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PHOTOSYNTHETIC ACTIVITY AND EARLY GROWTH OF FOUR CACAO GENOTYPES AS INFLUENCED BY DIFFERENT SHADE REGIMES UNDER WEST AFRICAN DRY AND WET SEASON CONDITIONS

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SUMMARY

The physiological performance of four cacao clones was examined under three artificial shade regimes over the course of a year in Ghana. Plants under light shade had significantly higher photosynthetic rates in the rainy seasons whereas in the dry season there was a trend of higher photosynthetic rates under heavy shade. The results imply that during the wet seasons light was the main limiting factor to photosynthesis whereas in the dry season vapour pressure deficit was the major factor limiting photosynthesis through stomatal regulation. Leaf area was generally lower under heavier shade but the difference between shade treatments varied between clones. Such differences in leaf area allocation appeared to underlie genotypic differences in final biomass production in response to shade. The results suggest that shade for young cacao should be provided based on the current ambient environment and genotype.

INTRODUCTION

The use of shade trees is a common practice in the establishment of cacao (*Theobroma cacao* L.) (de Almeida and Valle, 2007; Raja Harun and Hardwick, 1988b). Gaining a better understanding of how light levels impact the early performance of cacao is an important component in optimising its growth during the establishment phase. Such knowledge is important because replanting of cacao is expected to become more challenging under predicted climate change scenarios, and yet this will be needed if the anticipated future demand for cocoa beans (International Cocoa Organization (ICCO), 2008) is to be met.

Cacao has a number of attributes that are typical of shade species. These include low photosynthetic rates (Daymond *et al.*, 2011) and a high sensitivity to photoinhibitory stress (Galyuon *et al.*, 1996a, b; Serrano and Biehl, 1996). There is a general view that shading (usually by other plants) is a requirement of young cacao plants to ameliorate physiological stress caused by high light intensities and to reduce evapotranspiration (Ahenkorah *et al.*, 1974). However, there is still considerable inconsistency in what is considered to be the optimal shading levels for cacao. For example, whereas cacao is also known to have low light saturation levels of around $400 \mu\text{mol m}^{-2} \text{s}^{-1}$ (Daymond

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et al., 2011; Raja Harun and Hardwick, 1988a), studies conducted by Da Matta *et al.* (2001) and Rada *et al.* (2005) suggest that some cacao genotypes may show increasing photosynthetic rates as the photosynthetically active radiation (PAR) increases up to $750 \mu\text{mol m}^{-2} \text{s}^{-1}$. Furthermore, differences between genotypes in canopy attributes (e.g. size of canopy and amount of self-shading) will impact on their response to shade.

Since there is a defined dry season in West Africa where the bulk of the world's cacao is cultivated, it is also important to understand whether genotypic responses to different shade regimes vary across seasons. Moreover, there is a paucity of information on the extent of genotypic response to different shading regimes, particularly across seasons.

To address these questions, the study described here assesses the response to shade of four contrasting cacao clones (SCA 6, T 79/501, P 30 [POS] and PA 150) all grafted on a common rootstock over a 12-month period. To circumvent possible complications resulting from inter-species competition (as is the case typically under field conditions where other plants are used as shade for cacao) the experiment was conducted in a nursery environment using shade netting to vary the intensity of incident light.

MATERIALS AND METHODS

Plant material

Three cacao clones T 79/501, PA 150 and SCA 6, representing, respectively, a vigorous, an intermediate and a low vigour genotype, and P 30 [POS] (a traditionally cultivated cacao genotype in Ghana, used in this work as control) were propagated by side grafting using a common seedling rootstock (T 60/887 \times Amelonado). Care was taken to insert the scion below the point of attachment of cotyledons of the rootstock to prevent chupons growing from below the graft union.

Experimental design

The experiment was conducted at the Cocoa Research Institute of Ghana, New Tafo, Eastern Region, Ghana. The experiment was designed as a split-plot with shade as the main plot factor and the four genotypes as sub-plots. There were five plants per sub-plot and each plot was replicated three times. Results were analysed (using Genstat) by analyses of variance, and significance among mean values was determined by least significant difference (LSD) values where $p = 0.05$.

