

**UNIVERSIDADE ESTADUAL DO MARANHÃO - UEMA**  
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**PROGRAMA DE PÓS-GRADUAÇÃO EM AGROECOLOGIA**

**LORENA SILVA CAMPOS**

**RESPOSTAS ECOFISIOLÓGICAS EM MILHO CULTIVADO COM GESSO E  
BIOMASSA DE LEGUMINOSA EM UM AGROSSISTEMA TROPICAL**

**SÃO LUÍS - MA**  
**2020**

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**Engenheira Agrônoma**

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Dissertação apresentada à Universidade Estadual do Maranhão, como parte das exigências do Programa de Pós-Graduação em Agroecologia, para obtenção do título de Mestre em Agroecologia.

Orientador: Prof. Dr. Tiago Massi Ferraz

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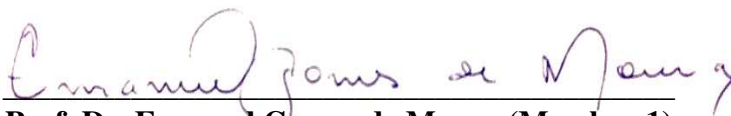
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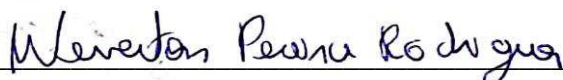
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*“A menos que modifiquemos a nossa maneira de pensar, não seremos capazes de resolver os problemas causados pela forma como nos acostumamos a ver o mundo”.*

Albert Einstein

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## RESUMO

Os modelos climáticos consistentemente projetam aumentos na variabilidade da temperatura nos países tropicais nas próximas décadas, com a Amazônia como um ponto de preocupação. Os limites para os quais os sistemas agrícolas podem se adaptar ao aquecimento global dependem, em parte, das melhorias das propriedades físicas do solo que favoreçam o aumento da absorção de água e nutrientes. Utilizamos gesso e biomassa de leguminosa para melhorar a zona radicular do solo e nossa hipótese é que tais alterações podem melhorar as condições físicas do solo tropical propenso à coesão e aumentar as respostas ecofisiológicas no milho. O estudo teve quatro repetições dos seguintes tratamentos: controle (C), gesso (G), nitrogênio (N), biomassa de leguminosa (L), nitrogênio + gesso (NG), biomassa de leguminosa + gesso (LG) e biomassa de leguminosa + nitrogênio + gesso (LNG). Os tratamentos foram submetidas ao regime sequeiro em 2018 e irrigação em regime de restrição hídrica em 2019. Os resultados das melhorias nos atributos do solo tiveram efeitos positivos nas respostas ecofisiológicas do milho, com uma redução no estresse das plantas principalmente no índice de performance ( $PI_{abs}$ ) que mostrou ser um índice sensível às adições de gesso e nitrogênio. Essas mudanças foram capazes de produzir diferenças na absorção de N, produção de biomassa e rendimento de grãos de milho, mesmo em regime de restrição de água. Sob regime de restrição de água, o rendimento de grãos de milho no tratamento LNG foi 41% maior do que no tratamento N e 151% maior do que no tratamento C. Além disso, a adição de nitrogênio diminuiu consideravelmente o estresse das plantas, com efeitos positivos nas respostas fisiológicas e no aumento da produção de biomassa. Esses resultados destacam o papel do nitrogênio em conjunto com a biomassa de leguminosa e o gesso na melhoria do solo, especialmente no contexto da adaptação às mudanças climáticas.

**Palavras-chave:** biomassa de leguminosa; manejo do solo; fotossíntese; milho; adaptação às mudanças climáticas

## ABSTRACT

Climate models consistently project increases in temperature variability in tropical countries over the coming decades, with the Amazon a particular hotspot of concern. The extent to which agricultural systems can adapt to global warming depends in part on physical improvements to soil that enhance water and nutrient uptake. We use gypsum and leguminous residue to improve the soil root zone, hypothesizing that such changes may improve the physical conditions of tropical soil prone to hardness and enhance the ecophysiological responses in maize. The study consisted of four replicates using the following treatments: control (C), gypsum (G), nitrogen (N), legume residue (L), nitrogen + gypsum (NG), legume residue + gypsum (LG) and legume residue + nitrogen + gypsum (LNG). The study plots were rainfed in 2018 and used a restricted irrigation regime in 2019. The soil improvements studied had positive effects on the physiological parameters of maize, with a reduction in stress mainly in performance index ( $PI_{abs}$ ) which indicated the sensitivity of maize plants to gypsum and nitrogen addition. More specifically, we observed changes in N uptake, biomass production and maize grain yield, even in the restricted irrigation regime. Under a restricted irrigation regime, maize grain yield in the LNG plots was 41% higher than in N plots and 151% higher than the C plots. Interestingly, additions of nitrogen considerably decreased plants stress, improving physiological parameters and increasing biomass production. These results highlight the role for added nitrogen alongside soil improvement measures such as leguminous residues and gypsum, especially in the context of climate change adaptation.

**Keywords:** legume residue; soil management; photosynthesis; maize; climate change adaptation.

# **CAPÍTULO I**

## 1 INTRODUÇÃO GERAL

Com o aquecimento global é previsto que o rendimento das culturas nas regiões de clima tropical e subtropical seja reduzido como resultado de uma aceleração do desenvolvimento fenológico (TURNER e RAO, 2013). Modelos climáticos projetam diferenças nas características climáticas regionais entre os dias atuais e o aquecimento global de 1,5°C a 2°C. Essas diferenças incluem aumentos na temperatura média na maioria das regiões terrestres e oceânicas, nos extremos de calor na maioria das regiões habitadas, na ocorrência de chuva intensa em diversas regiões e na probabilidade de seca e déficits de chuva em algumas regiões (IPCC, 2019)

O déficit hídrico é a causa mais grave de redução da produção agrícola global e as mudanças climáticas tendem a agravar esse efeito. Além disso, existe o desafio de aumentar a oferta de alimentos nos cenários de estresse em cerca de 60% para atender o aumento da população mundial, que excederá 9 bilhões de pessoas até 2050 (GRASSINI et al., 2015). Com os adventos climáticos, o desenvolvimento da agricultura terá o desafio de buscar a economia de água e o aumento da eficiência do seu uso. Os ambientes tropicais que apresentam períodos de estiagem estão mais sujeitos a fatores estressantes, com a consequente diminuição da produção de biomassa (TURNER e RAO, 2013).

Outro fator de elevada importância que induz a diminuição e limita o crescimento e a produção das culturas é a baixa disponibilidade de nutrientes (VANCE et al., 2003). Em condições meteorológicas tropicais, devido à interação negativa entre a inadequada condição do solo e a alta demanda evaporativa da atmosfera, as culturas nem sempre alcançam seu potencial de produtividade, tornando esses agrossistemas inviáveis, favorecendo a insegurança alimentar (MOURA et al., 2013).

A maioria dos solos da região tropical não possui alta resiliência intrínseca à degradação física devido ao baixo conteúdo de elementos agregadores, como carbono, cálcio e magnésio (MOURA et al., 2018). Nesse solo, também conhecido como solo coeso (DANIELLS, 2012), as raízes das plantas podem explorar apenas uma fina camada superficial para sustentar o desenvolvimento e a produtividade das culturas. Portanto, a melhoria das propriedades físicas do solo para aumentar o aproveitamento da água no solo será crucial para diminuir o efeito deletério do aquecimento global sobre a produtividade das culturas na agricultura tropical (EBSA, 2019). O plantio direto com o uso de biomassa de leguminosas é um sistema em que

utilizam-se ramos e folhas de leguminosas aplicados à superfície para inicialmente servir de cobertura e posterior retorno de nutrientes ao solo durante a decomposição (TSONKONVA et al., 2012). Essa camada protetora permite que o solo permaneça úmido reduzindo a evaporação a partir da superfície por mais tempo o que favorece a disponibilidade de água para as culturas e com isso possa aumentar a taxa fotossintética (KADER et al., 2019).

