parameters**

320 **Soil parameters**

321 We defined five horizons for soil parameterization: 0-15cm, 15-35cm, 35-55cm, 55-85cm, and from 85
322 to maximum sampling depth (that varied per experimental plot, see Table 1). Soil analysis carried-out in
323 the experimental plots (see section 2.3.1) informed the soil parameters "norg" (soil total Nitrogen in the
324 topsoil) (**Table 1**). pH and soil C:N ratio were set constant across all experimental plot to 5.8 and 13,
325 respectively, in line with the soil analysis carried out across all experimental plots in 2017. Soil moisture
326 at field capacity and wilting point were first estimated with pedo-transfer functions (Lidon and Francis,
327 1983) based on the texture measured in 2017 in a soil profile in one location close to the experiment
328 (**Table S1**).

329 **Plant parameters**

330 Model parameters related to crop geometry reflect the assumption that the two species in the intercrop
331 occupy distinct rows. In our experimental design, cowpea was planted on the same row as sorghum.
332 This mismatch between reality and model assumptions was easily overcome by considering that the
333 spacing between plants on a given row was the spacing between rows, and that the spacing between
334 rows was the spacing between plants on a given row (see Fig S2).

335 Model parameters related to grain yield formation were directly derived from the measurements made in
336 the experimental plots (Table 3). The greatest experimental value of grain weight among the 16
337 experimental plots was considered to inform the parameter that set maximum number of grains
338 (*nbgrmax*). Maximum height (*hautmax*) was set as measured after flowering in the experiment, i.e. 4 m
339 for sorghum and 0.5 m for cowpea. A few other parameters were taken from the literature (e.g.
340 temperature thresholds for development and photosynthesis) (Table 3).

341 2.7.2 Calibration procedure

342 STICS model has not been previously calibrated for cowpea and tropical sorghum. Initial parameter
343 values for cowpea were obtained from the spring pea plant file (Corre-Hellou et al., 2009). For sorghum,
344 initial parameters were taken from the STICS sorghum plant file as obtained with the 8.5 version of the
345 model, which corresponds to temperate fodder sorghum (Constantin et al., 2015). For sorghum, we
346 calibrated cultivar-dependent parameters for the two varieties used in the experiment.

347 To calibrate the thermal constants associated with development stage, the model was forced to go
348 through the observed dates of maximum leaf area index, flowering and maturity, in order to compute the
349 temperature sums corresponding to the intervals between the stages. We assumed that flowering
350 coincided with the beginning of grain filling. Parameters related to photoperiod (*phobase* and *phosat*) for
351 the two sorghum varieties (V1 sensitive to photoperiod and V2 less sensitive to photoperiod) were set
352 according to Traore (2015). *Phobase* and *phosat* are daylength thresholds that define the period of crop
353 growth during which a photoperiod slowing effect is applied to crop development. Photoperiod sensitivity
354 Index (*sensiphot*) was obtained by trial-and-error to minimize the gap between observed and simulated
355 flowering date (Table 3).

356 LAI is a central state variable in the model. Parameter related to leaves development (*dlaimax*) was
357 adjusted by trial-and-error to minimize the difference between observed and simulated LAI. Similarly
358 aboveground biomass growth parameters (*efcroijuv*, *efcroiveg*, *efrcoirepro*) were also adjusted by trial
359 and error to minimize the difference between the observed and simulated aboveground biomass. The
360 radiative transfer parameter "*ktrou*" used for radiation transfer from sorghum to cowpea was set to 0.30
361 for sorghum. This value was obtained using a trial-and-error calibration method to minimize the
362 differences between observed and simulated sorghum and cowpea biomass in simulation units with
363 intercropping. Default values for shape and base height (*shape*, *hautbase* parameters) as well as the
364 ratio between the thickness and width of the crop shape (*rapform* parameter) were considered.

365 The fraction of stable organic nitrogen in soil (*finert*) was set to zero. This appeared necessary for the
366 model to simulate amounts of soil mineral N and plant nitrogen uptake consistent with the observed crop
367 N uptake in the treatments without fertilization. The moisture at field capacity and at wilting point, that
368 were first set with the help of pedo-transfer functions (see section 2.7.1) were then adjusted for each
369 experimental plot (see final values in Table 1) to minimise the difference between observed and
370 simulated soil water during the growing season.

371

372 For some parameters the simple trial and error approach used above would have been poorly effective
373 and we used instead the optimization tool implemented in the software package associated with Stics
374 model (Table 3). This optimizer proposes a parameter values that minimizes the gap between model
375 simulation and observations for the variables to be explained (Buis et al., 2011), using weighted least
376 squares (Makowski et al., 2006). Lower and upper parameters limits were set before optimization to
377 ensure that the optimized values correspond to reasonable physiological values. Crop parameters were
378 first calibrated using sole sorghum and cowpea simulation units. Then, these parameters were refined
379 using intercropping simulation units. The parameters Kmabs2 (*i.e.* the affinity constant of N uptake by
380 the roots for the low uptake system) and Vmax2 (*i.e.* the maximum specific N uptake rate with the high
381 affinity transport system) were optimized to minimize the difference between observed and simulated
382 plant N uptake. Parameters related to grain formation such as the slope of the relationship between
383 grain number and growth rate (*cgrain*), the number of grains produced when growth rate is zero
384 (*cgrainv0*) were also optimized by minimizing the difference between the simulations and the
385 observations of the number of grains (Table 3). The rate of increase of the harvest index vs. time
386 (*viticarbT*) was adjusted to minimize the difference between simulations and observations for grain yield.

387 **2.7.2 Model evaluation**

388 Model error was estimated with the root means square error (*RMSE*), which has the same unit as the
389 considered variable, and its relative value in % (*rRMSE*). We used these statistical criteria to assess the

390 accuracy of STICS simulations for maximum leaf area index (*Laimax*), aboveground biomass, plant N
391 uptake and grain yield of sole and intercropped sorghum and cowpea.

$$392 \quad RMSE = \sqrt{\frac{1}{n} \sum_{i=1}^n (O_i - S_i)^2} \quad (5)$$

$$393 \quad rRMSE = \frac{RMSE}{\bar{O}} \times 100 \quad (6)$$

394 Where O_i et S_i are the observed and simulated values for the i th measurement, n is the number of
395 observations and \bar{O} is the average of the observed values.

396 Model efficiency (*EF*) was also calculated:

$$397 \quad EF = 1 - \frac{\sum_{i=1}^n (O_i - S_i)^2}{\sum_{i=1}^n (O_i - \bar{O})^2} \quad (7)$$

398 Model efficiency (*EF*) (Willmott et al., 1985) varies from infinite negative value to 1 (perfect
399 performance). A negative value indicates that the mean of observations is a better predictor than the
400 model, and a zero value indicates that the model does not outperform the mean of observed values.
401 The combined analysis of these three indicators gives a comprehensive assessment of model accuracy.

402

403 **3. Results**

404 **3.1 Weather data**

405 Compared to that of 2018, growing season in 2017 was cooler and wetter in terms of cumulated rainfall,
406 but with a greater number of dry days and greater solar radiation overall. The average minimum
407 temperature observed in 2017 (17.3°C) was 5°C lower than that observed in 2018 (22.3°C) during the
408 growing season (Figure 1). Average Maximum temperature in 2017 (34.2°C) was 2.1°C lower than
409 average maximum temperature in 2018 (36.3°C). Average growing season temperature in 2017
410 (27.9°C) was 1.1°C lower than in 2018 (30.0°C). Average global solar radiation during the growing
411 season in 2017 (23.1 MJ m⁻²) was greater than in 2018 (18.0 MJ m²). Cumulative rainfall recorded in

412 2017 (808 mm) (Fig. 1A) was 110 mm greater than in 2018 (698 mm) (Fig. 1B). On the other hand,
413 there were more dry days (i.e. days with no rainfall) in 2017 (134 days) than in 2018 (122 days).

414

415 **3.2. Model calibration**

416 **3.2.1. Sorghum and cowpea phenology**

417 Model simulations of sorghum phenology led to a RMSE of 3.4 days for beginning of grain filling and 3.1
418 days for physiological maturity (Fig. 2a, 2b). RMSE for cowpea phenology was 3.9 days for grain filling
419 and 10.8 days for physiological maturity (Fig. 2a, 2b).

420 The model reproduced the relatively high sensitivity of V1 to photoperiod reasonably well: a 24-day
421 delay in sowing (i.e. the difference between early and late sowing in the experiments) shortened the
422 observed length of the vegetative phase (from planting to start of grain filling) by 17 days. The model
423 simulated a 14-day decrease in the length of vegetative phase when sowing was delayed. V2 was less
424 sensitive to photoperiod: a 24-day delay in sowing created only a 9-day observed shortening of the
425 vegetative phase. The model simulated a 6-day decrease in the length of the vegetative phase when
426 sowing was delayed.

427

428 **3.2.2. Simulation of in-season soil water, LAI and aboveground biomass (AGB)**

429 ***Sole crops with calibration dataset***

430 In-season soil water was well reproduced by the model, especially the increase in soil water up to field
431 capacity during vegetative phase (Fig. 3). However, soil water after the end of the growth cycle (a period
432 with low rainfall and high soil evaporation) was overestimated by the model due to the underestimation
433 of soil evaporation.

434 A reasonably good agreement between model and observations for the key plant growth variables was
435 reached after calibration. However, it was not possible to calibrate LAI and biomass growth with an
436 equally satisfactory fit for the two sowing dates treatments and the two sorghum cultivars (Fig. 3). For

437 sole sorghum with early sowing date (D1) and nitrogen input (F1), LAI simulated by model matched the
438 observations for the improved variety (V2) and slightly overestimated them for the local variety (V1) (Fig.
439 3a, 3c). With late sowing (D2) and nitrogen input (F1), the model overestimated observed maximum LAI
440 for both sorghum varieties (Fig. 3b, 3d). With early sowing (D1) and without nitrogen input (F0), the
441 model overestimated observed LAI for V1 (Fig. 3e) but agreed with observed LAI for V2 under the same
442 fertilization conditions (Fig. 3g). With late sowing (D2) and without nitrogen input (F0), the model
443 overestimated LAI for both sorghum varieties (Fig. 3f, Fig. 3h). LAI simulations for cowpea in sole crop
444 matched closely the observations (Fig. 3i).