The experiment was conducted inside a $35 \text{ m} \times 30\text{-m}$ nursery shade house that was covered artificially on all sides with a single-layer of shade netting, supported by galvanized pipes and rafters. Shade levels were measured twice in each of the three seasons by means of a 'SunScan' System SS1 (Delta-T, UK). The experiment incorporated three shade regimes: 'light' (32.5%), 'medium' (55%) and 'heavy' (76%) shades, created, respectively, by means of (i) the single layer of shade netting, (ii) the erection of an additional single-netted or (iii) double-netted cage over the growing areas.

Prior to imposition of the shade treatments, a drainage system was put into place. The plants were put into the treatments on 8 March 2007. Growing pots [black polythene bags ($32 \text{ cm} \times 37.5 \text{ cm}$), perforated with 28 evenly distributed holes (0.7 cm in diameter)] and filled with top-soil] were placed in pits on top of a layer of river sand.

Each pot contained a young clonal cacao plant of one of the four cacao genotypes. More sand was added between the pots until the river sand layer was approximately 27-cm deep, leaving the top 15 cm of the pit empty. The perforations in the growing pots provided contact between the topsoil in the pots and the river sand in the pits, while the holes in the polythene sheet ensured drainage down the pit. Watering was not direct to the growing pots, but was applied to the river sand when its water content declined from field capacity.

Photosynthesis and leaf chlorophyll fluorescence measurements

Beginning in April 2007, a portable photosynthesis-measuring system incorporating an infrared gas analyzer (LCpro+, ADC Bioscience, UK) was used to measure photosynthesis once every month through the major rainy (April–July) and the minor rainy (August–October) seasons of 2007 and during the dry season (in November 2007 and January/February 2008) of the 2007–2008 crop year of Ghana. Measurements were carried out between 9:00 am and 11:00 am on three plants per treatment per block. A diurnal study was also carried out on 28 April 2008, in which an artificial light source provided a constant saturated light intensity ($1200 \mu\text{mol photons m}^{-2} \text{s}^{-1}$) to assess diurnal photosynthesis under constant light conditions. Measurements were carried out between 8:00 am and 4:00 pm.

Around midday on the same days that photosynthesis measurements were conducted, leaf chlorophyll fluorescence was measured using a leaf chlorophyll fluorescence meter (FP100; Photon Systems Instruments, the Czech Republic) on leaves that had been dark adapted for 30 min. A set of diurnal leaf chlorophyll fluorescence measurements was conducted on 28 April 2008 between 8 am and 4 pm.

Microclimate measurements

Small data loggers (Tinytag, Gemini Data Loggers, Chichester, UK) placed in Stephenson screens were used to record the temperature and relative humidity at 15-min intervals in the three shade environments of the trial.

Determination of leaf area and plant dry weight

At the end of the experiment (in May 2008) two out of the three plants in each plot (= 6 per treatment) that had been previously selected for photosynthesis measurements were harvested destructively to determine the leaf area (using an automatic leaf area meter, Delta-T Devices Ltd., Burwell, Cambridge, UK) and subsequently plant organ dry weight after drying at 80 °C for 60 h (using a ventilated drying oven, OV-165, Thermo Scientific, UK).

RESULTS

Microclimatic data

Table 1 summarises the temperature and relative humidity conditions recorded under the different shade regimes during the three seasons, and Table 2 summarises the mean sunshine hours across the seasons.

Table 1. Mean seasonal relative humidity (RH), temperature (T) and vapour pressure deficit (VPD) under different shade regimes over the experimental period.