Segundo Kluge et al. (2015), o dióxido de carbono ( $\text{CO}_2$ ) entra na folha ao mesmo tempo que a água, na forma de vapor, é perdida através dos estômatos da epiderme. Como planta C4, o milho (*Zea mays* L.) possui alta eficiência fotossintética mesmo com intensidade luminosa elevada. No entanto, quando ocorre o déficit hídrico, os estômatos se fecham para conservar água, reduzem a difusão de  $\text{CO}_2$  nos locais de fixação no mesófilo foliar nas proximidades da enzima ribulose 1,5-bifosfato carboxilase oxigenase (Rubisco), o que diminui a fotossíntese pois esta é indiretamente afetada pela sensibilidade estomática e, assim, consequentemente poderá haver perdas na produtividade (SILVA et al., 2013). Entretanto, o rendimento em determinadas culturas, não depende apenas da taxa fotossintética, mas também do comprimento da fase ativa da fotossíntese das folhas e da capacidade de realocação para os grãos (FUHRER, 2003). Boretti e Florentine (2019) alertam que embora o aumento da concentração de dióxido de carbono no contexto do aquecimento global possa contribuir para um maior crescimento das plantas, sob condições meteorológicas tropicais isso pode não ocorrer devido ao déficit hídrico.

A cobertura morta de biomassa de árvores leguminosas tem sido especialmente recomendada como alternativa para o manejo dos solos coesos por diminuir a evaporação, mantendo a umidade do solo mesmo sob alta taxa de evaporação da água (MOURA et al., 2018; KADER et al., 2019). No entanto, o efeito a curto prazo da cobertura morta tende a se concentrar nas camadas superficiais do solo (BLANCO-CANQUI et al., 2007).

Para aumentar a captação de água pelas raízes em camadas mais profundas do solo, o uso do gesso tem sido recomendado devido à sua alta solubilidade e rápido enriquecimento por cálcio das camadas inferiores do perfil (SUMNER, 2009). Entretanto, para garantir a produção das culturas agrícolas destinadas a alimentar uma população em crescimento e em cenários de crescente falta de água, é necessário ampliar o conhecimento acerca das práticas que minimizem a perda e aumente o aproveitamento da água nos cultivos a fim de evitar o estresse pela falta de água na planta.

O nitrogênio (N) faz parte dos principais componentes do sistema fotossintético, tais como clorofilas, rubisco e fosfoenolpiruvato carboxilase – PEPcase. O incremento na disponibilidade de nitrogênio geralmente traz efeitos benéficos na taxa de assimilação de

carbono, já que esse nutriente faz parte dos principais componentes do sistema fotossintético (CORREIA et al., 2005). Em solo tropical intemperizado, os serviços fornecidos pelos ecossistemas naturais, como a ciclagem de nutrientes na zona das raízes e fixação do nitrogênio são essenciais para a manutenção da estrutura e aumento da fertilidade do solo. Esses processos impulsionam a eficiência pela qual as plantas adquirem o N, que foi apontado como um dos mais importantes para melhorar a eficiência de absorção de raízes (Chapman et al. 2012). Isso é particularmente verdadeiro em condições tropicais úmidas, onde a eficiência do uso do nitrogênio é geralmente muito baixa devido às altas taxas de volatilização da amônia e ao aumento da remoção da água de drenagem devido às altas temperaturas e altas precipitação (Jabloun et al. 2015).

A hipótese deste trabalho é que a utilização de gesso e biomassa de leguminosa poderá melhorar as condições físicas da zona radicular do solo tropical e elevar as respostas ecofisiológicas do milho, como estratégia para aumentar a adaptação às mudanças climáticas na agricultura tropical. Portanto, este estudo teve como objetivo avaliar como as respostas ecofisiológicas do milho são afetadas pelos efeitos combinados do gesso com biomassa de leguminosa como cobertura morta no solo. O objetivo final é contribuir para a adaptação da agricultura familiar ao aquecimento global previsto na periferia da Amazônia.

## **2 REVISÃO DE LITERATURA**

### **2.1 Milho: Características gerais**

O milho é uma planta monoica, membro da Família Poaceae e pertence ao gênero *Zea*, um grupo de gramíneas anuais e perenes, nativas do México e da América Central. É um dos principais cereais cultivados no mundo e os grãos podem ser consumidos diretamente, sem a necessidade de processamento para remover a “casca”, como é feito com outros cereais, como arroz e trigo. A partir do milho são originados vários produtos destinados à alimentação humana e animal. Além disso, é matéria prima para a agroindústria. Em termos nutricionais, o milho possui uma composição rica em carboidratos, principalmente na forma de amido, além de proteínas, lipídios, vitaminas e minerais (OLIVEIRA et al., 2014; LANGNER et al., 2019).

O rendimento de grãos de milho é determinado pelo resultado da interação de genótipo, manejo e fatores ambientais. Os fatores ambientais mais importantes são radiação solar, precipitação e temperatura. Esses fatores não podem ser controlados pelo produtor e variam com a época do ano (FAGERIA et al., 2006). A fotossíntese durante a fase de crescimento e a

eficiência com que o fotossintato é realocado para os grãos são processos fundamentais na determinação do desenvolvimento da planta de milho e conseqüentemente no seu rendimento. Práticas de manejo como plantio direto, irrigação, suprimento de nutrientes e manejo de pragas são utilizados para maximizar o rendimento econômico, mas as respostas a essas práticas variam entre os ambientes.

O milho tem um grande impacto na economia dos países desenvolvidos assim como dos países em desenvolvimento. É cultivado em todas as regiões do território brasileiro e sua produção ocorre em diferentes épocas, a depender das condições climáticas das regiões. Devido a uma gama de cultivares disponíveis no mercado com grande adaptabilidade é possível o cultivo do milho em épocas distintas. Considerado como uma das principais culturas da agricultura brasileira, o milho está presente em diversos sistemas de cultivo como em consórcio com gramíneas forrageiras para compor sistemas integrados de produção lavoura-pecuária, em cultivo consorciado com espécies arbóreas leguminosas, em esquema de rotação de culturas no sistema de plantio direto, etc. (CONTINI et al., 2019). A produtividade média do milho no Brasil está por volta de 5,3 t/ha e encontra-se abaixo da produtividade mundial que é de aproximadamente 5,65 t/ha (USDA, 2019).

## **2.2 Parâmetros fisiológicos**

### **2.2.1 Trocas gasosas no milho**

A fotossíntese é o processo usado pelas plantas para capturar energia da luz solar e convertê-la em energia bioquímica, que é posteriormente usada para suportar quase toda a vida na Terra. Esse processo é complexadamente regulado por fatores biofísicos, bioquímicos, fisiológicos e estruturais das folhas assim como alguns órgãos fotossintetizantes. Apesar de muitos avanços alcançados neste campo, nossa compreensão dos processos regulatórios da fotossíntese ainda é insatisfatório (EVANS, 2013). Os fatores que causam a variação genética na fotossíntese também permanecem amplamente desconhecidos (FLOOD et al., 2011).

O milho pode suportar luz de alta intensidade ao meio-dia e o padrão diurno de sua taxa fotossintética corresponde à curva da intensidade da luz diurna com um pequeno atraso, ou seja, a sua taxa fotossintética tem uma curva de pico único. Plantas com esse padrão de taxa de fotossíntese na forma de um pico podem abrir mais os estômatos e sofrer menos fotoinibição do que aquelas com padrões de taxa de dois picos, como as de arroz por exemplo, que sofre



fotoinibição ao meio-dia, quando a alta intensidade luminosa causa fechamento dos estômatos e baixa eficiência nos cloroplastos (VAN et al., 2011; FENG et al., 2017).

Um importante mecanismo regulador da fotossíntese é o controle dos níveis de proteínas enzimáticas. Durante um ciclo diurno com intensidade luminosa variável, as folhas otimizam a assimilação fotossintética de CO<sub>2</sub> para se adaptar ao ambiente, ajustando os níveis de enzimas. Porém, entre as proteínas relacionadas à fotossíntese identificadas no estudo feito por Feng et al. (2017), as de antenas da fotossíntese das folhas de milho permanecem em níveis quase constantes durante todo o ciclo diurno, indicando que a captação de luz na clorofila não muda com a intensidade de luz variável. Os autores concluem que as folhas de milho não ajustam os níveis de proteína da antena da fotossíntese em resposta à mudança da intensidade da luz. No entanto, proteínas no fotossistema exibem expressão diferencial durante o dia.