445 AGB was underestimated (especially around flowering) in simulations for both sorghum varieties with
446 early sowing (D1) and N input (F1) (Fig. 3a, Fig. 3c). With late sowing (D2) and N input (F1), AGB
447 simulations were in closer agreement with in-season and harvest observations for V1 and V2 (Fig. 3b,
448 3d). Without nitrogen supply (F0), the same trends were observed for early sowing i.e. sorghum AGB
449 was overestimated for both varieties (especially around flowering) (Fig. 3e, 3g) but underestimated at
450 harvest for late sowing (Fig. 3f, 3h). The simulation of cowpea AGB closely matched the observation
451 (Fig. 3i).

452

453 ***Intercrops with calibration dataset***

454 Simulated in-season soil water for intercrops corresponded fairly well to the observations (Fig.4 a, 4b,
455 4c, 4d, 4m, 4n, 4o, 4p). As with sole crops, soil water at the end of the season was overestimated.
456 Simulated LAI of (i) total canopy (sorghum plus cowpea), (ii) sorghum, and (iii) cowpea matched the
457 observations for variety V2 (Fig. 4q) but model simulations overestimated the observations for variety
458 V1 (Fig. 4e) with early sowing (D1) and nitrogen input (F1). With late sowing (D2) and nitrogen input
459 (F1), the model overestimated LAI of i) total canopy (sorghum and cowpea), (ii) sorghum, and (iii)
460 cowpea for V1 and V2 (Fig 4f, 4r). Without N input (F0), the model also overestimated the LAI of (i) total

461 canopy (sorghum and cowpea), (ii) sorghum, and (iii) cowpea for the two sorghum varieties, in early and
462 late sowing (Fig. 4g, 4s, 4h, 4t).

463 With early sowing (D1) and nitrogen input (F1), the model underestimated sorghum AGB throughout
464 crop cycle for V1 (Fig. 4i), and matched more closely the observations for V2 (Fig 4u). With late sowing
465 (D2) and nitrogen application (F1), AGB simulations matched the observations for both sorghum
466 varieties (Fig. 4j, 4v). With early sowing (D1) and without nitrogen input (F0), the model simulated
467 accurately AGB throughout crop cycle for V1 and V2 (Fig 4k, 4w). With late sowing (D2) and without
468 nitrogen input (F0), the model overestimated sorghum AGB at the end of crop growth (Fig. 4l, 4x). For
469 cowpea, the model accurately simulated AGB throughout the crop cycle whether grown with V1 or V2,
470 with and without N inputs (Fig. 4i, 4u, 4j, 4v, 4k, 4w, 4l, 4x).

471

472 ***Sole and intercrops in evaluation dataset***

473 In sole and intercropping conditions, though the soil was filled with water during crop early growth, water
474 stocks rapidly decreased below 50% of maximum available plant water capacity, indicating possible
475 water stress (Figure S3 and Figure S4). For all sowing dates, for sole crops and intercrops, model
476 simulations overestimated observed LAI (Fig. S3, Fig. S4). For sole crops, AGB simulations were in
477 close agreement with the observations during the season but overestimated the observations at harvest
478 for the local variety (V1) regardless of sowing dates (Fig. S3a, S3b). For the improved variety (V2),
479 model simulations of AGB were in good agreement with observations (Fig. S3c) for early sowing. With
480 late sowing, AGB was overestimated at the end of the cycle (Fig. S3d). With intercropping, AGB
481 simulation displayed similar trends as observed with sole cropping (Fig. S4, e, f, k, l).

482

483 **3.2.3 Simulation of aboveground biomass, plant N, number of grains and grain yield at harvest**

484 Sorghum and cowpea AGB at harvest varied widely in sole cropping and intercropping, due to
485 differences in sorghum variety, sowing date and fertilizer inputs: aboveground biomass ranged from 3.5

486 t/ha to 9.6 t/ha for sorghum and from 0.4 t/ha to 2.5 t/ha for cowpea for the calibration dataset. The
487 model reproduced this variability with rRMSE of 23 % and EF of 0.81 (Fig. 5a, Table 4). EF of model
488 simulation was smaller in evaluation compared with calibration (Table 4), and rRMSE was greater,
489 indicating a loss in the accuracy of model simulation of AGB with the independent calibration dataset.
490 The model underestimated plant N at harvest (Fig. 5c): rRMSE was 35% and EF was negative (-0.05).
491 With regard to grain number per m² of sorghum (Fig. 5e), rRMSE was 29% (Table 4), and remained
492 fairly stable with evaluation dataset (rRMSE = 25%). The variability in observed grain yield across crops
493 and experimental treatments was well reproduced by the model (Fig. 5g). Although rRMSE was large
494 (49%), EF was 0.56 (Table 4, Fig.5). Model accuracy in simulating grain yield remained fairly stable with
495 the evaluation dataset (rRMSE = 14% with the evaluation dataset), though EF became negative (Table
496 4).

497

498

499

500

501 **3.3 Simulation of the impact of intercropping, fertilizer and variety on crop yield and LER**

502 Observed sorghum and cowpea grain yield decreased with intercropping compared with sole cropping
503 (see figure 6a, 6c). The model accurately depicted this trend (see Figure 6b, 6d), though it
504 underestimated the greater negative impact of intercropping that was observed for cowpea grain yield.
505 In the experiment, the improved variety (V2) outperformed the traditional variety (Figure 6e) for grain
506 yield, whereas the two varieties did not show marked differences in AGB yield (Figure S5e). The model
507 successfully simulated this trend (Figure 6f). AGB of sorghum increased with N input, but not cowpea
508 biomass (see Figure S5i, k), and this trend was also adequately represented by the model (Figure S5j,
509 l). Grain yield of sorghum also increased with N input, while cowpea yield decreased with N input
510 (Figure 6i, 6k), and this was also well depicted by the model, although the model overestimated

511 sorghum and cowpea grain yield with no N input (Figure 6j, 6l). AGB and grain yield of sorghum and
512 cowpea decreased with late sowing compared with early sowing (Figure 6m, 6o). The model failed to
513 reproduce such a drop in yield with late sowing (see Figure 6n, Figure 6p). AGB and grain yield of
514 cowpea and sorghum were smaller in 2017 than in 2018 (Figure 6q, 6s and Figure S5q, S5s). The
515 model reproduced such behaviour, although the simulated effect of year on the AGB and grain yield of
516 cowpea and sorghum was less marked than in the reality (Fig 6r, 6t and Figure S5r, S5t).

517 There was an interaction between cropping system (sole vs intercropping) and growing season. In 2017,
518 the observed negative impact of intercropping on sorghum yield was weaker than in 2018 (Fig S6). The
519 model failed in reproducing such effect and it simulated a more pronounced detrimental effect of
520 intercropping on sorghum yield in 2017 than in 2018. In 2018, the model attributed sorghum and
521 cowpea yield decrease with intercropping to competition for nitrogen: while water stress was identically
522 inexistant in sole- and intercropping, the simulated N stress for sorghum during reproductive phase was
523 stronger (i.e. lower indicator value) in intercropping than in sole cropping (see Figure S7). In 2017, the
524 model simulated a relatively strong water stress during reproductive stage for both intercropping and
525 sorghum sole crop, but this did not impacted LAI as strongly as in the reality and translated into a strong
526 simulated yield reduction only in the intercropping situation whereas the simulated yield of sole sorghum
527 was at the same level as in 2018 where no water or nitrogen stress was simulated (Figure S6, S7).

528

529 Despite the decrease in both sorghum and cowpea yield with intercropping (i.e. partial LER below one),
530 overall observed intercropping LER was always above one, with regard to AGB (Figure 7A). The model
531 proved accurate in simulating this behavior, as simulated LER was always above one (Figure 7B).
532 Decrease in observed cowpea AGB with intercropping was greater than the decrease in sorghum AGB,
533 and the model was satisfactorily depicting this trend. With regard to grain yield, LER was above one
534 without N input, and below one with N input (due to a stronger decrease in cowpea grain yield) (Figure

535 7C). Because the model did not capture the stronger decrease in cowpea grain yield with N input and
536 intercropping, the simulations could not reproduce these LER values below one (Figure 7D).

537

538 **4. Discussion**

539

540 *Promising features of the calibrated model*

541 The calibrated intercrop model reproduced the key feature of the tested additive intercropping: LER for
542 AGB was always above one, thanks to a moderate decrease in sorghum AGB and despite a more
543 pronounced decrease in cowpea AGB. This decrease in cowpea AGB with intercropping was probably
544 linked to the competition for solar radiation that is primarily intercepted by sorghum canopy. This gives
545 confidence in the ability of the Stics intercrop model to predict the behaviour of intercropping based on
546 two plant species with contrasting heights. Chimonyo et al., (2016) found a similar promising feature of
547 the APSIM model.