Season	Major rainy season			Minor rainy season			Dry			Mean		
Shade level	RH (%)	T (°C)	VPD (kPa)	RH (%)	T (°C)	VPD (kPa)	RH (%)	T (°C)	VPD (kPa)	RH (%)	T (°C)	VPD (kPa)
32.5%	70.0	36.9	1.9	64.9	31.9	1.7	50.2	36.8	3.1	61.7	35.2	2.2
55%	72.8	34.8	1.5	73.1	31.2	1.2	51.1	35.2	2.8	65.7	33.7	1.8
76%	75.0	32.2	1.2	76.8	31.0	1.0	58.5	33.3	2.1	70.1	32.2	1.4
Mean	72.6	34.6	1.5	71.6	31.4	1.3	53.3	35.1	2.7	65.8	32.7	1.8

Table 2. Mean seasonal sunshine hours over the experimental period.

	Major rainy season	Minor rainy season	Dry season	Mean
Mean sunshine hours	5.2	4.4	6.7	5.4

As a result of overcast conditions that characterised much of the minor rainy season (Table 2), mean aerial temperatures were approximately 3.3 °C lower during the minor rainy season compared with the major rainy season and 3.8 °C lower than during the dry season. Also, mean temperature under the heavy shade was approximately 3 °C lower than under the light shade treatment (Table 1).

The mean relative humidity averaged over two rainy seasons was 72.1% compared with a mean value of 53.3% for the dry season. The mean relative humidity under heavy shade was 70.1% compared with 61.7% under light shade (Table 1). Vapour pressure deficit was lower during the rainy seasons than during the dry season, the difference being the greatest under the light shade regime.

Seasonal variation in stomatal conductance

There was a significant ($p < 0.001$) season \times shade interaction on stomatal conductance. During the dry season, plants under heavy shade had higher stomatal conductance. In contrast, in the rainy seasons, heavily shaded plants had lower stomatal conductance rates compared with plants under lighter shade. Over the entire experimental period, differences in stomatal conductance between cacao clones were not significant. However, a significant ($p < 0.001$) season \times clone interaction was observed. While all the clones had lower stomatal conductance rates in the dry season than in the rainy seasons, the difference was more pronounced for the clone PA 150 (Figure 1).

Seasonal variation in photosynthesis

Similar to the observation in stomatal conductance, there was a seasonal influence on photosynthetic rate ($p < 0.001$). The response to shade treatments varied between seasons, hence the season \times shade interaction was significant ($p < 0.001$). The lowest rates were recorded in the dry season, in which little variation was observed between shade treatments (Figure 2). In that season, plants under medium shade had the