Sabe-se que as plantas C<sub>4</sub>, como o milho, têm maior assimilação fotossintética de CO<sub>2</sub> e maior produtividade biológica do que as plantas C<sub>3</sub> em condições tropicais (BROWN, 1999). Isso é obtido por um mecanismo de concentração de CO<sub>2</sub> que atua em plantas C<sub>4</sub>. Nas folhas de plantas C<sub>4</sub>, as células do mesófilo e da bainha são diferenciadas e envolvem os feixes vasculares. Na fotossíntese C<sub>4</sub>, o CO<sub>2</sub> atmosférico que entra nos estômatos é fixado pela PEP<sub>case</sub> nas células do mesófilo. Os ácidos C<sub>4</sub> produzidos são transportados para as células da região interna da bainha do feixe vascular (células BS), onde são descarboxilados pela descarboxilase do ácido C<sub>4</sub>. O CO<sub>2</sub> liberado é fixado pela rubisco. Esse processo bioquímico aumenta a concentração de CO<sub>2</sub> em torno da Rubisco nas células da bainha do feixe vascular e reduz a fotorrespiração (HATCH, 1987).

A abertura estomática está diretamente ligada à regulação das trocas gasosas, de modo que por meio da variação na abertura estomática é possível controlar o fluxo de CO<sub>2</sub> e água da folha. Conforme descrito por Kluge et al. (2015), o CO<sub>2</sub> entra na folha e a água, na forma de vapor, é perdida através dos estômatos da epiderme. Os estômatos podem fechar se o suprimento de água for insuficiente, podendo reduzir o suprimento de CO<sub>2</sub> para a fotossíntese.

Fatores como luz, temperatura, concentração de CO<sub>2</sub>, teor de nitrogênio da folha e umidade do solo afetam a atividade fotossintética dos vegetais, sendo que, o processo de abertura e fechamento dos estômatos está relacionado principalmente com a intensidade e qualidade de luz, concentração de CO<sub>2</sub> e o teor de água na folha. Assim, a abertura estomática e a área foliar influenciam diretamente a produtividade do vegetal, o primeiro porque controla a absorção de CO<sub>2</sub> e o segundo porque determina a interceptação luminosa (COSTA e MARENCO, 2007). Lopes et al. (2009) avaliaram a taxa fotossintética líquida na cultura de milho no sudeste brasileiro em plantio convencional e direto com biomassa de triticale (X

*Triticosecale*) e obtiveram valor máximo de  $48 \mu\text{mol CO}_2\text{m}^{-2}\text{s}^{-1}$  por volta das 11h e valor mínimo de  $20 \mu\text{mol m}^{-2}\text{s}^{-1}$  por volta das 7h para ambos os tipos de plantio no estádio V8 e concluíram que esses manejos não afetaram essa variável.

A demanda por água da atmosfera pode ser avaliada pela diferença de déficit de pressão de vapor entre a folha e o ar ( $\text{DPV}_{\text{folha-ar}}$ ), que envolve a temperatura da folha e a temperatura do ar e essa variável pode afetar a saída de água da folha, por ação negativa sobre a condutância estomática (FRANKS e FARQUHAR, 2007; REIS et. al., 2009). De acordo com Taiz et al. (2017), quanto maior o  $\text{DPV}_{\text{folha-ar}}$ , maior a tendência da planta em perder água. Por intermédio da análise quantitativa do crescimento e de avaliações de trocas gasosas periódicas pode-se avaliar os efeitos dos tipos de manejo sobre as plantas, pois as mudanças nas características do crescimento são observadas em função do tempo, o que não é possível somente com o registro do rendimento do grão (URCHEI et al., 2000).

### 2.2.2 Fluorescência da clorofila

Embora a fotossíntese possa ser medida diretamente pela absorção de  $\text{CO}_2$ , para melhor compreensão, o desempenho fotossintético pode ser avaliado juntamente com outros parâmetros. Medidas da fluorescência da clorofila *a* apresentam-se como um complemento em estudos ecofisiológicos de plantas. O rendimento da fluorescência da clorofila nos informa o nível de excitação da energia no sistema de pigmentos que comandam a fotossíntese e é uma excelente ferramenta de monitoramento para tolerância de plantas de milho a condições adversas, principalmente estresse hídrico (MAGALHÃES et al., 2013).

Em outras palavras, as moléculas de clorofila são capazes de reemitir, na forma de fluorescência, uma porcentagem da luz absorvida que não é transformada em energia química. Assim, a intensidade da fluorescência pode ser correlacionada com a atividade do transporte fotossintético de elétrons (CARELLI, 2017). Um dos parâmetros mais frequentemente empregados para esta medida é o rendimento quântico máximo do fotossistema II (relação  $F_v/F_m$ ), que fornece informações sobre a proporção da luz absorvida pela clorofila no PSII usada em processos fotoquímico (LEPEDUŠ et al., 2012).

Pinto et al. (2012) afirmaram que reduções significativas nos valores dos rendimentos quânticos máximos do fotossistema II, bem como nos valores do índice de desempenho - PI, indicam efeito de fotoinibição. Tais mudanças podem representar perda de eficiência fotoquímica pelas plantas e, como consequência, a diminuição da fotossíntese. Valores baixos do índice de desempenho indicam que apenas uma pequena proporção dos fótons absorvidos

foi usada no transporte de elétrons, a maioria sendo dissipada como calor. O índice de performance (PI) vem sendo utilizado com mais intensidade para avaliar plantas sob condições adversas. Ele reflete a funcionalidade de ambos os fotossistemas (I e II) e fornece informações quantitativas sobre o estado atual do desempenho da planta. (ŽIVČÁK et al., 2008).

O índice de desempenho é obtido com emprego de fluorímetros, que integra três variáveis independentes sendo: a densidade dos centros de reação ativos (RC) por unidade de moléculas de clorofilas no sistema antena do PSII (RC/ABS); a eficiência do elétron que reduziu Quinona A (Qa) e entrou na cadeia transportadora de elétrons (ET/(TR-ET)); a relação entre a capacidade de redução de Qa pelo fóton absorvido, e a energia dissipada na forma de calor ( $F_v/F_0$ ). Silva et al. (2013) afirmam que a alta correlação entre os sintomas de estresse e a redução da performance fotossintética em plantas reforça a utilidade do índice de performance como indicador de estresse.

### **2.3 Estresse em plantas de milho**

A definição de estresse é complexa porém é um conceito de grande importância para a ecologia e agricultura. Uma motivação para o estudo do estresse na ecofisiologia das plantas é a consideração sobre os padrões de distribuição e desempenho das plantas ao longo dos gradientes ambientais (OSMOND et al., 1987). Nas culturas, os efeitos estressantes dos ambientes afetam o rendimento de grãos a partir da diminuição dos componentes de rendimento. O estresse durante o crescimento vegetativo afeta o número de grãos devido ao fato de que o tamanho da espiga e o número de óvulos formados são determinados durante esse estágio (ABENDROTH et al., 2011).

O déficit hídrico é a causa mais grave e limitante de redução da produção agrícola global e as respostas das plantas à escassez de água são diferentes e dependem da intensidade e do tempo de estresse. O déficit hídrico em baixa intensidade leva à redução da transpiração, falha na translocação da água das raízes para os rebentos, presença de menos pigmentos fotossintéticos, limitação dos produtos fotossintéticos (HOSSEINZADEH et al., 2017). O déficit hídrico em alta intensidade resulta na formação de espécies reativas de oxigênio (ROS) nas plantas, que tem um efeito prejudicial sobre a proteína D1 do PSII, conteúdo de clorofila, transporte de elétrons e produção de moléculas como o trifosfato de adenosina (ATP) (PAGTER et al., 2005). Lu et al. (2002) relataram que o milho sob estresse hídrico teve uma diminuição em  $F_v/F_m$  em relação aos tratamentos sob estresse não hídrico. Sob condições de estresse hídrico, a proteína D1 do PSII e os centros de reação do PSII são destruídos. Bolhàr-

Nordenkamp et al. (1989) relatam que a razão  $F_v/F_m$  varia entre 0,75 e 0,85 quando a planta está com seu aparato fotossintético intacto e uma queda nesta razão reflete a presença de danos nos centros de reação do PSII por algum estresse.