548 The grain yield advantage of the improved variety over the local variety, and the yield advantage of N
549 input over no fertilization, were also fairly well simulated. The model simulated less accurately but
550 reasonably well the observed differences in yield resulting from the differences in water stress
551 occurrences between our two experimental years, under intercropping conditions. The model can help
552 identify complex interactions between growing season characteristics, N inputs, and the competitions at
553 stake in the intercropping system. While competition for water clearly prevailed in the intercropping
554 situations of 2017, the simulations suggested that competition for nitrogen occurred in the intercropping
555 situations in 2018, with clearly less water stress. Such competition for nitrogen can be surprising -
556 because cowpea is a legume that fixes atmospheric nitrogen. Cowpea did fix nitrogen in the experiment.
557 The observed N uptake for sole sorghum without N fertilizer was 90 kgN/ha (i.e. which indicates that at
558 least 90 kgN/ha was provided by the mineralization of soil organic matter). Sole cowpea with no fertiliser
559 input achieved a N uptake above 200 kgN/ha. This indicates a possible N₂ fixation of at least 110 kgN
560 ha⁻¹ (on top of the 90 kg N/ha provided by the soil organic matter mineralisation). However, N

561 competition can still occur when mineral fertilizer are applied. High level of soil mineral N can depress
562 the amount of atmospheric N₂ that is fixed by the legume (Sprent et al., 1988). This decrease in legume
563 N₂ fixation with mineral fertilizer input has been observed in wheat-pea and barley pea-experiments
564 (Bedoussac and Justes, 2010; Corre-Hellou et al., 2009), and this process is accounted for by the Stics
565 model (Brisson et al., 2009). Detailed measurement of the amount of N₂ fixed by the legume (using ¹⁵N
566 natural abundance method), in sole and intercropping, would be required to validate this simulated
567 competition for nitrogen in 2018.

568

569 *Avenues to improve model calibration*

570 Despite the promising features of the model, prediction error of crop AGB and grain yield remained
571 rather large (in the range 29-49%), compared with other studies. Chimonyo et al. (2016) obtained
572 rRMSE values for grain yield and aboveground biomass in the range 6-15% when simulating sorghum-
573 cowpea intercropping in South Africa. Masvaya (2019) obtained rRMSE for grain yield in the range 18-
574 29% when simulating grain yield of sorghum and cowpea with sole and intercropping in semi-arid
575 Zimbabwe.

576 In this study, the calibrated intercrop model failed to accurately predict the low yield of cowpea with
577 intercropping, and thus overestimated LER for grain. In Stics, the number of grain (a key yield
578 component) is determined by biomass growth during a short period before the beginning of grain filling
579 (Falconnier et al., 2019). Accurately setting the length of this period, and set the parameters that define
580 the relation between biomass growth and the final number of grain requires measurement of biomass
581 growth during that period, along with measurement of the final number of grain, for a range of
582 contrasting experimental conditions (e.g. with/without water and nitrogen stress) (Affholder et al., 2013).
583 The latter was missing in our experimental setting, and more detailed measurement of biomass growth
584 and grain number would certainly be necessary to improve model accuracy. The resulting robustness of
585 yield prediction under water stress during reproductive stage was insufficient to predict correctly the

586 impact of water stress in some cropping situations. Caution should therefore be taken in the
587 interpretation of the simulated inter-annual variation in grain yield of intercropping when using historical
588 weather records and/or future climatic series. The model also overestimated grain yield of sorghum and
589 cowpea with no N input. This is possibly related to the poor simulation of N uptake by both sorghum and
590 cowpea. Measurement of initial soil mineral N prior to sowing and during crop growth would possibly
591 help improve the calibration of the mineralization of soil organic matter and crop N uptake. These
592 improvements in model calibration with regard to yield component and yield with no N input are required
593 to improve our confidence in the prediction of LER for grain. This will be crucial if the model is to be
594 used for virtual experiments where sustainable intensification pathways with intercropping are tested.
595 Late planting of sorghum resulted in smaller yield, and the model could not reproduce this behaviour.
596 Late sowing of photoperiodic plant decrease the length of their growing cycle, thus directly lowering the
597 amount of radiation intercepted and the final yield (Tovignan et al., 2016). Such effect should be smaller
598 with the less-photoperiodic variety for which late planting does not translate into shorter growing cycle.
599 But the improved variety was also impacted by late sowing. Possibly, the late plantings were negatively
600 impacted by excess water in the plots (anoxia), a process that can be taken into account by the Stics
601 model, but that we did not calibrate due to lack of appropriate data on soil water conductivity under
602 saturated conditions. Low nitrogen availability due to runoff (sowing during the period of intensive
603 rainfall), or armyworm damage on sorghum could also possibly contribute to the discrepancy between
604 observations and model simulations. Armyworm damage was observed on sorghum plants during the
605 2018 season but their impact on yield was not assessed. The Stics model does not simulate the impact
606 of pests on the crop.

607

608 *A calibrated intercrop model to explore options for sustainable intensification in land constrained sub-*
609 *Saharan Africa*

610

611 Our experimental design investigated the crucial features of sustainable intensification, namely the
612 integration of legumes with intercropping, the use of mineral fertilizer, combined with improved varieties
613 (Vanlauwe et al., 2014). The calibrated intercrop model was able to mimic the key main effects and
614 interactions at play when investigating the performance of these options. Hence, it can be a crucial tool
615 to explore pathways toward sustainable intensification of agriculture in land constrained sub-Saharan
616 Africa. Intercropping is generally a better option with low fertilizer inputs (Bedoussac et al., 2015; Yu et
617 al., 2015). Our experimental results also support such claim. This contrasts with the now widely
618 acknowledged need for sustainable intensification of agriculture with more mineral fertilizer in sub-
619 Saharan Africa (Jayne et al., 2018). The predicted increase in climate variability and climate change in
620 sub-Saharan Africa (Sultan and Gaetani, 2016) may alter such picture, as intercropping is also well
621 known to stabilize yield (Weih et al., 2021). Crop models and crop models ensembles are great tools to
622 investigate the interactions between crop management and climate variability and change in sub-
623 Saharan Africa (Falconnier et al., 2020). The increasing availability of promising calibrated intercrop
624 model (e.g. this study, Chimonyo et al. (2016); Masvaya (2019)) offer the prospect of ensemble
625 simulation of the performance of intercropping in the face of climate variability and change, in order to
626 guide the design of adaptive cropping system involving intercropping.

627

628 **5. Conclusion**

629 Our study showed that the locally calibrated intercropping version of the STICS soil-crop model
630 displayed some promising features to explore the performance of sorghum-cowpea intercropping under
631 tropical rainfed environment. In the experiment, sorghum and cowpea aboveground biomass and yield
632 decreased with intercropping compared with sole cropping, but observed LER for aboveground biomass
633 remained above one, regardless of experimental treatment (fertiliser, sowing date and sorghum variety).
634 The calibrated model accurately depicted this overall trend related to LER for aboveground biomass,
635 and satisfactorily accounted for the impact of fertiliser and sorghum variety. It can therefore be a useful

636 tool to understand the competitions that occur between the intercropped species in this additive cereal-
637 legume intercropping systems in tropical rainfed environment, under the framework of sustainable
638 intensification and integrated soil fertility management. Despite this promising feature of the model,
639 large inaccuracies in the simulated leaf area index and N uptake were observed. The detrimental impact
640 of late sowing on sorghum yield, possibly because of anoxia, was also poorly accounted for. Though the
641 model can be used to explore the impact of inter-annual climate variability on intercropping
642 performance, these limitations will constraint the ability of the model to accurately and meaningfully
643 portray the interactions at stake when crop management varies strongly. We therefore advocate for
644 increased, continuous and detailed experimental effort on cereal-legume intercropping systems (e.g.
645 measurement of soil mineral N, N₂ fixation, and grain number for a great range of contrasted cropping
646 situations), in order to improve the calibration of parameters related to water and N stress. This will
647 improve model ability to deal with intercropping, a key management option for sustainable intensification
648 of smallholder farming in the global South.

649 **Acknowledgement:**

650 The Agriculture and Climate Risk Management program: Tools and Research in Africa "Agricora" and
651 the "Eco-Fert-Clim" project funded the field work of this study. The authors are grateful to Remi Vezy
652 who provided useful scripts and R functions that greatly help analyze simulation outputs, and also to
653 Eric Justes for useful insights on intercrop model calibration.

654

655 **Reference**

- 656 Affholder, F., Parrot, L., Jagoret, P., 2014. Acquis et perspectives de l'intensification écologique, in:
657 Sourrisseau, J.-M. (Ed.), *Agricultures familiales et mondes à venir*. Versailles, France, pp. 303–
658 316.
- 659 Affholder, F., Poeydebat, C., Corbeels, M., Scopel, E., Tiftonell, P., 2013. The yield gap of major food
660 crops in family agriculture in the tropics: Assessment and analysis through field surveys and
661 modelling. *Field Crops Res.* 13.
- 662 Allen, R.G., PEREIRA, L.S., RAES, D., SMITH, M., 1998. Crop evapotran-spiration-Guidelines for
663 computing crop water requirements-FAO Irrigation and Drainage Paper N° 56. FAO 1–15.

664 Andrews, D.J., Kassam, A.H., 1976. The Importance of Multiple Cropping in Increasing World Food
665 Supplies, in: Papendick, R.I., Sanchez, P.A., Triplett, G.B. (Eds.), ASA Special Publications.
666 American Society of Agronomy, Crop Science Society of America, and Soil Science Society of
667 America, Madison, WI, USA, pp. 1–10. <https://doi.org/10.2134/asaspecpub27.c1>

668 Baldé, A.B., Scopel, E., Affholder, F., Corbeels, M., Da Silva, F.A.M., Xavier, J.H.V., Wery, J., 2011.
669 Agronomic performance of no-tillage relay intercropping with maize under smallholder
670 conditions in Central Brazil. *Field Crops Res.* 124, 240–251.
671 <https://doi.org/10.1016/j.fcr.2011.06.017>

672 Basinger, J.M., Kluitenberg, G.J., Ham, J.M., Frank, J.M., Barnes, P.L., Kirkham, M.B., 2003. Laboratory
673 Evaluation of the Dual-Probe Heat-Pulse Method for Measuring Soil Water Content. *Soil Sci.*
674 *Soc. Am.* 2, 389–399. <https://doi.org/10.2136/vzj2003.3890>

675 Bedoussac, L., Journet, E.-P., Hauggaard-Nielsen, H., Naudin, C., Corre-Hellou, G., Jensen, E.S., Prieur,
676 L., Justes, E., 2015. Ecological principles underlying the increase of productivity achieved by
677 cereal-grain legume intercrops in organic farming. A review. *Agron. Sustain. Dev.* 35, 911–
678 935.