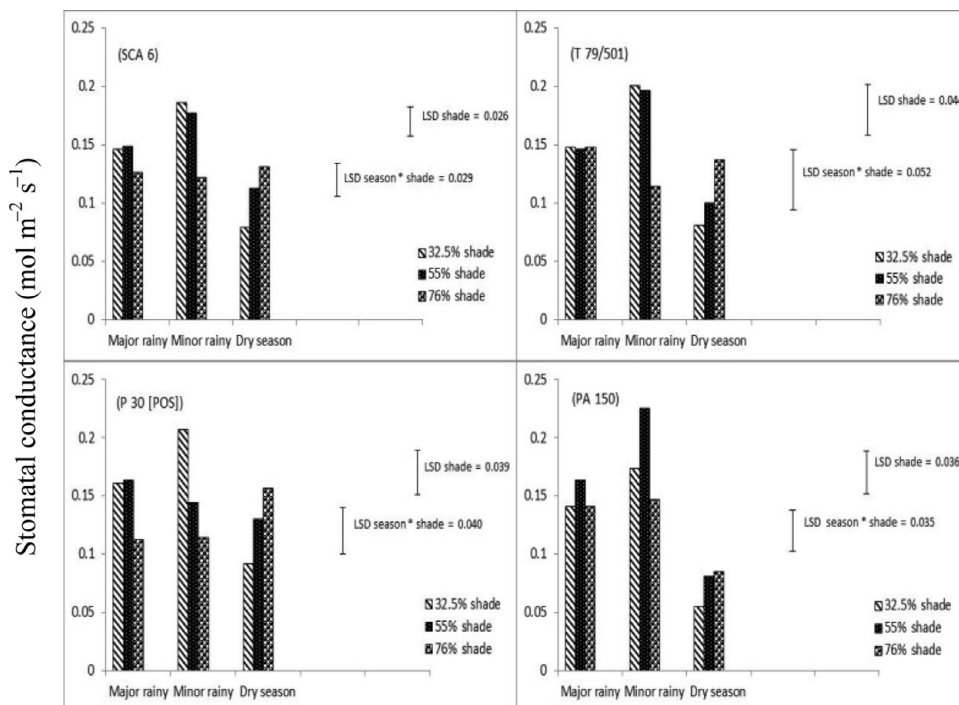


Figure 1. Stomatal conductance of cacao under varying shade and in different seasons. (Each bar represents the mean of four measurements on nine plants per treatment).

highest photosynthetic rates (mean = $0.82 \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), while those under light shade had the lowest photosynthetic rates (mean = $0.50 \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$). Photosynthetic rates were higher in the rainy seasons compared with the dry season. In the rainy seasons photosynthesis was the highest under light shade and the lowest under heavy shade. The average photosynthetic rates under light, medium and heavy shade treatments were 3.3 , 1.6 and $1.2 \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$, respectively, in the minor rainy season, and 6.0 , 3.0 and $1.1 \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$, respectively, in the major rainy season. Variation in photosynthesis between shade treatments was consistent across all genotypes with no significant differences between genotypes (Figure 2).

Diurnal pattern of photosynthesis

Photosynthetic rates were significantly higher ($p = 0.004$) during the first half of the day (between 8:00 am and 12:00 noon) than during the second half. The highest rates of photosynthesis were recorded between 10:00 am and 12:00 noon, after which the rates declined steadily until the end of the day (Figure 3). Photosynthetic rates declined with increasing shade intensity ($p < 0.001$). There was no interaction between time of day and shade level on diurnal pattern of photosynthesis. There was also no significant difference in photosynthetic rates between the cacao genotypes throughout the course of the day.

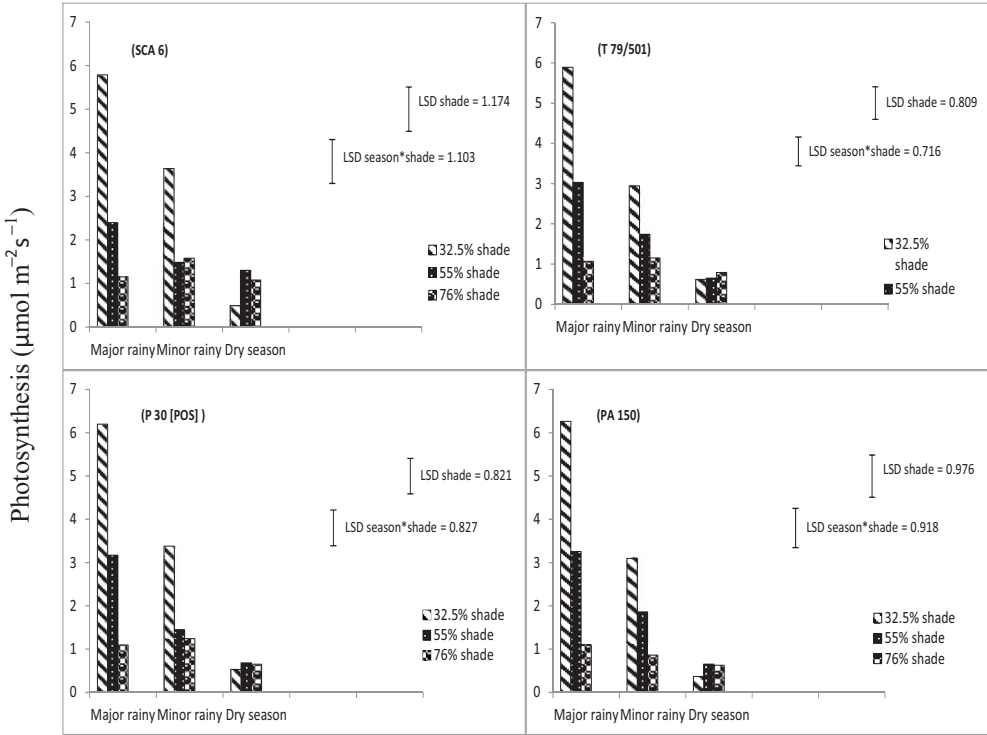


Figure 2. Photosynthesis of cacao under varying shade and in different seasons. (Each bar represents the mean of four measurements on nine plants per treatment).