Como planta C4, o milho possui alta eficiência fotossintética mesmo com intensidade luminosa elevada. No entanto, a fotossíntese é particularmente sensível ao déficit hídrico, pois os estômatos tendem a fechar para conservar água interna, com isso reduzem a difusão de  $CO_2$  nos locais de fixação no mesófilo foliar nas proximidades da enzima rubisco. Esse processo diminui a fotossíntese e, conseqüentemente, reduz a produtividade. Entretanto, os estômatos podem ser regulados com base no nível de déficit hídrico fechando apenas parcialmente, levando a alguma fixação de carbono durante condições de seca e assim um aumento na eficiência do uso da água (SILVA et al., 2013).

O resultado da escassez de água é importante principalmente nos seguintes estádios de desenvolvimento da planta de milho: I - iniciação floral e surgimento da inflorescência, período em que o número potencial de grãos é estabelecido; II - época da fertilização, quando o potencial de produção é determinado e a presença da água evita o dessecamento do grão de pólen e assegura o desenvolvimento e a penetração do tubo polínico; III- enchimento dos grãos, quando há o acréscimo na deposição de matéria seca, processo diretamente ligado à fotossíntese. O estresse nesta fase culminaria na menor produção de fotossintatos, o que acarretaria menor quantidade de matéria seca nos grãos (MAGALHÃES et al., 2003).

Os efeitos da condutância estomática na tolerância do déficit de água na planta são complexos. Em algumas dicotiledôneas C4, modificações na arquitetura do xilema, incluindo vasos mais estreitos e mais curtos, diminuem a condutividade do caule por unidade de área foliar, o que fornece proteção contra cavitação e, portanto, melhora a tolerância à seca (SAGE et al., 2012). Pode-se supor que a maior eficiência no uso da água conferida pela fisiologia das C4 permitiu uma evolução no sentido de reduções na condutividade do caule por unidade de área foliar (CHRISTIN e OSBORNE, 2014).

## **2.4 Manejo do Solo: O uso de biomassa vegetal e gesso**

O manejo do solo tem grandes efeitos na melhoria das suas propriedades físicas e químicas na qualidade das terras agrícolas e na eficiência fotossintética do milho (SUN et al., 2019). Estratégias para facilitar a penetração e alcance da raiz no solo e para aumentar a absorção de nutrientes incluem o seguinte: I) uso apropriado da cobertura morta; II) a aplicação

de biomassa para aumentar a matéria orgânica do solo; III) a adição de cátions básicos, como cálcio e magnésio; e IV) estabelecimento de equilíbrio iônico adequado do solo.

As coberturas mortas podem atrasar o endurecimento do solo e favorecer o crescimento das raízes, reduzindo a evaporação da superfície do solo (MULUMBA E LAL, 2008). Além disso, a aplicação contínua de biomassa, como ocorre com a cobertura morta, pode aumentar as frações lábeis da matéria orgânica, criando agregados que, embora instáveis, melhoram a porosidade e, portanto, o ambiente do solo para o crescimento radicular (PASSIOURA et al., 1991). Infelizmente, nas regiões tropicais, o acúmulo de matéria orgânica, que poderia mitigar os efeitos negativos da coesão do solo na região da raiz, é prejudicado por condições que favorecem a rápida deterioração da biomassa incorporada (HIJBEEK et al., 2018).

O uso do gesso também vem sendo recomendado para os solos tropicais, pois o cálcio compreende um nutriente importante para o crescimento radicular das plantas. Para estender a capacidade das raízes nas camadas mais profundas do solo, pesquisas sugerem o uso de cálcio e magnésio como “agentes flocculantes” para melhorar a estrutura do solo e reduzir a dispersão da argila (ANIKWE et al., 2016).

A aplicação de gesso com biomassa de leguminosas modifica a zona radicular, aumentando o teor de cálcio e matéria orgânica e reduzindo a resistência do solo (MOURA et al., 2018). Alguns autores enfatizam interações positivas entre o cálcio e a matéria orgânica composta a partir do resíduo aplicado como cobertura morta (WUDDIVIRA e CAMPS-ROACH, 2007). A formação de pontes de cátions com produtos derivados da decomposição da biomassa aplicada pode ser formada entre partículas carregadas negativamente, essencialmente ligando moléculas orgânicas em conjunto com os minerais (WHITTINGHILL e HOBBIE, 2012). A relação entre a melhoria da capacidade de suporte do solo, os fatores físicos e fisiológicos que sustentam o crescimento das plantas deve ser esclarecida para apoiar os esforços para evitar o desperdício de água e nutrientes e garantir a viabilidade dos sistemas agrícolas tropicais.

Em estudo realizado por Kusumah et al. (2019), a média total da produção de milho com cobertura morta aplicada de gliricídia (*Gliricida sepium*) foi de 4,43 ton/ha. A produção obtida foi muito maior em comparação à média da produção de milho dos agricultores de Merauke (Indonésia) que nos últimos 5 anos atingiram apenas 3,53 ton/ha. Os autores relatam que a maioria dos cultivos é de sequeiro e as altas temperaturas resultam em maior evaporação no solo, de modo que o estresse causado pela seca ocorre com frequência. Aguiar et al. (2018) obtiveram 4,1 ton/ha de rendimento de grãos quando utilizaram biomassa de gliricídia e clitoria

(*Clitoria fairchildiana*) com ureia no cultivo de milho em sistema de aleias no município de Chapadinha, MA, Brasil. O tratamento que recebeu somente ureia teve rendimento de grãos de 0.6 ton/ha. Sena et al. (2020) obtiveram 70% a mais de rendimento na produtividade de milho no leste maranhense no tratamento com biomassa de gliricidia e adição de ureia (5,3 ton/ha) em comparação ao solo desnudo com adição de ureia (3,1 ton/ha).

## 2.5 Aquecimento global

O clima é governado por influências naturais, mas as atividades humanas têm um impacto sobre ele também. O principal impacto que os humanos exercem sobre o clima é através da emissão de gases de efeito estufa. Gases capazes de absorver a energia radiante são chamados gases de efeito estufa (GEE). O desmatamento é outro exemplo de atividade que influencia o clima. As emissões globais antropogênicas de GEE têm aumentado desde os tempos pré-industriais, com um aumento de 90% entre 1970 e 2011, impulsionados pelo uso de combustíveis fósseis, mudanças no uso da terra e intensificação da agricultura (CLARKE et al., 2015).

O aquecimento global e as mudanças climáticas devido a processos naturais e fatores antropogênicos são as principais e mais importantes questões ambientais que afetarão o mundo (ALEXANDROV e HOOGENBOOM, 2000). Alimentar um mundo com nove bilhões de pessoas até 2050 exigirá um aumento substancial na produção de alimentos. A produção agrícola pode ser aumentada através da expansão da área cultivada ou pela intensificação da produção na área em uso. Em termos de eficácia da mitigação, a intensificação tende a ser preferível à expansão. A expansão pode causar emissões substanciais a partir da conversão de terras com altos estoques de carbono. Se bem administrada, a intensificação pode evitar a conversão de novas áreas em terras cultivadas, porque pressupõe maior produção agrícola na mesma área de terra (DICKIE et al., 2014).

Embora os pesquisadores ofereçam cenários diferentes, o consenso é de que a produtividade das culturas e dos animais pode diminuir devido às altas temperaturas e ao estresse relacionado à seca, mas esses efeitos serão variados entre diferentes regiões (ALTIERI et al., 2015). Alguns trabalhos baseados em modelagens, afirmam que temperaturas mais quentes podem fazer com que muitas culturas cresçam mais rapidamente, mas podem também reduzir o rendimento de outras muitas culturas. Para qualquer cultura em particular, o efeito do aumento da temperatura dependerá da temperatura ideal para crescimento e reprodução; em

áreas onde o aquecimento excede a temperatura ideal de uma espécie, os rendimentos daquela espécie podem diminuir (LOBELL e FIELD, 2011).