679 Bedoussac, L., Justes, E., 2010. The efficiency of a durum wheat-winter pea intercrop to improve yield
680 and wheat grain protein concentration depends on N availability during early growth. *Plant*
681 *Soil* 330, 19–35. <https://doi.org/10.1007/s11104-009-0082-2>

682 Bonny, S., 2011. Ecologically intensive agriculture: Nature and challenges. *Cah. Agric.* 20, 451–462.
683 <https://doi.org/10.1684/agr.2011.0526>

684 Boote, K.J., Jones, J.W., Pickering, N.B., 1996. Potential Uses and Limitations of Crop Models. *Agron.*
685 *J.* 88, 704–716. <https://doi.org/10.2134/agronj1996.00021962008800050005x>

686 Bremner, J.M., Mulvaney, C.S., 1982. Nitrogen-Total, in: Klute, A. (Ed.), *Methods of Soil Analysis Part*
687 *2.* American Society of Agronomy, Soil Science Society of America, Madison, WI, USA, pp.
688 595–624. <https://doi.org/10.2134/agronmonogr9.2.2ed.c31>

689 Brisson, N., Bussi re, F., Ozier-Lafontaine, H., Tournebize, R., Sinoquet, H., 2004. Adaptation of the
690 crop model STICS to intercropping. Theoretical basis and parameterisation. *Agron. EDP Sci.*
691 24, 409–421. <https://doi.org/10.1051/agro:2004031>. hal-00886023

692 Brisson, N., Launay, M., Mary, B., Beaudoin, N., 2009. Conceptual Basis, Formalisations and
693 Parameterization of the Stics Crop Model. Editions Quae, Paris.

694 Brisson, N., Mary, B., Ripoche, D., Jeuffroy, M.H., Ruget, F., Nicoullaud, B., Gate, P., Devienne-Barret,
695 F., Antonioletti, R., Durr, C., Richard, G., Beaudoin, N., Recous, S., Tayot, X., Plenet, D., Cellier,
696 P., Machet, J.-M., Meynard, J.M., Del colle, R., 1998. STICS: a generic model for the
697 simulation of crops and their water and nitrogen balances. I. Theory and parameterization
698 applied to wheat and corn. *Agronomie* 18, 311–346. <https://doi.org/10.1051/agro:19980501>

699 Buis, S., Wallach, D., Guillaume, S., Varella, H., Lecharpentier, P., Launay, M., Gu rif, M., Bergez, J.-E.,
700 Justes, E., 2011. The STICS Crop Model and Associated Software for Analysis,
701 Parameterization, and Evaluation, in: Ahuja, L.R., Ma, L. (Eds.), *Advances in Agricultural*
702 *Systems Modeling.* American Society of Agronomy and Soil Science Society of America,
703 Madison, WI, USA, pp. 395–426. <https://doi.org/10.2134/advagriscystmodel2.c14>

704 Carsky, R.J., Douthwaite, B., Manyong, V.M., Sanginga, N., Schulz, S., 2003. Am lioration de la gestion
705 des sols par l’introduction de l gumineuses dans les syst mes c r aliers des savanes
706 africaines. *Cah. Agric.* 12, 227–233.

707 Cassman, K.G., 1999. Ecological intensification of cereal production systems: Yield potential, soil
708 quality, and precision agriculture. *Proc. Natl. Acad. Sci.* 96, 5952–5959.
709 <https://doi.org/10.1073/pnas.96.11.5952>

710 Chantereau, J., Fran ois Cruz, J., Ratnadass, A., Trouche, G., 2013. *Le Sorgho*, Qu e, ed. CTA, Postbus
711 380, 6700 AJ Wageningen, Pays-Bas, Versailles-Gembloux.

712 Chimonyo, V.G.P., Modi, A.T., Mabhaudhi, T., 2016. Simulating yield and water use of a sorghum–
713 cowpea intercrop using APSIM. *Agric. Water Manag.* 177, 317–328.
714 <https://doi.org/10.1016/j.agwat.2016.08.021>

715 Connolly-Boutin, L., Smit, B., 2015. Climate change, food security, and livelihoods in sub-Saharan
716 Africa. *Clim. Change* 385–399. <http://dx.doi.org/10.1007/s10113-015-0761-x>

717 Constantin, J., Willaume, M., Murgue, C., Lacroix, B., Therond, O., 2015. The soil-crop models STICS
718 and AqYield predict yield and soil water content for irrigated crops equally well with limited
719 data. *Agric. For. Meteorol.* 206, 55–68. <https://doi.org/10.1016/j.agrformet.2015.02.011>

720 Cooper, P.J.M., Dimes, J., Rao, K.P.C., Shapiro, B., Shiferaw, B., Twomlow, S., 2008. Coping better with
721 current climatic variability in the rain-fed farming systems of sub-Saharan Africa: An essential
722 first step in adapting to future climate change? *Agric. Ecosyst. Environ.* 126, 24–35.
723 <https://doi.org/10.1016/j.agee.2008.01.007>

724 CORAF/WECARD, 2008. Plan opérationnel 2008-2012. Déployer des systèmes agricoles innovants en
725 Afrique de l’Ouest et du Centre. Dakar, Sénégal, Coraf/Wecard, P10 60.

726 Corre-Hellou, G., Faure, M., Launay, M., Brisson, N., Crozat, Y., 2009. Adaptation of the STICS
727 intercrop model to simulate crop growth and N accumulation in pea–barley intercrops. *Field
728 Crops Res.* 113, 72–81. <https://doi.org/10.1016/j.fcr.2009.04.007>

729 Diedhiou, A., Bichet, A., Wartenburger, R., Seneviratne, S.I., Rowell, D.P., Sylla, M.B., Diallo, I., Todzo,
730 S., Touré, N.E., Camara, M., Ngatchah, B.N., Kane, N.A., Tall, L., Affholder, F., 2018. Changes
731 in climate extremes over West and Central Africa at 1.5 °C and 2 °C global warming. *Environ.
732 Res. Lett.* 13, 065020. <https://doi.org/10.1088/1748-9326/aac3e5>

733 Falconnier, G.N., Corbeels, M., Boote, K.J., Affholder, F., Adam, M., MacCarthy, D.S., Ruane, A.C.,
734 Nendel, C., Whitbread, A.M., Justes, E., Ahuja, L.R., Akinseye, F.M., Alou, I.N., Amouzou, K.A.,
735 Anapalli, S.S., Baron, C., Basso, B., Baudron, F., Bertuzzi, P., Challinor, A.J., Chen, Y., Deryng,
736 D., Elsayed, M.L., Faye, B., Gaiser, T., Galdos, M., Gayler, S., Gerardeaux, E., Giner, M., Grant,
737 B., Hoogenboom, G., Ibrahim, E.S., Kamali, B., Kersebaum, K.C., Kim, S.-H., van der Laan, M.,
738 Leroux, L., Lizaso, J.I., Maestrini, B., Meier, E.A., Mequanint, F., Ndoli, A., Porter, C.H.,
739 Priesack, E., Ripoche, D., Sida, T.S., Singh, U., Smith, W.N., Srivastava, A., Sinha, S., Tao, F.,
740 Thorburn, P.J., Timlin, D., Traore, B., Twine, T., Webber, H., 2020. Modelling climate change
741 impacts on maize yields under low nitrogen input conditions in sub-Saharan Africa. *Glob.
742 Change Biol.* 26, 5942–5964. <https://doi.org/10.1111/gcb.15261>

743 Falconnier, G.N., Journet, E.-P., Bedoussac, L., Vermue, A., Chlébowski, F., Beaudoin, N., Justes, E.,
744 2019. Calibration and evaluation of the STICS soil-crop model for faba bean to explain
745 variability in yield and N₂ fixation. *Eur. J. Agron.* 104, 63–77.
746 <https://doi.org/10.1016/j.eja.2019.01.001>

747 Gbakatchetche, H., Sanogo, S., Camara, M., Bouet, A., 2010. Effet du paillage par des résidus de pois
748 d’angole (*cajanus cajan* l.) sur le rendement du riz (*oryza sativa*) pluvial en zone forestière de
749 CÔTE D’IVOIRE. *Agron. Afr.* 131–137.

750 Gowing, J.W., Palmer, M., 2008. Sustainable agricultural development in sub-Saharan Africa: the case
751 for a paradigm shift in land husbandry. *Soil Use Manag.* 24, 92–99.
752 <https://doi.org/10.1111/j.1475-2743.2007.00137.x>

753 Jayne, T.S., Mason, N.M., Burke, W.J., Ariga, J., 2018. Review: Taking stock of Africa’s second-
754 generation agricultural input subsidy programs. *Food Policy* 75, 1–14.
755 <https://doi.org/10.1016/j.foodpol.2018.01.003>

756 Knörzer, H., Grözinger, H., Graeff-Hönninger, S., Hartung, K., Piepho, H.-P., Claupein, W., 2011.
757 Integrating a simple shading algorithm into CERES-wheat and CERES-maize with particular
758 regard to a changing microclimate within a relay-intercropping system. *Field Crops Res.* 121,
759 274–285. <https://doi.org/10.1016/j.fcr.2010.12.016>

760 Li, L., Yang, S., Li, X., Zhang, F., Christie, P., 1999. Interspecific complementary and competitive
761 interactions between intercropped maize and faba bean. *Plant Soil* 105–114.

762 Li, L., Zhang, F., Li, X., Christie, P., Sun, J., Yang, S., Tang, C., 2003. Interspecific facilitation of nutrient
763 uptake by intercropped maize and faba bean. *Nutr. Cycl. Agroecosystems* 61–71.

764 Lidon, B., Francis, F., 1983. Valorisation agricole des ressources pluviométriques : synthèse de
765 l’atelier IRAT-CIEH tenu à Ouagadougou (Burkina) du 27/11 au 4/12/1982. Montpellier :
766 GERDAT-IRAT, 180 p.