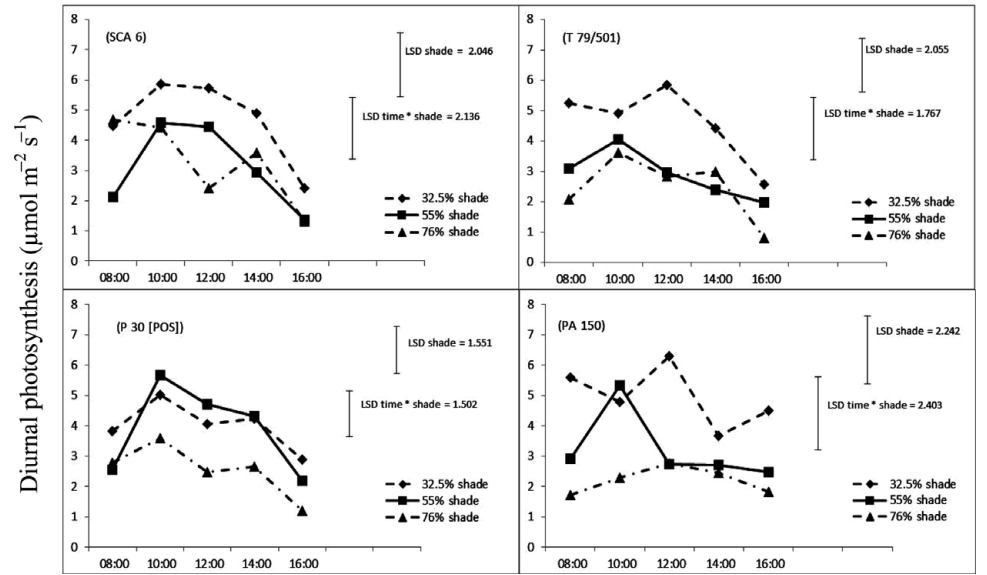


Figure 3. Diurnal pattern of photosynthesis of cacao under different shade levels. (Each line represents the mean for nine plants per shade treatment).

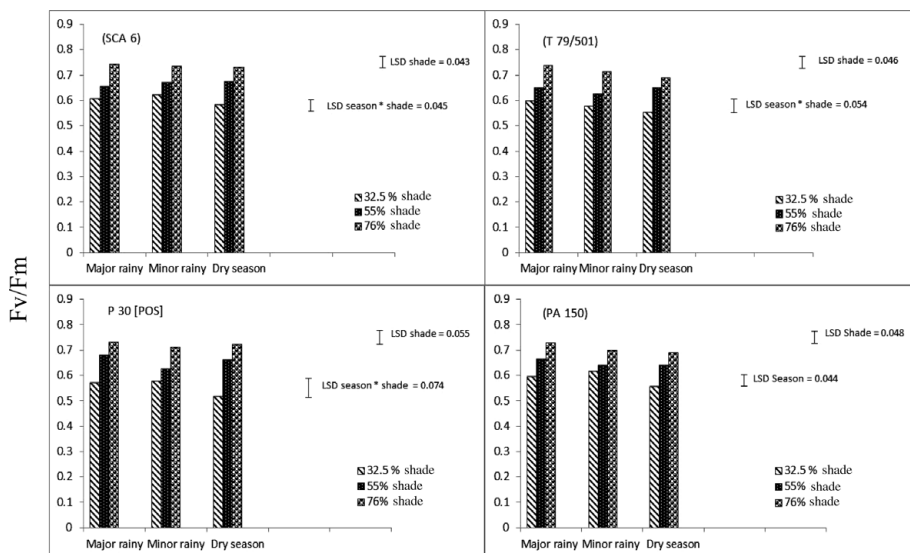


Figure 4. Leaf chlorophyll fluorescence of cacao under different shade levels. (Each bar represents the mean of four measurements on nine plants per shade treatment).