Meza et al. (2008) relataram que o milho pode ser afetado pelas mudanças climáticas, com reduções de produtividade entre 10% e 30%, dependendo do cenário das mudanças climáticas e do tipo de cultivar utilizada. As mudanças climáticas também afetarão outras variáveis relevantes da cultura como a taxa de desenvolvimento e a eficiência do milho para completar seu ciclo de crescimento em períodos mais curtos. Mudanças na precipitação sazonal total ou em seu padrão de variabilidade também afetarão a produção agrícola, mas grande parte dos modelos afirma que a maioria dos impactos é causada principalmente por tendências de temperatura e não por precipitação. As mudanças no rendimento das culturas de sequeiro serão impulsionadas por mudanças na precipitação e na temperatura, enquanto as mudanças no rendimento em áreas irrigadas serão impulsionadas apenas pelas mudanças de temperatura (ALTIERI et al. 2015).

Projeções futuras através de modelo climático mostram uma redução de até 46% na produtividade de milho na agricultura de sequeiro da China (TONG et al., 2016). Para o nordeste do Brasil, os resultados de um trabalho desenvolvido por Martins et al. (2019) apontam que os efeitos das mudanças climáticas na produtividade do milho na agricultura de sequeiro será negativo. As projeções sugerem que essas mudanças podem afetar a produção de milho nos dois cenários avaliados, resultando em um declínio na produção entre 30 e 60%. Os autores argumentam que para mitigar a situação a utilização de cultivares de ciclo longo trarão maior vantagem juntamente com a introdução da irrigação durante os períodos mais secos. Afirmam ainda que a irrigação e o desenvolvimento de novas variedades de milho seriam essenciais para sustentar a produtividade em cenários adversos de mudanças climáticas. No entanto, a manutenção de tais níveis de produção requer um aumento significativo no consumo de água (até 140%). Ahmed et al. (2019) sugerem o desenvolvimento de cultivares tolerantes ao calor e a modificação nas atuais tecnologias de produção agrícola para compensar os efeitos negativos das mudanças climáticas.

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## **CAPÍTULO II**

1 **Ecophysiological response of maize to soil management in the Amazonian periphery in the**  
2 **context of climate change adaptation**

3  
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37

38 **Abstract**

39

40 Climate models consistently project increases in temperature variability in tropical countries over the  
41 coming decades, with the Amazon a particular hotspot of concern. The extent to which agricultural  
42 systems can adapt to global warming depends in part on physical improvements to soil that enhance  
43 water and nutrient uptake. We use gypsum and leguminous residue to improve the soil root zone,

44 hypothesizing that such changes may improve the physical conditions of tropical soil prone to hardness  
45 and enhance the ecophysiological responses in maize. The study consisted of four replicates using the  
46 following treatments: control (C), gypsum (G), nitrogen (N), legume residue (L), nitrogen + gypsum  
47 (NG), legume residue + gypsum (LG) and legume residue + nitrogen + gypsum (LNG). The study plots  
48 were rainfed in 2018 and used a restricted irrigation regime in 2019. The soil improvements studied had  
49 positive effects on the physiological parameters of maize, with a reduction in stress which indicated the  
50 sensitivity of maize plants to gypsum and nitrogen addition. More specifically, we observed changes in  
51 N uptake, biomass production and maize grain yield, even in the restricted irrigation regime. Under a  
52 restricted irrigation regime, maize grain yield in the LNG plots was 41% higher than in N plots and  
53 151% higher than the C plots. Interestingly, additions of nitrogen considerably decreased plants stress,  
54 improving physiological parameters and increasing biomass production. These results highlight the role  
55 for added nitrogen alongside soil improvement measures such as leguminous residues and gypsum,  
56 especially in the context of climate change adaptation.

57

58 Keywords: legume residue; soil management; photosynthesis; maize; climate change adaptation.

## 59 **Introduction**

60

61 Climate models consistently project increases in temperature variability in tropical countries over the  
62 coming decades, with the Amazon a particular hotspot of concern (Bathiany et al. 2018). A number of  
63 harmful effects on agricultural production are predicted due to global warming, including extreme  
64 events such as heatwaves, excessive drying of soils, increased rainfall intensity and higher total  
65 precipitation (Gupta et al. 2020, Kamphorst et al. 2020). All of these mechanisms contribute to soil  
66 degradation in the humid tropics, particularly the leaching of base cations and increased rates of soil  
67 organic matter decomposition (Senna et al. 2020).

68 Although increased CO<sub>2</sub> concentrations [CO<sub>2</sub>] is predicted to impact on C3 plants to greater extent,  
69 recent studies have demonstrated positive effects of increased [CO<sub>2</sub>] in C4 plants such as maize (*Zea*  
70 *mays* L.) especially under adequate water supply (Meng et al. 2014). Therefore, improving the physical  
71 properties of soil to increase soil water uptake will be crucial to decrease the deleterious effect of global  
72 warming on tropical crop productivity (Ebsa 2019).

73 Overall, soils in Amazon region do not have a high intrinsic resilience against physical degradation due  
74 to the low content of aggregator elements like soil carbon, calcium and magnesium (Moura et al. 2018).  
75 In the natural conditions of the Amazonian periphery, the soil-vegetation system is resilient against  
76 physical degradation due to complete soil cover at ground level combined with the perforating activity  
77 of the soil fauna (Brinkman and Sombroek 1996). In agroecosystems, strategies using mulch application  
78 as a ground cover increase organic matter and avoid soil degradation by reducing cohesion and  
79 improving the root zone (Sena et al. 2020). Mulching with plant residues decreases the rate of

80 evaporation from the soil, delaying soil moisture loss, and improving rootability (Blanco-Canqui and  
81 Lal 2007). Such improvements, however, may be limited to the soil surface (Moura et al. 2012).

82 One solution to extend soil rootability to lower depths is using calcium in the form of gypsum, which  
83 acts as a “flocculating” agent to improve the soil structure by reducing the dispersion of clay at deeper  
84 layers (Sumner 2009). In addition, polyvalent cations like calcium derived from gypsum may interact  
85 with organic matter from decomposing plant residue, which further contributes to a favorable  
86 environment for root growth (Ellerbrock and Gerke 2018). These bonds prevent the biological,  
87 chemical, and physical breakdown of organic matter, leading to reduced losses of organic matter and  
88 soil base cations (Whittinghill and Hobbie, 2012).

89 Although environmental stresses are one of the most important factors that limit crop production, the  
90 responses of plants to water scarcity can be different depending on the intensity and time of stress  
91 (Kamphorst et al. 2020). Low-intensity water deficit leads to reduced transpiration, failure to translocate  
92 water from roots to shoots, decreased photosynthetic pigments and limiting production of photosynthetic  
93 products (Hosseinzadeh et al. 2017). High-intensity water deficit creates reactive oxygen species (ROS)  
94 in plants with detrimental effect on the D1 protein of photosystem II, chlorophyll content, electron  
95 transport, and the production of molecules such as adenosine triphosphate (ATP) (Pagter et al. 2005).  
96 Therefore, improvements to soil physical characteristics that enhance water and nutrient uptake are an  
97 important component of climate change adaptation for tropical agricultural systems (Moura et al. 2012).  
98 We hypothesize that improvements in the soil root zone from using gypsum along with leguminous  
99 residue might improve the physical conditions of tropical soil prone to hardening and enhance crop  
100 physiological performance. Therefore, this study aimed to evaluate the combined effects of gypsum with  
101 leguminous residue on changes in soil properties and how these changes could affect maize  
102 physiological performance. The ultimate objective is to contribute to family farm adaptation to global  
103 warming predicted to the Amazonian periphery.

## 104 **Materials and Methods**

### 105 *Study Site*

106

107 The field experiment was conducted in 2018 and 2019 at the Universidade Estadual do Maranhão, São  
108 Luís, Maranhão, Brazil (2°30'4" S, 44°18'33" W). The region's climate is hot and humid equatorial. Its  
109 rainy season extends from January to June and its dry season, with a pronounced water deficit, which  
110 lasts from July to December. The annual average temperature is approximately 27°C, the maximum  
111 temperature is 37°C, and the minimum temperature is 23°C. The total rainfall during the experimental  
112 periods of October to January of each year was 528.6 mm in 2018 and 53.6 mm in 2019 (Fig. 1). The



113 soil in the experimental area is an sandy loam Arenic Hapludult (Smith 2014) with hard-setting  
114 characteristics (Daniells 2012).