767 Makowski, D., Naud, C., Jeuffroy, M.-H., Barbottin, A., Monod, H., 2006. Global sensitivity analysis for
768 calculating the contribution of genetic parameters to the variance of crop model prediction.
769 *Reliab. Eng. Syst. Saf.* 91, 1142–1147. <https://doi.org/10.1016/j.res.2005.11.015>
770 Masvaya, E.N., 2019. Risk and crop production intensification options for semi-arid southern
771 Zimbabwe (Thesis). Wageningen University. <https://doi.org/10.18174/471607>
772 Pouya, M.B., Bonzi, M., Gnankambary, Z., Traoré, K., Ouédraogo, J.S., Somé, A.N., Sédogo, M.P.,
773 2013. Current soil fertility management practices and their effects on the cotton production
774 and soil on the cotton farms of Central and Western Burkina Faso. *Cah. Agric.* 22, 282–292.
775 <https://doi.org/10.1684/agr.2013.0643>
776 Rezig, M., Sahli, A., Jeddi, F.B., Harbaoui, Y., 2010. Adopting intercropping system for potatoes as
777 practice on drought mitigation under Tunisian conditions. *Options Méditerranéennes* 329–
778 334.
779 Ripoché, A., Crétenet, M., Corbeels, M., Affholder, F., Naudin, K., Sissoko, F., Douzet, J.-M., Tittonell,
780 P., 2015. Cotton as an entry point for soil fertility maintenance and food crop productivity in
781 savannah agroecosystems—Evidence from a long-term experiment in southern Mali. *Field*
782 *Crops Res.* 177, 37–48. <https://doi.org/10.1016/j.fcr.2015.02.013>
783 Sinclair, T.R., Horie, T., 1989. Leaf Nitrogen, Photosynthesis, and Crop Radiation Use Efficiency: A
784 Review. *Crop Sci.* 29, 90–98. <https://doi.org/10.2135/cropsci1989.0011183X002900010023x>
785 Sprent, J.I., Stephens, J.H., Rupela, O.P., 1988. Environmental effects on nitrogen fixation, in:
786 Summerfield, R.J. (Ed.), *World Crops: Cool Season Food Legumes*, Current Plant Science and
787 Biotechnology in Agriculture. Springer Netherlands, Dordrecht, pp. 801–810.
788 https://doi.org/10.1007/978-94-009-2764-3_64
789 Stern, R.D., Cooper, P.J.M., 2011. ASSESSING CLIMATE RISK AND CLIMATE CHANGE USING RAINFALL
790 DATA – A CASE STUDY FROM ZAMBIA. *Exp. Agric.* 47, 241–266.
791 <https://doi.org/10.1017/S0014479711000081>
792 Stern, R.D., Dennett, M.D., Garbutt, D.J., 1981. The start of the rains in West Africa. *J. Climatol.* 1, 59–
793 68. <https://doi.org/10.1002/joc.3370010107>
794 Sultan, B., Gaetani, M., 2016. Agriculture in West Africa in the Twenty-First Century: Climate Change
795 and Impacts Scenarios, and Potential for Adaptation. *Front. Plant Sci.* 7, 20.
796 <https://doi.org/10.3389/fpls.2016.01262>
797 Sultan, B., Guan, K., Kouressy, M., Biasutti, M., Piani, C., Hammer, G.L., McLean, G., Lobell, D.B., 2014.
798 Robust features of future climate change impacts on sorghum yields in West Africa. *Environ.*
799 *Res. Lett.* 9, 104006. <https://doi.org/10.1088/1748-9326/9/10/104006>
800 Tittonell, P., Giller, K.E., 2013. When yield gaps are poverty traps: The paradigm of ecological
801 intensification in African smallholder agriculture. *Field Crops Res.* 143, 76–90.
802 <https://doi.org/10.1016/j.fcr.2012.10.007>
803 Tovignan, T.K., Fonceka, D., Ndoye, I., Cisse, N., Luquet, D., 2016. The sowing date and post-flowering
804 water status affect the sugar and grain production of photoperiodic, sweet sorghum through
805 the regulation of sink size and leaf area dynamics. *Field Crops Res.* 192, 67–77.
806 <https://doi.org/10.1016/j.fcr.2016.04.015>
807 Traore, A., 2015. Analyse de l'évolution à long terme de l'écart de rendement du sorgho dans une
808 rotation coton-sorgho-arachide plus ou moins intensifiée Approche par modélisation
809 (Mémoire de Master). SUPAGRO Montpellier, Montpellier France.
810 Traore, B., Corbeels, M., van Wijk, M.T., Rufino, M.C., Giller, K.E., 2013. Effects of climate variability
811 and climate change on crop production in southern Mali. *Eur. J. Agron.* 49, 115–125.
812 <https://doi.org/10.1016/j.eja.2013.04.004>
813 Tsubo, M., Mukhala, E., Ogindo, H., Walker, S., 2004. Productivity of maize-bean intercropping in a
814 semi-arid region of South Africa. *Water SA* 29, 381–388.
815 <https://doi.org/10.4314/wsa.v29i4.5038>
816 Vandermeer, J., 1989. *The ecology of intercropping*, Cambridge University Press. ed. Cambridge.

817 Vanlauwe, B., Coyne, D., Gockowski, J., Hauser, S., Huising, J., Masso, C., Nziguheba, G., Schut, M.,
818 Van Asten, P., 2014. Sustainable intensification and the African smallholder farmer. *Curr.*
819 *Opin. Environ. Sustain.* 8, 15–22. <https://doi.org/10.1016/j.cosust.2014.06.001>
820 Weih, M., Karley, A.J., Newton, A.C., Kiær, L.P., Scherber, C., Rubiales, D., Adam, E., Ajal, J.,
821 Brandmeier, J., Pappagallo, S., Villegas-Fernández, A., Reckling, M., Tavoletti, S., 2021. Grain
822 Yield Stability of Cereal-Legume Intercrops Is Greater Than Sole Crops in More Productive
823 Conditions. *Agriculture* 11, 255. <https://doi.org/10.3390/agriculture11030255>
824 Willey, R.W., 1979. Intercropping-its importance and research needs. Part 1. Competition and yield
825 advantages. *Field Crops Abstr* 32, 1–10.
826 Willey, R.W., Rao, M.R., 1980. A Competitive Ratio for Quantifying Competition Between Intercrops.
827 *Exp. Agric.* 16, 117–125. <https://doi.org/10.1017/S0014479700010802>
828 Yang, C., Huang, G., Chai, Q., Luo, Z., 2011. Water use and yield of wheat/maize intercropping under
829 alternate irrigation in the oasis field of northwest China. *Field Crops Res.* 124, 426–432.
830 <https://doi.org/10.1016/j.fcr.2011.07.013>
831 Yu, Y., Stomph, T.-J., Makowski, D., van der Werf, W., 2015. Temporal niche differentiation increases
832 the land equivalent ratio of annual intercrops: A meta-analysis. *Field Crops Res.* 184, 133–
833 144. <https://doi.org/10.1016/j.fcr.2015.09.010>
834

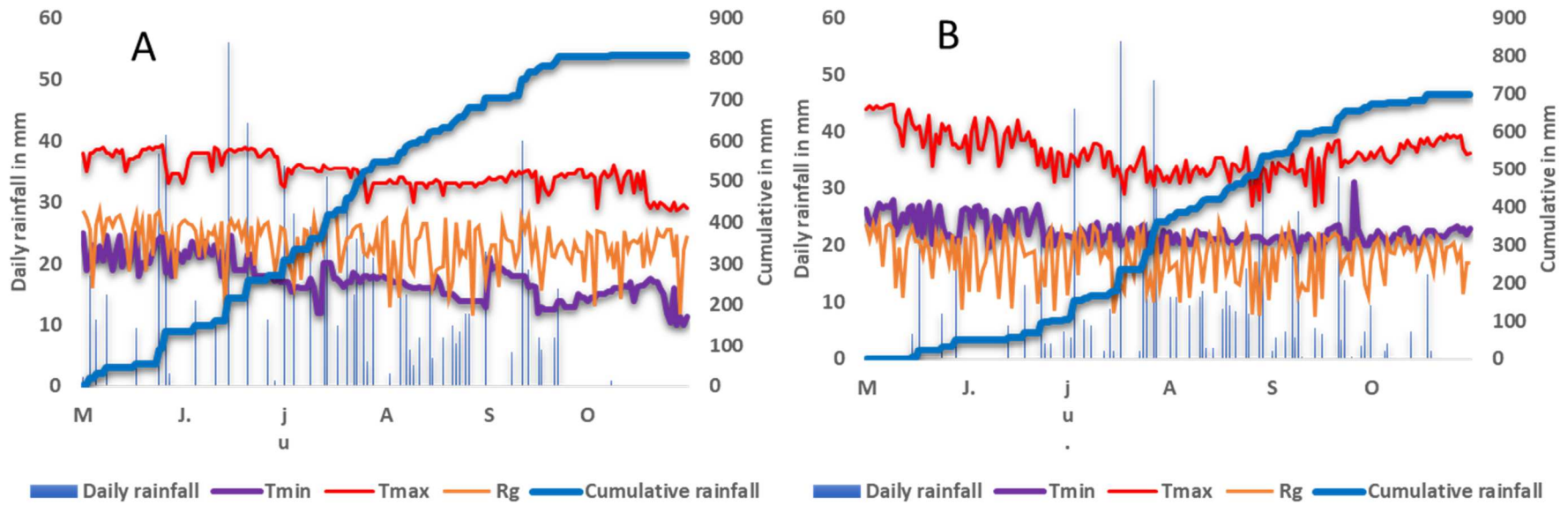


Figure 1: Daily and cumulative rainfall, daily temperature and daily global radiation (Rg) for 2017 (A) and 2018 (B) at the N'Tarla agricultural research station, southern Mali