Seasonal variation in leaf chlorophyll fluorescence

Variable (Fv)/maximum (Fm) fluorescence values increased with greater shade level through all seasons ($p < 0.001$). Also, there was a slight reduction in Fv/Fm during the dry season, which was more pronounced under the light shade treatment. Thus, the season \times shade interaction was significant ($p = 0.013$). Small but significant ($p = 0.006$) differences were observed between the cacao genotypes in their Fv/Fm values, which were consistent between shade levels and between seasons (Figure 4). The respective Fv/Fm ratios for T 79/501, P 30 [POS], PA 150 and SCA 6 were 0.643, 0.644, 0.648 and 0.669 respectively.

Diurnal pattern of chlorophyll fluorescence

Values of Fv/Fm measured over a single day declined towards the mid part of the day and then increased during the second half of the day ($p = 0.001$). It was also consistently higher ($p < 0.001$) under heavier shade (Figure 5) although the nature of the response to shade differed between clones. Differences between the clones were significant ($p < 0.001$), varying from 0.650 for PA 150 to 0.679 for T 79/501. Whereas the time \times clone interaction was not significant, the shade \times clone interaction was significant ($p < 0.001$), reflecting the fact that suppression of Fv/Fm under higher light conditions was greater for some clones (e.g. SCA 6) than others.

Leaf production

Plants under heavy shade produced the lowest final leaf area per plant (Figure 6). The shade \times genotype interaction was significant ($p < 0.04$) such

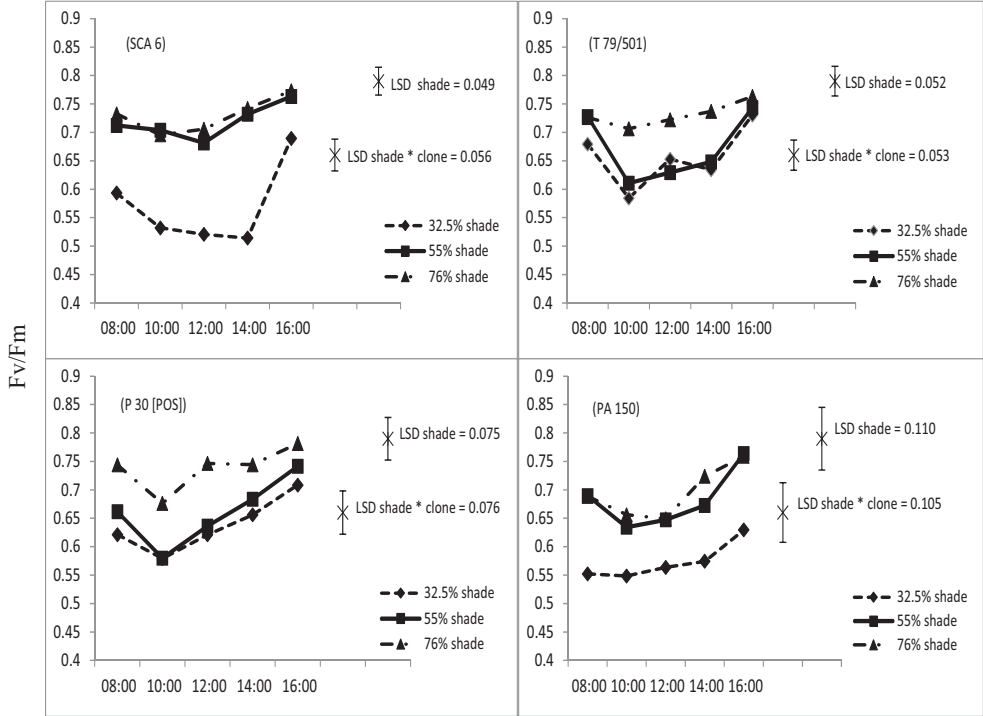


Figure 5. Diurnal pattern of leaf chlorophyll fluorescence of cacao under varying shade. (Each line represents the mean for nine plants per shade treatment).