115

116 **Fig. 1** Precipitation, daily maximum, mean and minimum temperature and relative humidity at the study  
117 site, 2018–2019

### 118 *History of the experimental area*

119

120 The area was fallow since 1990 and supported a native species of grass, which was removed using  
121 glyphosate. In January 2011, lime was applied to the surface at a rate of 1 t ha<sup>-1</sup>. During this period,  
122 natural gypsum was applied at a rate of 6 t ha<sup>-1</sup> to plots designated for this treatment. Maize (cultivar  
123 AG 7088) was sown in 2011–2013 and 2015–2017. Gypsum was applied again in 2016 at a rate of 4 t  
124 ha<sup>-1</sup>. Residues from *Gliricidia sepium* (*Gliricidia*) was applied in the form of fresh branches at 12 t ha<sup>-1</sup>  
125 each year.

### 126 *Field trial and treatments*

127

128 The experiment was conducted under no-tillage mulch system, with maize planted and harvested in  
129 2018 and 2019. We used Pioneer 30f35, a commercial maize cultivar. Each experimental plot measured  
130 4 x 8 m, with a total of 28 plots in a randomized block arrangement. Plots consisted of 5 rows, spaced  
131 0.80 m apart with an intra-plant spacing of 0.25 m. The planting density was 50,000 plants ha<sup>-1</sup>.  
132 *Gliricidia* mulch was applied at a rate of 12 t ha<sup>-1</sup> per year. The study had four replicates of each of the  
133 following treatments: control (C), gypsum (G), nitrogen (N), legume residue (L), nitrogen + gypsum  
134 (NG), legume residue + gypsum (LG) and legume residue + nitrogen + gypsum (LNG). Chemical  
135 fertilizer was applied to maize at a rate of 100 kg ha<sup>-1</sup> of P in the form of triple superphosphate, 180 kg  
136 ha<sup>-1</sup> of K as potassium chloride, 5 kg ha<sup>-1</sup> of Zn in the form of zinc sulphate and 180 kg ha<sup>-1</sup> of N in the  
137 form of urea. Phosphorus and zinc were applied as basal fertilizer once before planting. Nitrogen (only  
138 in the plots that received nitrogen) and potassium were applied at a rate of 60 kg ha<sup>-1</sup> at planting, 60 kg  
139 ha<sup>-1</sup> at the V6 stage (six leaves with collars visible) and 60 kg ha<sup>-1</sup> at the V11 stage (11 leaves with collars  
140 visible). The plots used a rainfed system in 2018 and a restricted irrigation regime in 2019. In 2019,  
141 water was supplied by drip tape irrigation with one tape per row with emitters spaced 25 cm apart  
142 delivering 1.25 L h<sup>-1</sup> for 4 h, resulting in a total of 20 mm of water applied per irrigation.

## 143 **Measured Parameters and Analyses**

### 144 *Soil Samples*

145  
146 We collected soil samples for chemical analysis at the end of the second-year experiment at depths of  
147 0–20 and 20–40 cm. A total of four simple samples per plot were collected using a Dutch auger with a  
148 3 inch diameter. Samples from each plot were passed through a 2 mm screen mesh and air-dried before  
149 the analyses. Each sample was analysed using an exchangeable ion resin (Raij et al. 1986) as an extractor  
150 for Ca, Mg, and K, which were measured using a Varian 720-ES ICP Optical Emission Matter Analysis  
151 Spectrometer (Walnut Creek, CA, USA).

### 152 *Soil penetration resistance (SPR) and soil moisture*

153  
154 The SPR measurements were collected at depths of 0–0.10, 0.10–0.20, 0.20–0.30, and 0.30–0.40m with  
155 three replicates per plot. In 2019, these measures were taken 4 days after irrigation (or 4 days without  
156 water) and immediately before the irrigation (or 7 days without water). SPR was measured using a digital  
157 penetrometer (Falker, Porto Alegre, Brazil) with 1 cm gradations. The bars of critical level defined by  
158 Hazelton and Murphy (2016) was used to construct the graph of the soil penetration strength. We  
159 calculated soil moisture content using the gravimetric method with the following formula: soil moisture  
160 (g/g) = (weight of fresh soil – weight of oven-dried soil)/weight of oven-dried soil.

### 161 *Physical fractionation of soil organic matter*

162  
163 The soil organic matter in the 0–20 cm layer of the soil profile was physically fractionated following  
164 Cambardella and Elliott (1992). After drying, a soil sample was ground in a porcelain mortar, after  
165 which an aliquot was collected, weighed, and analysed for its C content, which represented the soil  
166 particulate organic carbon (POC) in the particulate organic matter, according to the Walkley and Black  
167 (1934) method. An aliquot of the 2 mm sieved subsample was ground in a porcelain mortar and weighed  
168 and analysed for total organic carbon (TOC). Soil mineral-associated carbon (MOC) was calculated as  
169 the difference between TOC and POC. The total organic carbon stock (TOCS) of each of the 0–30 cm  
170 layers was calculated with the following expression (Veldkamp 1994): C stock = (TOC x  $\rho_s$  x E) /100,  
171 where C stock = organic C stock at a given depth ( $t\ ha^{-1}$ ); OC = organic C content at the sampled depth  
172 ( $g\ kg^{-1}$ );  $\rho_s$  = soil bulk density ( $kg\ dm^{-3}$ ); and E = thickness of the layer (30 cm).

173

174

### 175 ***Plant and nitrogen analysis***

176

177 Total N content were measured in 2018 and 2019 at the physiological maturity stage. Five plants from  
178 each plot were randomly selected. All plant material was dried at 60°C for 6 days to a constant weight  
179 in an oven and was then ground for analysis. Total nitrogen content in each sample was determined from  
180 the mass of dried plant matter by digestion in sulfuric acid (H<sub>2</sub>SO<sub>4</sub> – H<sub>2</sub>O<sub>2</sub>), according to the standard  
181 distillation method described by Cottenie (1980).

### 182 ***Gas exchanges***

183

184 Gas exchange measurements were performed using a portable LI-6400XT open gas exchange system  
185 (Li-Cor Inc., Lincoln, NE, USA) that included a 6400-02B LED light source. These two accessories  
186 were used to maintain a constant CO<sub>2</sub> concentration of 400 ± 10 μmol mol<sup>-1</sup> and a constant incident  
187 photosynthetic active radiation (PAR) of 1500 μmol m<sup>2</sup> s<sup>-1</sup> in the leaf chamber during instantaneous  
188 measurements. A flow rate to chamber was 500 μmol s<sup>-1</sup>. Net photosynthetic rate ( $P_N$ ), stomatal  
189 conductance ( $g_s$ ) and transpiration (E) were obtained from the instantaneous measurements. The  
190 measurements were taken on three randomly selected and marked plants on the middle rows of each  
191 plot in 2018 and 2019, between 7:30 am and 9:30 am at the V6 (six leaves with collars visible) and VT  
192 (tasseling) stages on the third fully matured leaf from the top. In 2018, the first measurement was done  
193 6 days after rainfall and the second measure was performed 1 day after rainfall. In 2019 the first and  
194 second measures were done 6 days after the irrigation (or 6 days without water). The instantaneous water  
195 use efficiency (WUE) at leaf level was calculated as the ratio between photosynthesis and transpiration  
196 (instantaneous WUE =  $P_N / E$ ).

### 197 ***Fluorescence measurements***

198

199 The performance index ( $PI_{abs}$ ) was measured at the same time and on the same leaves as the gas exchange  
200 measurements. Maize leaves were adapted to darkness for 30 minutes using a light-with-holding clips.  
201 Chlorophyll *a* fluorescence was measured with a portable non-modulated fluorimeter Pocket-PEA  
202 (*Plant Efficiency Analyser* - Hansatech, Norfolk, UK). After the adaptation of leaves to darkness, a  
203 single strong light pulse lasting one second (3500 μmol m<sup>2</sup> s<sup>-1</sup>) was applied using three light-emitting  
204 diodes (650 nm). The measurements were taken on three randomly selected and marked plants in the  
205 middle rows of each plot.