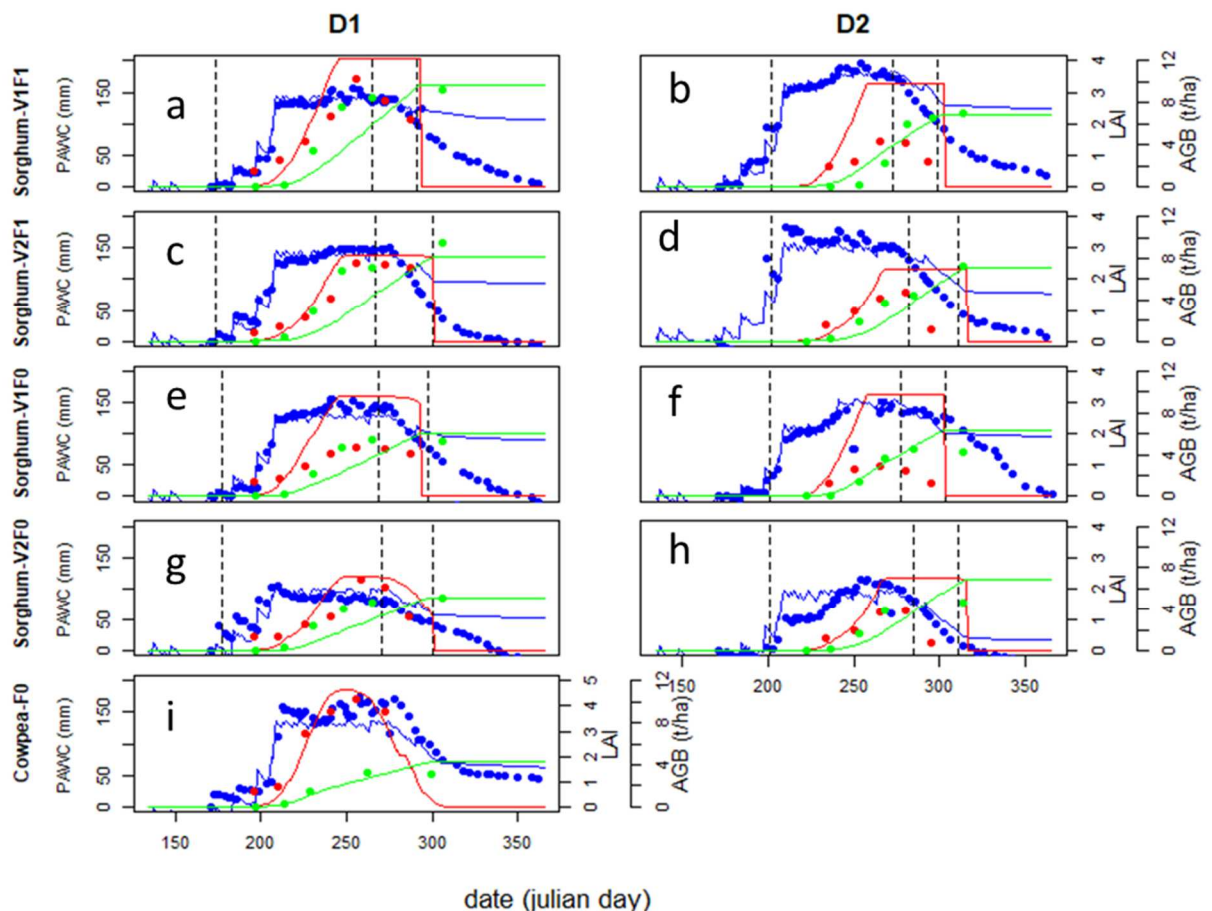


Fig. 2: Observed plant available soil water to maximum measurement depth (PAWC) (blue dots), LAI (red dots), and aboveground biomass (green dots) for the eight sole sorghum simulation units and one sole cowpea simulation unit used for model calibration. The lines are model simulations: blue for soil, red for LAI and green for AGB. Local variety (V1), improved variety (V2), early sowing (D1), late sowing (D2), no fertilizer (F0), 38 kg N ha⁻¹ (F1) at Ntarla research station in 2018. Vertical dotted bars indicate from left to right sowing date, beginning of grain filling and physiological maturity.

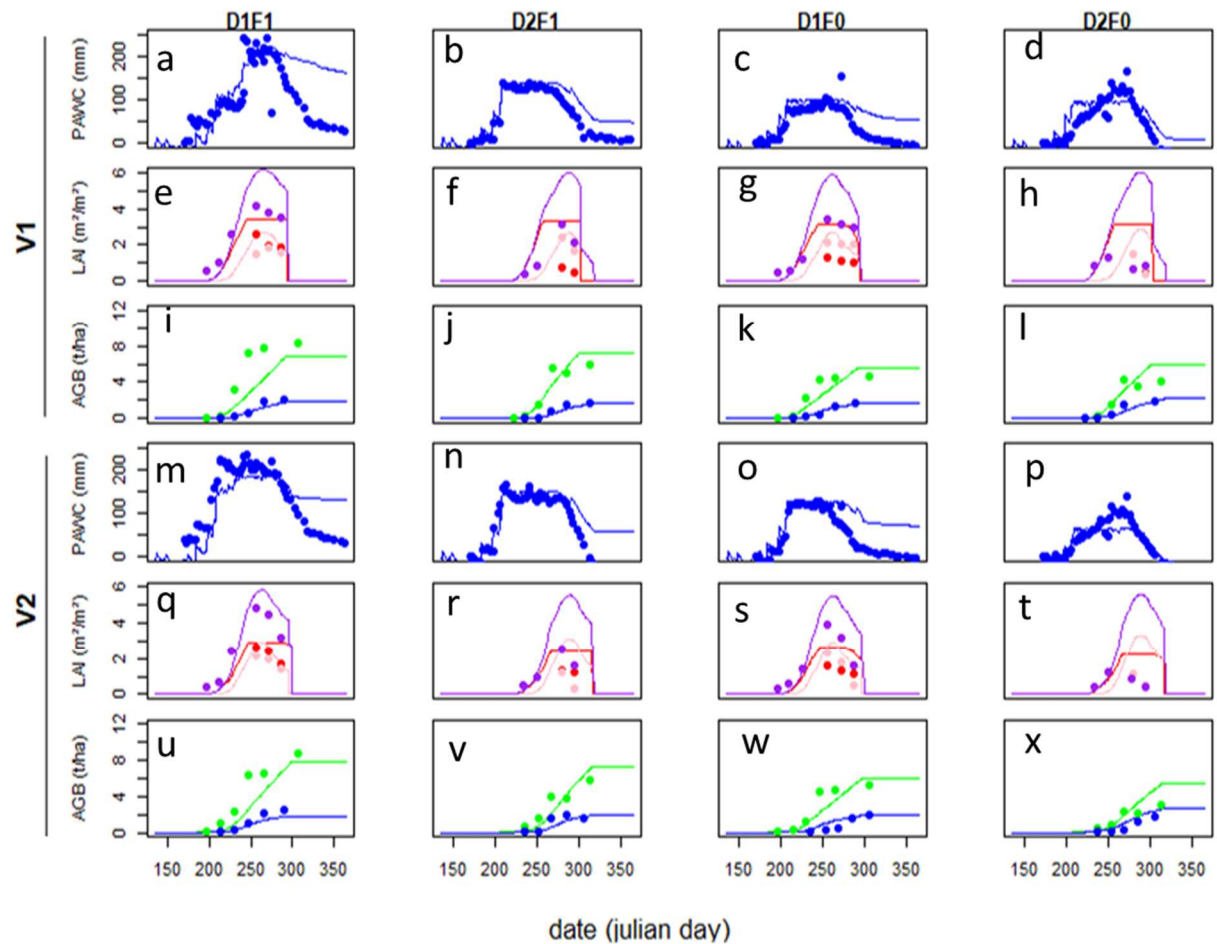


Fig. 3: Observation (blue dots) and simulation (blue line) of plant available soil water to maximum measurement depth (PAWC) (Fig. a, b, c, d, m, n, o, p); observation (purple dots for sorghum + cowpea, red dots for sorghum in intercropping, pink and transparent dots for cowpea in intercropping) and simulations (lines) of LAI (Fig. e, f, g, h, q, r, s, t); observation (green dots for sorghum, blue dots for cowpea) and simulations (green lines for sorghum and blue lines for cowpea) of aboveground biomass of sorghum intercropped with cowpea (Fig. i, j, k, l, u, v, w, x). Local variety (V1), improved variety (V2), early sowing (D1), late sowing (D2), 38 kgN ha⁻¹ (F1), 0 kgN ha⁻¹ (F0) at Ntarla research station in 2018.