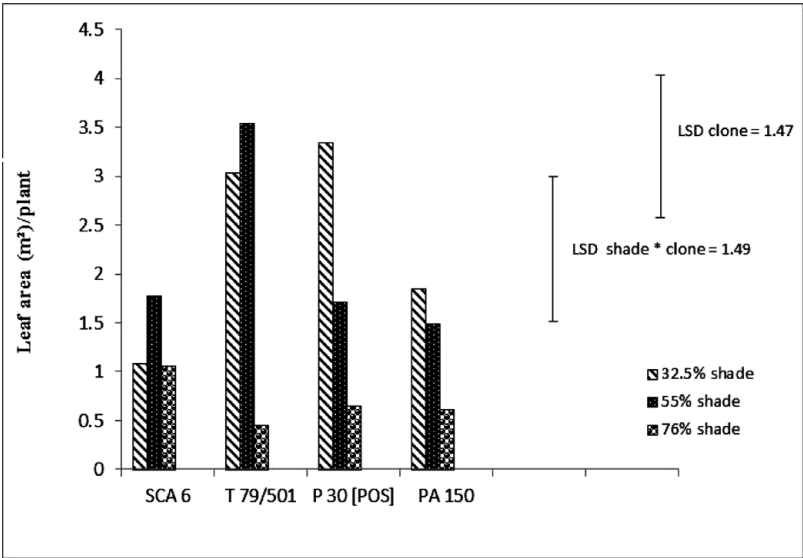


Figure 6. Final leaf area per plant under different shade levels. (Each bar represents the mean value for six plants per treatment).

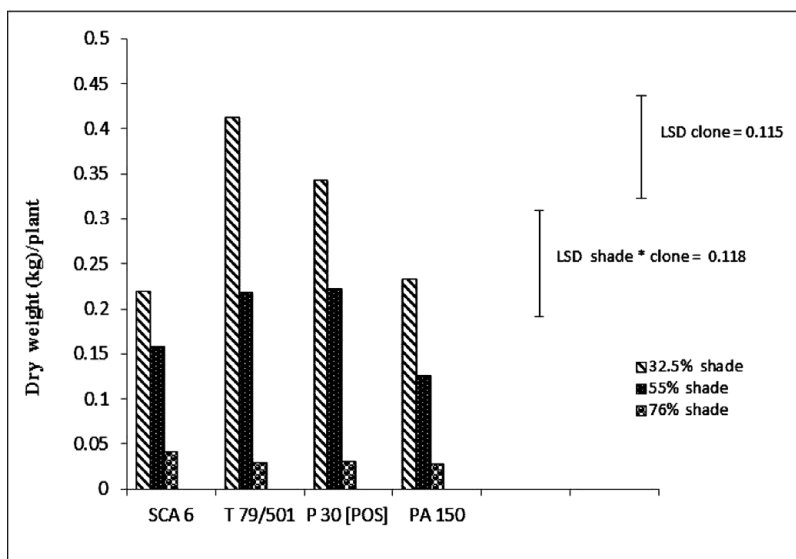


Figure 7. Mean final stem dry weight of cacao under different shade levels. (Each bar represents the mean dry weight of six plants per treatment).

that SCA 6 and T 79/501 gave the highest leaf areas under the medium shade, while P 30 [POS] and PA 150 gave the highest leaf areas under the light shade treatment.

Stem dry weight

For all genotypes, stem dry matter accumulation was significantly ($p = 0.003$) lower under increasing shade. The response to shade differed between clones, it being the greatest for clone T 79/501 and the least for SCA 6 (Figure 7).