### 206 **Statistical Analysis**

207

208 An analysis of variance (ANOVA) was performed on the data to evaluate treatment effects and means  
 209 were compared using the Tukey's HSD test ( $p < 0.05$ ) using R program. The data from the each year  
 210 were analysed separately because of significant ( $p < 0.05$ ) season year interactions.

## 211 **Results**

212

### 213 **Changes in SOC and SBC**

214

215 Total organic carbon (TOC) was higher in the N (11%), G (13%) and L (60%) treatments compare to  
 216 the control, while the LNG treatment was 81% higher (Table 1). These results were primarily driven by  
 217 mineral associated carbon (MOC), whose values and differences between treatments were higher than  
 218 particulate organic carbon (POC).

219

220 **Table 1** Soil organic carbon (SOC) fractions: particulate organic carbon (POC), mineral associated-  
 221 organic carbon (MOC) and total organic carbon (TOC). Notes: Values followed by different letters in  
 222 the same line indicate a significant difference at the 5% level by the Tukey's HSD test. C: control; N:  
 223 nitrogen; G: gypsum; NG: nitrogen+gypsum; LG: legume residue+gypsum; L: legume residue; LNG:  
 224 legume residue+nitrogen+gypsum

225

226 Calcium (Ca) content was also increased by legume and gypsum in the soil profile. In the 0–20 cm layer,  
 227 the effect of the legume to calcium content was greater than the gypsum effect (Table 2). There was no  
 228 significant difference between the L, NG, LG, and LNG treatments, though they all had higher calcium  
 229 levels than the other treatments. Notably, calcium levels were higher in plots with only legume mulch  
 230 than in plots with only gypsum ( $L > G$ ). Similar results were obtained for the sum of base cations (SBC).  
 231 In contrast, magnesium (Mg) content was lowest in the G and NG treatments, in which Mg content was  
 232 three times lower than in the L plots. Potassium (K) content was higher in L and LNG than in N and  
 233 NG.

234 In contrast, gypsum was more effective in increasing Ca levels in the 20–40cm layer than legume  
 235 residue, which was reflected in higher overall Ca levels in treatments with gypsum. The increase in  
 236 SBC was also greater in gypsum treatments than in those with legume residue. Indeed, SBC was lower  
 237 in the L plots than in treatments with gypsum, though not as low as the C and N treatments.

238

239 **Table 2** Exchange base cations ( $\text{mmol}_c \text{ m}^{-3}$ ) and sum of base cation (SBC). Notes: Values followed by  
 240 different letters in the same line indicate a significant difference at the 5% level by the Tukey's HSD  
 241 test. Notes: C: control; N: nitrogen; G: gypsum; NG: nitrogen+gypsum; LG: legume residue+gypsum;  
 242 L: legume residue; LNG: legume residue+nitrogen+gypsum

243

244 Legume residue application positively affected soil moisture, with higher levels than the control in the  
 245 0–20 cm layer, even after 7 days without rain (Table 3). In this layer, only treatments with residue

246 maintained soil moisture above the critical level of 5%. On the whole, soil moisture in the 20–40 cm  
 247 layer was higher in the L and LNG treatments than in treatments without legume residue.

248

249 **Table 3** Gravimetric soil moisture (g/g) 7 days without water in 2019. Notes: Values followed by  
 250 different letters in the same line indicate a significant difference at the 5% level by the Tukey's HSD  
 251 test. C: control; N: nitrogen; G: gypsum; NG: nitrogen+gypsum; LG: legume residue+gypsum; L:  
 252 legume residue; LNG: legume residue+nitrogen+gypsum

253

254 Our results showed that impact of water scarcity on treatments without legume residue was greater than  
 255 in treatments with mulch cover. This can be seen in the penetration resistance curves on the 4<sup>th</sup> and 7<sup>th</sup>  
 256 days after irrigation. Figure (2a) and (2b) provide insights on the intensity (distance between curves of  
 257 the same treatment) and duration of water stress (time spent in the stress range). After four days without  
 258 irrigation, soil penetration resistance was under 1 MPa up to 20 cm and lower than 2 MPa up to 30 cm  
 259 in treatments with legume residues (Fig. 2a). Meanwhile, SPR was higher than 1 MPa in the 10–20 cm  
 260 layer (Fig. 2b) in treatments without legume mulch. After seven days without irrigation, SPR was higher  
 261 than critical level of 2.0 MPa in the 0–30 cm layer in the LG and L treatments, as well as all treatments  
 262 without leguminous mulch. Only in the LNG treatment SPR was lower than 1 MPa in the 0–20cm layer  
 263 and lower than 2 MPa in the 0–30 cm layer.

264

265 **Fig. 2** Response of different treatments with legume residue (a) and without legume residue (b) in soil  
 266 penetration resistance (MPa). Notes: C: control; G: gypsum; N: nitrogen; L: legume residue; NG:  
 267 nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum; 4: four days  
 268 without water; 7: seven days without water

269

## 270 **Maize physiological response to soil improvement**

271

272 There were no significant differences in stomatal conductance ( $g_s$ ) in 2018 at the VT stage, though  $g_s$   
 273 was higher in LG ( $0.53 \text{ mol m}^{-2} \text{ s}^{-1}$ ) than in G ( $0.34 \text{ mol m}^{-2} \text{ s}^{-1}$ ) at the V6 stage (Fig. 3a). In 2019, the  
 274 LNG treatment had a higher  $g_s$  at the V6 stage than all other treatments, with no difference between the  
 275 remaining treatments (Fig. 3b). At VT stage, however, the NG, LG, and LNG treatments were greater  
 276 than the control. There was no difference between the other treatments.

277 The net photosynthetic rate ( $P_N$ ) was also different in each year of the experiment. In 2018,  $P_N$  ranged  
 278 from 38.72 to 52.17  $\mu\text{mol m}^{-2} \text{ s}^{-1}$ , while the range was larger in 2019, varying between 13.45 and 39.45  
 279  $\mu\text{mol m}^{-2} \text{ s}^{-1}$ . There was no difference between treatments at the VT stage in 2018, while G ( $38.72 \mu\text{mol}$   
 280  $\text{m}^{-2} \text{ s}^{-1}$ ) had lower  $P_N$  than LG ( $44.64 \mu\text{mol m}^{-2} \text{ s}^{-1}$ ) at the V6 stage (Fig. 3c). In contrast, in 2019 LNG  
 281 had the highest photosynthetic rate at the V6 stage, with no difference among the others treatments,  
 282 except for LG ( $36.16 \mu\text{mol m}^{-2} \text{ s}^{-1}$ ) being higher than L ( $31.49 \mu\text{mol m}^{-2} \text{ s}^{-1}$ ) (Fig. 3d). At the VT stage,  
 283 all treatments with added urea had a higher  $P_N$  than the remaining treatments and L had a higher  $P_N$  than  
 284 C. All of the treatments had low  $P_N$  values at the VT stage (with means under  $23 \mu\text{mol m}^{-2} \text{ s}^{-1}$ ).

285 **Fig. 3** Stomatal conductance,  $g_s$  ( $\text{mol m}^{-2} \text{s}^{-1}$ ) in 2018 (a) and in 2019 (b) and net photosynthetic rate,  $P_N$   
 286 ( $\mu\text{mol m}^{-2} \text{s}^{-1}$ ) in 2018 (c) and 2019 (d). Notes: Different lower-case letters at the V6 stage and different  
 287 capital letters at the VT stage in each graph indicate significant differences at the 5% level per Tukey's  
 288 HSD test. Notes: C: control; G: gypsum; N: nitrogen; L: legume residue; NG: nitrogen+gypsum; LG:  
 289 legume residue+gypsum; LNG: legume residue+nitrogen+gypsum

291 There was large variation in the performance index ( $PI_{\text{abs}}$ ) between 2018 and 2019. In 2018 at V6 stage,  
 292  $PI$  was higher in LNG than in C, G, and N, while at the VT stage,  $PI$  was higher in NG and LNG than  
 293 in C and G (Fig. 4a). In 2019, there were no significant differences among treatments at the V6 stage,  
 294 but at VT the NG and LNG treatments were had higher  $PI$  values than the other treatments (Fig. 4b).