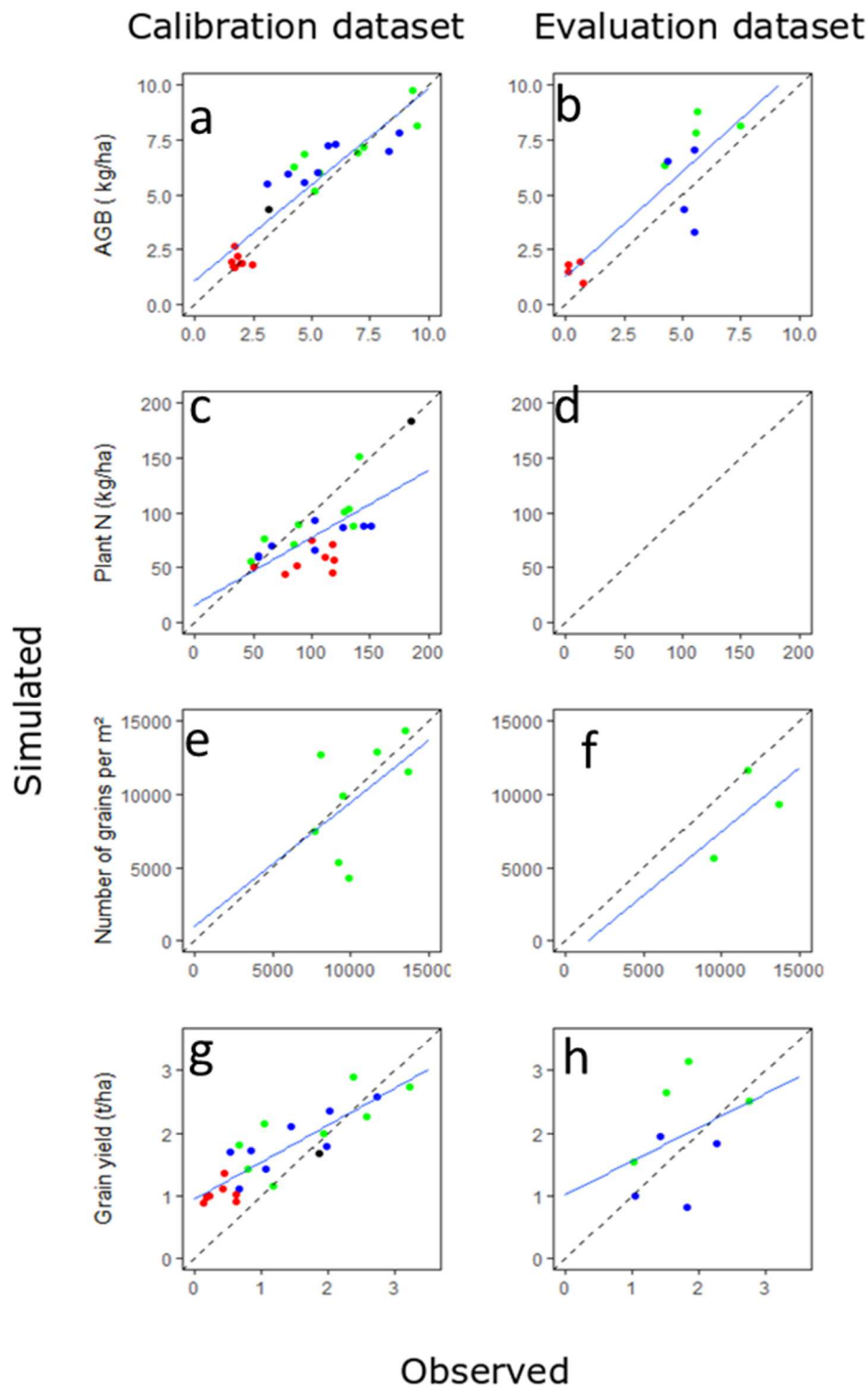


Fig. 4: Observed and STICS simulated aboveground biomass at harvest (AGB), aboveground plant N at harvest (Plant N), number of grains per m² of sorghum at harvest and grain yield for calibration and evaluation datasets (see Table1). There were no observation for cowpea grain yield in 2017 and sorghum and cowpea plant N in the evaluation dataset. The dotted black line is the 1:1 line. The blue line is the regression of simulated values against observed values. Red dots are intercropped cowpea, black dots are sole cowpea, green dots are sole sorghum, blue dots are intercropped sorghum.

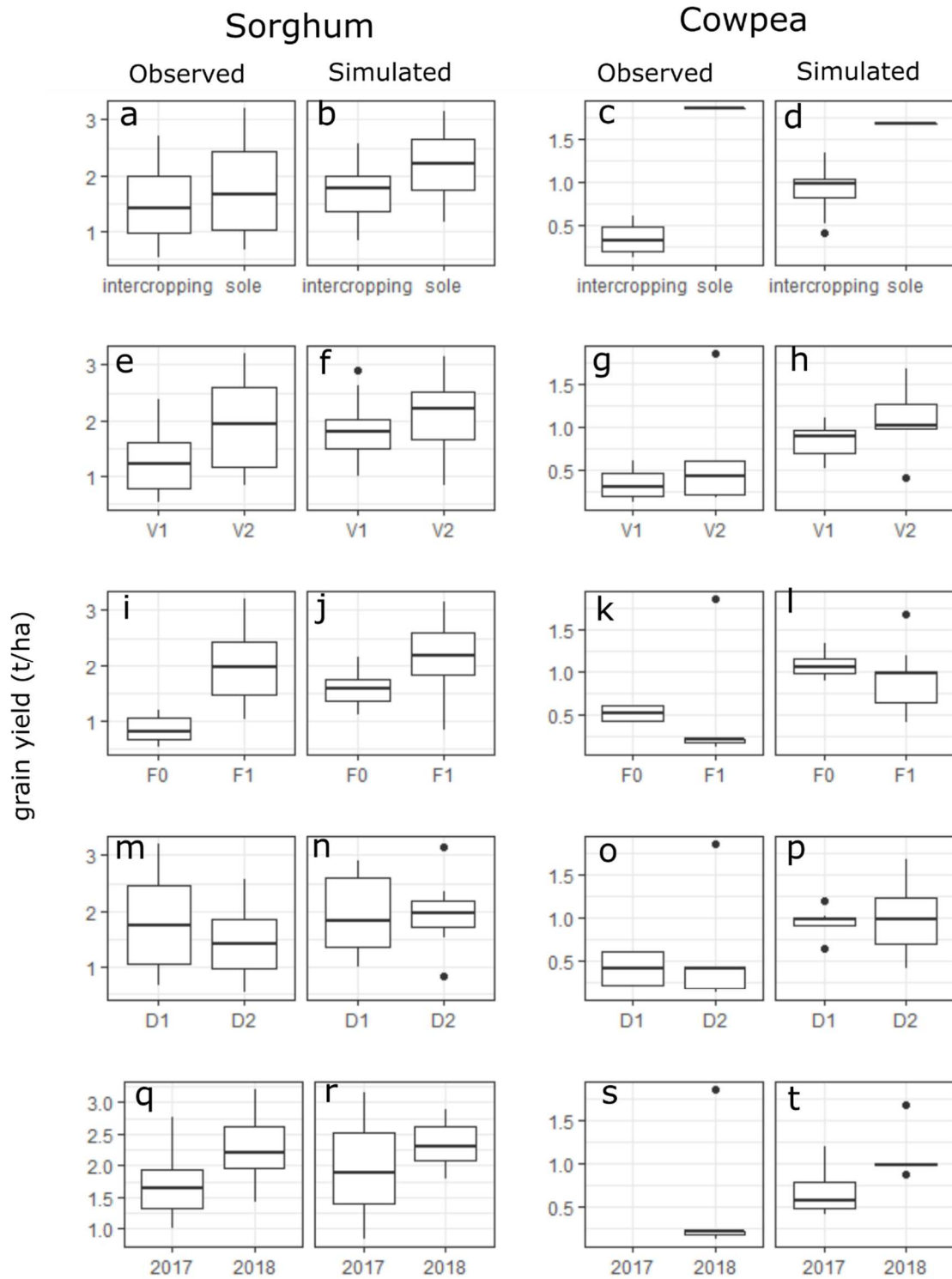


Figure 5: Boxplots of the effect of intercropping, sorghum variety (V1: local, V2: improved), N input (F0 = 0 kgN/ha⁻¹, F1 = 8 kg N/ha⁻¹ in 2017 and 38 kgN/ha⁻¹ in 2018) and sowing date (D1: early, D2: late) on observed and simulated grain yield. Calibration and evaluation datasets were pooled together. In 2017, there was no sole cowpea plot and no data for cowpea grain yield.