DISCUSSION

Photosynthetic rates were the highest under light shade in the rainy seasons for all four genotypes studied. During the dry season, however, photosynthetic rates were much lower and the differences between treatments were smaller, the trend being opposite to that seen in the wet seasons, i.e. rates were generally higher under heavier shade. Since soil water was not limiting in this experiment, the seasonal variation in photosynthetic rate was most likely modulated by changes in vapour pressure deficit. Previous studies have shown that stomatal conductance in cacao is limited by vapour pressure deficit above a particular threshold (Raja Harun and Hardwick, 1988a; Sena Gomes *et al.*, 1987). Therefore, during the rainy seasons when vapour pressure deficit would not severely limit stomatal opening, light was the main limiting factor (light levels are typically lower in the minor rainy seasons, hence the lower photosynthetic rates observed). In contrast, during the dry season, vapour pressure deficit appeared to be a limiting factor, particularly for plants grown under the low shade (high light) treatment. Thus, the ameliorating effects of higher shade in the dry season (resulting

in a slightly lower vapour pressure deficit) resulted in a higher photosynthetic rate under this treatment.

Some evidence for photoinhibition was observed as Fv/Fm values were lower under higher light conditions. However, this photoinhibition was not so significant as to prevent the highest photosynthetic rate being expressed under the light shade regime during the wet seasons. In the dry season the difference between shade treatments was slightly greater, presumably due to the fact that ambient light levels were higher at this time of the year.

There was an evidence for photoinhibition from the diurnal chlorophyll fluorescence measurements typically from late morning until afternoon. This was especially the case under low shade conditions and was consistent with the observations of Serrano and Biehl (1996). By the end of the day, Fv/Fm values had typically recovered to, or were higher than early morning values, suggesting that the photoinhibition observed was partially reversible. Nevertheless, for all four clones, values of Fv/Fm remained lower under the high light conditions at both beginning and end of the day, implying that non-reversible inhibition was also present. The results also imply that in terms of photoinhibition clones SCA 6 and PA 150 were the most sensitive to high light intensities.

When considering net assimilation over the course of a day, although a constant saturating irradiance was applied, photosynthetic rates varied through time with higher rates occurring in the morning than in the afternoon. This was observed in spite of the fact that for most treatments Fv/Fm recovered through the course of the afternoon. Since vapour pressure deficits are typically higher in the afternoon, this may have been the cause of the reduction in photosynthetic rate. Previous studies have suggested that even in the case of mature cacao, some overhead shade should be provided, otherwise the photosynthetic rate of cacao is likely to decrease (Galyuon *et al.*, 1996b; Hutcheon, 1981; Raja Harun and Hardwick, 1988b). While this is a factor to consider in shading, the results of the present study suggest that the amelioration of high vapour pressure deficit is also a major consideration.

The larger leaf area observed with increasing light is not only reflective of higher photosynthetic rates but also implies that there was no greater leaf abscission under high light as earlier observed by Ofori-Frimpong *et al.* (2007). Genotypic differences in leaf area under the different shade levels suggest differential partitioning of assimilates in response to light. This differential partitioning was probably the main factor underlying the different magnitudes of response to shade in terms of final biomass between the four cacao clones studied. Furthermore, since there were no big differences in photosynthetic rates between genotypes, the differential partitioning of assimilate to the leaf component was an important factor in the observed overall genotypic difference in biomass.

The observation that both season and time of the day influence the photosynthetic performance of cacao under different light regimes may explain the considerable inconsistency observed in what is regarded as the optimal shade level for the growth of cacao. The fact that over the entire period, plants under light shade had photosynthetic rates that were over three times faster than those under the heavy shade treatment

(which was also reflected in the plants' growth) demonstrates that, with enough soil water and in the absence of severe mirid infestations (Babin *et al.*, 2010), young cacao can be grown under less shade than has been generally been recommended (usually around 50% shade; Opoku-Ameyaw *et al.*, 2010).

The use of overhead shade may, nevertheless, be used by the farmer for other reasons, such as provision of an alternate crop or aiding in pest control. This study has demonstrated genotypic differences in early growth under shade and therefore some genotypes are likely to be more suited than others to growth under shade. This work has also shown that for young cacao, shade should be provided on the basis of the current ambient environment and that good shade management needs to take into account changes in the aerial environment from wet season to dry season.

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