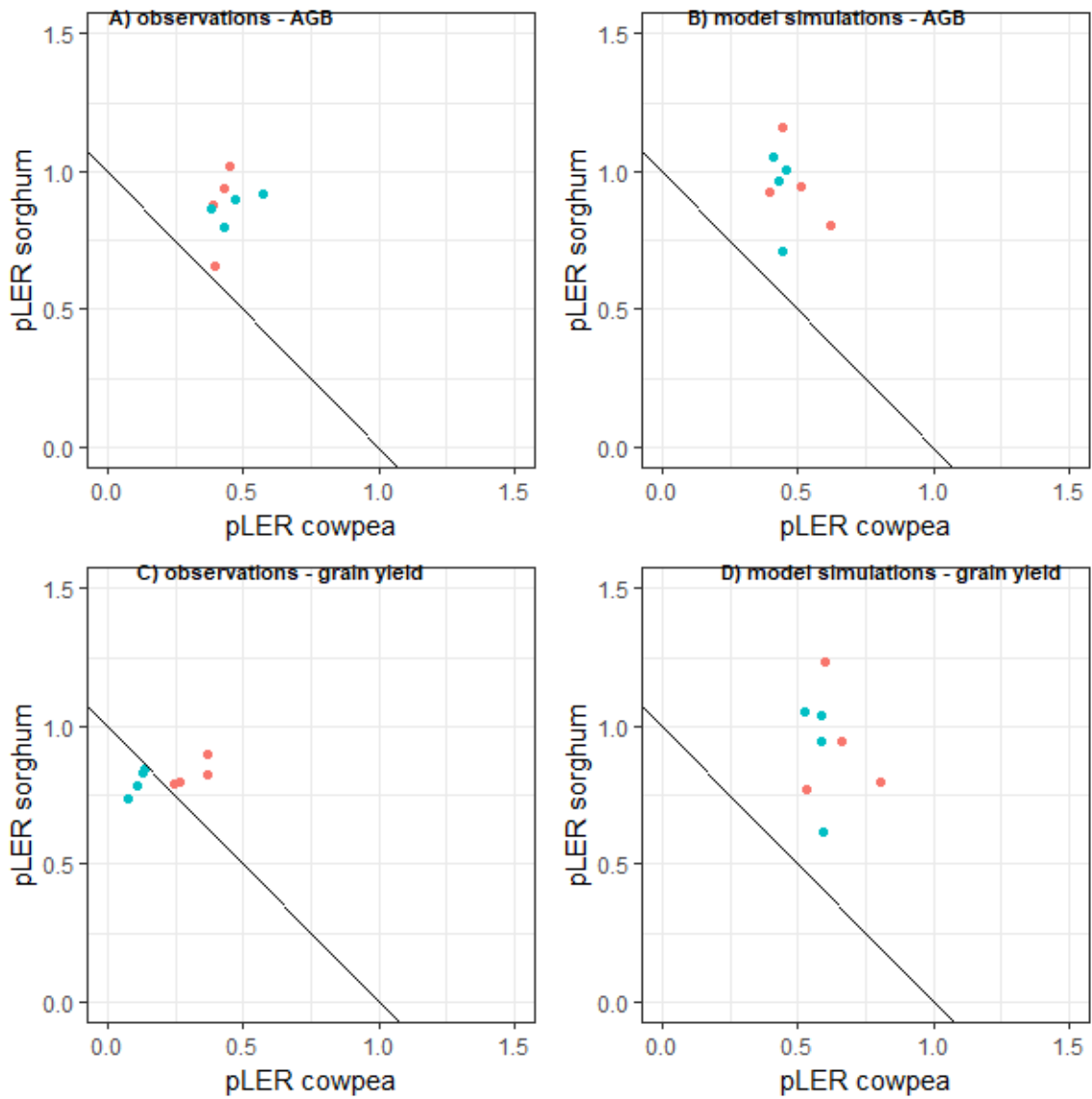


Figure 6: Cowpea and sorghum pLER as observed in the experiment (A for aboveground biomass and C for grain yield) and as simulated by the model (B for aboveground biomass and D for grain yield). Blue dots are F1 (38 kgN ha⁻¹) treatments, and red dots are F0 (0 kgN ha⁻¹) treatments. The black line indicates a LER of 1. Only 2018 experimental plots and simulation units were considered, as in 2017 pLER of cowpea could not be computed because there was no experimental plot with sole cowpea.

Table 1: Description of the experimental plots at the Agronomic Research Station of N'Tarla used for model calibration and evaluation. V1 = Photoperiod-sensitive local sorghum variety, V2 = Improved sorghum variety less sensitive to the photoperiod, D1 = early sowing, D2 = late sowing, F1 = 38 kg N/ha in 2018 and f=8 kg N/ha in 2017, F0 = 0 kg N/ha. Soil water was not measured in 2017.

Dataset	Year	Cropping system	Variety	Sowing date	Fertilizer application	Maximum depth of soil water measurement (cm)	Soil organic nitrogen content in the upper layer in %	Name of simulation unit	Field capacity (%).						
									0 - 15 cm	15 - 35 cm	35- 55 cm	55-85 cm	85- maximum depth		
Calibration	2018	Sole sorghum	V1	D1	F0	115	0.01	V1D1F0	13.7	19	19	19	19		
					F1	145	0.02	V1D1F1	13.7	19	19	19	19		
				D2	F0	155	0.01	V1D2F0	13.5	19	24	24	24		
			F1		155	0.008	V1D2F1	22	25	25	28	19			
			V2	D1	F0	95	0.007	V2D1F0	13.7	19	19	18	18		
					F1	155	0.011	V2D1F1	13.7	19	19	19	19		
		D2		F0	155	0.016	V2D2F0	13.7	19	19	19	22			
			F1	155	0.018	V2D2F1	13.7	19	21	21	21				
		Sorghum intercropped with cowpea	V1	D1	F0	145	0.015	V1D1F0-inter	13.7	17	17	17	19		
					F1	155	0.015	V1D1F1-inter	13.7	23	23	23	23		
				D2	F0	155	0.011	V1D2F0-inter	13.7	19	18	18	18		
			F1		135	0.011	V1D2F1-inter	13.7	19	19	19	20			
			V2	D1	F0	145	0.011	V2D1F0-inter	13.7	17	17	17	17		
					F1	155	0.013	V2D1F1-inter	13.7	22	22	22	22		
		D2		F0	135	0.012	V2D2F0-inter	13.7	20	20	20	20			
			F1	145	0.012	V2D2F1-inter	13.7	22	22	22	22				
		Evaluation	2017	Sole Cowpea	Sangaraka	D1	F0	135	0.01	Niebpur	13.7	19	21	21	21**
				Sole sorghum	V1	D1	f	-	0.02*	NT17V1D1	13.7	19	19	19	19**
f	-						0.007*	NT17V1D2	22	25	25	28	19**		
V2	D1			f	-	0.011*	NT17V2D1	13.7	19	19	19	19**			
				f	-	0.018*	NT17V2D2	13.7	19	21	21	21**			
Sorghum intercropped with cowpea	V1			D1	f	-	0.015*	NT17V1D1-inter	13.7	23	23	23	23**		
					f	-	0.011*	NT17SV1D2-inter	13.7	19	19	19	20**		
	V2			D1	f	-	0.013*	NT17V2D1-inter	13.7	19	19	19	20**		
					f	-	0.012*	NT17V2D2-inter	13.7	22	22	22	22**		

* Not measured in 2017, the values measured in 2018 in the experimental plot were considered. ** maximum depth of soil water measurement for 2018 was considered

Table 2: Description of the crop development stages included in the model

Development stage	Step acronym used in model	Description
Vegetative stages	ILEV	Emergence
	AMF	Maximum acceleration of leaf growth, end of juvenile phase
	ILAX	Maximum leaf area index (LAI), end of leaf growth
Reproductive stages	Flo	Flowering
	IDRP	Start of grain filling
	IMAT	Physiological maturity

Table 3: Values of sorghum parameter as calibrated in the STICS crop model for experiments at N'Tarla in Mali

Parameter			Value				
Process	acronym	Description	Target variable	V1	V2	Cowpea	Source
Emergence	tdmin	basal temperature for crop development		8	8	6.2	(Folliard et al. 2004) ; Fao., 2012 ; (Luo 2011).
Crop development	sensiphot	index of photoperiod sensitivity (1=insensitive)		0.4	0.6	-	Trial and error calibration
	phobase	basal photoperiod controlling photoperiod slowing effect		14	14	-	Traoré A., 2015
	phosat	saturating photoperiod controlling photoperiod slowing effect		12,75	12.75	-	Traoré, A., 2015
	stlevamf	cumulative thermal time between emergence and end of juvenile phase	Leaf area index	180	718	881	Test of a range of values
	stamflax	cumulative thermal time between end of juvenile phase and maximum LAI		305	314	687	Trial and error calibration
	stlevdrp	cumulative thermal time between emergence and beginning of grain filling		685	1077	1609	Trial and error calibration
Leaves	dlaimaxbrut	maximum rate of the setting up of LAI		0,0015200	0.01	0.0035	Trial and error calibration
	durvieF	maximal lifespan of an adult leaf		480	280	240	Trial and error calibration
Shoot growth	efcroijuv	maximum radiation use efficiency during the juvenile phase	Aboveground biomass	2.1836	2.1877	1.2	Trial and error calibration
	efcroirepro	maximum radiation use efficiency during the grain filling phase	Aboveground biomass	3.8572	2.8372	1.3559	Trial and error calibration
	efcroiveg	maximum radiation use efficiency during the vegetative stage	Aboveground biomass	3.8049	2.8414	1.7465	Trial and error calibration
	temin	basal temperature for photosynthesis	-	11	11	7.2	(Folliard et al. 2004) ; Fao., 2012 ; (Luo 2011)
	teopt	optimal temperature for photosynthesis	-	25	25	27	(Folliard et al. 2004) ; Fao., 2012 ; (Luo 2011)
	temax	maximal temperature for photosynthesis	-	45	45	40	(Folliard et al. 2004) ; Fao., 2012 ; (Luo 2011)
Nitrogen fixation	fixmaxgr	maximal N symbiotic fixation rate per unit of grain growth rate	N ₂ fixed*	-	-	9.5	Trial and error calibration
	fixmaxveg	maximal N symbiotic fixation rate per unit of vegetative growth rate	N ₂ fixed*	-	-	30	Trial and error calibration
Nitrogen uptake	Kmabs2	affinity constant of N uptake by roots for the low uptake system	N uptake	40000	37672.32	25000	numerical optimization
	Vmax2	maximum specific N uptake rate with the high affinity transport system	N Uptake	0.1	0.00878	0.06	numerical optimization
Yield formation	cgrain	slope of the relationship between grain number and growth rate	Number of grains	0.06	0.07	0.084	Trial and error calibration
	nbggrain	Duration in days of the period during which the number of grains can be reduced by stresses	Number of grains	15	15	15	Traoré, A., 2015
	cgrainv0	number of grains produced during the nbgrains before beginning of grain filling	Number of grains	0	0	0.069	Trial and error calibration
Yield formation	vitircarbe	rate of increase of the C harvest index vs time	grain yield	0.009	0.015	0.01462	Trial and error calibration
	nbgrmax	maximum number of grains	grain yield	25000	60000	1200	Measurement
	pgrainmaxi	maximum weight of one grain	grain yield	0.027	0.0247	0.25	Measurement
	lmax	maximum harvest index	grain yield	0.3	0.51	0.42	Measurement

*N₂ fixed by the legume was not measured, but estimated for sole cowpea as the difference between the N uptake of sole sorghum without fertilizer and N uptake of sole cowpea.

Table 4: Root mean square error (RMSE), relative root means square error (rRMSE), and efficiency (EF) of STICS simulation of plant variables in the N'Tarla experiment in 2017 and 2018. See Table 1 for details on the calibration and evaluation datasets.

Crops	Variable	EF (-)		nRMSE (%)	
		Calibration	Evaluation	Calibration	Evaluation
Sorghum and cowpea	AGB	0.81	0.58	23	43
	Plant N	-0.05		35	
	Number of grains per m ²	-0.28	-2.17	29	25
	Grain yield	0.56	-0.56	49	41
Sorghum only	AGB	0.54	-0.66	21	85
	Plant N	0.26	-	30	-
	Number of grains per m ²	-0.28	-2.17	29	25
	Grain yield	0.38	-0.56	41	41
Cowpea only	AGB	-0.09	-0.33	25	99
	Plant N	-0.64	-	42	-
	Number of grains per m ²	-	-	-	-
	Grain yield	0.20	-4.66	87	66