295 **Fig. 4** Performance Index ( $PI_{\text{abs}}$ ) in 2018 (a) and 2019 (b). Notes: Different lower-case letters at the V6  
 296 stage and different capital letters at the VT stage in each graph indicate significant differences at the 5%  
 297 level per Tukey's HSD test. Notes: C: control; G: gypsum; N: nitrogen; L: legume residue; NG:  
 298 nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum

300 As for instantaneous water use efficiency (A/E) there was no difference between treatments at the V6  
 301 stage in 2018 and 2019 (Fig. 5a). However, at VT stage in 2018 instantaneous water use efficiency was  
 302 higher in LNG than in N, G and C. In 2019 it was higher in the treatments LNG, NG and N than in all  
 303 other treatments (Fig. 5b).

304 **Fig. 5** Instantaneous leaf water use efficiency (Ins WUE) in 2018 (a) and 2019 (b). Notes: Different  
 305 lower-case letters at the V6 stage and different capital letters at the VT stage in each graph indicate  
 306 significant differences at the 5% level per Tukey's HSD test. Notes: C: control; G: gypsum; N: nitrogen;  
 307 L: legume residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume  
 308 residue+nitrogen+gypsum

### 311 Nitrogen uptaken and maize grain yield (GY)

312  
 313 Legume residue affected total N uptake when used alone, though only in the rainfed system (2018), in  
 314 which the LG and L treatments had higher N uptake than C and G (Fig. 6a). The LNG and NG treatments  
 315 had higher N uptake than in the other treatments, while total N was not different from the LG and L  
 316 treatments than in plots with only urea (N treatment). In 2019, all treatments with added urea (N, NG,  
 317 and LNG) had considerably higher nitrogen uptake than other treatments, with no difference between  
 318 the remaining treatments. In 2018, GY was the lowest in the G, C, and N treatments (Fig. 6b). There  
 319 were no significant differences between the L, NG and LG treatments, while LNG had the highest GY  
 320 of any treatment. In 2019, added urea had the greatest effect on GY. In the N treatment, GY was up to  
 321 78% higher than in treatments without added urea. The LNG treatment had the highest yield, which was  
 322 151% greater than the control and 41% higher than the N treatment.

323  
 324 **Fig. 6** Effects of different treatments on total N ( $\text{g kg}^{-1}$ ) in 2018 (=1) and 2019 (=2) (a) and maize grain  
 325 yield (GY) in 2018 (1) and 2019 (2). Notes: Different letters in each graph indicate significant

326 differences at the 5% level per the Tukey's HSD test. C: control; G: gypsum; N: nitrogen; L: legume  
327 residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum

## 328 **Discussion**

329

### 330 **Soil improvement and nitrogen uptake**

331

332 Our results showed that interactions between legume residue, gypsum and N application can improve  
333 the soil root zone, reducing the effect of drought conditions on physiological parameters in maize while  
334 also increasing nitrogen uptake. The observed soil improvements can be explained through specific  
335 interactions between polyvalent cations and compounds derived from the decomposition of legume  
336 residues (Ellerbrock and Gerke 2018). In turn, synthetic N inputs increase microbial activity and thereby  
337 increase soil organic matter transformation and microbial biomass if N is a soil limiting resource (Chen  
338 et al. 2014). Gypsum and N application appear to have complementary effects that help capture the  
339 benefits of leguminous mulch application to ensure an adequate content of soil organic carbon. The  
340 predominance of mineral-associated organic carbon in our samples may be due to the fast decomposition  
341 of POC the more labile SOC fraction (Hijbeek et al. 2018).

342 Treatments with mulch had higher moisture levels and lower SPR. Straw mulches are commonly used  
343 because of their ability to reduce soil evaporation by preventing capillarity (Jimenez et al. 2017).  
344 Furthermore, the increase in water holding capacity may be directly related to increased soil organic  
345 carbon observed in treatments with legume mulch. The presence of SOM plays a key role in altering the  
346 pore size distribution of sandy loam soils, as a result of decreased bulk density (Robin et al. 2018). In  
347 turn, the increased number of small pores and higher specific surface area increases water holding  
348 capacity at low tensions such as field capacity, increasing water availability for plants (Moura et al.  
349 2009).

350 In addition to the positive effect of increased soil moisture, decreased impedance is due to a dilution  
351 effect caused by mixing of the biomass with the denser mineral fraction of the soil (Maltas et al. 2018).  
352 In soil prone to cohesion, increased SOM decreases SPR indirectly by affecting the stability  
353 of soil aggregates (Tarkiewicz and Nosalewicz 2005). Added calcium has both direct (increasing  
354 flocculation and aggregation in the subsoil) and indirect (improving root activity, which leads to greater  
355 soil aggregation) effects on SPR (Sumner et al. 2009). Thus, the interaction between legume residues  
356 and added calcium can increase SOM, soil moisture and SBC while decreasing SPR. These effects  
357 encourage plant growth while increasing the availability of water in drought scenarios  
358 (Lynch and Wojciechowski 2015).

359

### 360 **Maize physiological response to soil improvement**

361 The changes in maize physiological parameters observed in this experiment are driven principally by  
362 variations in SPR. According to Ley et al. (1995) in soil that tend to harden, effects on root growth are  
363 common when the water potential approaches -100 kPa. In these circumstances, SPR is a better proxy  
364 for plant water stress rather than the permanent wilting point, as proposed by Letey (1985). Therefore,  
365 the decreased intensity and duration of water stress in the LNG treatment, as shown in Figure 2a, can  
366 account for its improved performance in all physiological parameters evaluated in this experiment. In  
367 this experiment, the LNG treatment resulted in higher stomatal conductance and net photosynthetic rate  
368 in both years, but the effect was greatest in 2019, the year in which drip irrigation was used. Soil  
369 improvement as reflected by lower penetration resistance and higher soil moisture can partially account  
370 for these results.

371 Environmental stress results in lower stomatal conductance and net photosynthetic rate, as shown in  
372 Figure 3, which is reflected in lower growth (Ripley et al. 2007). Under both soil and atmosphere water  
373 stress, this decrease in stomatal conductance may be attributed to limited root system capacity to meet  
374 leaf water demand (Lopes et al. 2019; Brodribb et al. 2020). The combined effect of gypsum and legume  
375 residue to decrease root stress in LG and LNG can explain higher  $g_s$  and  $P_N$  values in these treatments.  
376 Indeed, the reduction in photosynthetic electron consumption, an important consequence of the drought-  
377 induced decrease in  $CO_2$  supply, causes a decrease in photosynthetic efficiency and an increased  
378 requirement for additional electron sinks such as photorespiration (Ripley et al. 2017).

379 Differences in performance index ( $PI_{abs}$ ) shown in Figure 4 confirm the effectiveness of the LNG  
380 treatment to reduce stress in maize plants.  $PI_{abs}$  reflects the functionality of both photosystems I and II  
381 and gives us quantitative information on the current state of plant performance under stress conditions  
382 (Živčák, et al. 2008). Therefore, the  $PI_{abs}$  provides a proxy for plant vitality as well as indicating stress  
383 in plants even before visible symptoms appear on the leaves (Christen et al. 2007). Indeed, even in 2018,  
384  $PI_{abs}$  was able to detect higher vitality in LNG treatment at the earlier V6 stage. Again, lower penetration  
385 resistance, higher soil moisture and higher nitrogen contents in leaves can explain higher  $PI_{abs}$  due to  
386 lower stress in plants in LNG plots.

387 Differences in instantaneous water user efficiency were just verified in water restrict regime, when it  
388 was higher in treatments with N, demonstrating that maize performance could be increased with no  
389 change in water use rate. Water shortage can lead to N deficiency and a decline in transpiration  
390 efficiency strictly proportional to the N deficiency induced by drought. Therefore, interaction between  
391 water and N deficit could be analysed by the N uptake/transpiration ratio indicator (Kunrath et al. 2018).

392

### 393 **Nitrogen uptake and grain yield (GY)**

394

395 Due to differences in the rainfall regime, harnessing of N from urea and legume residue was more  
396 dependent on the water supply regime than the soil structure. This can be seen in the higher uptake of N  
397 from decomposing legume residue in 2018 but not in 2019, when legume residues had no effect on N



398 uptake. In contrast, under intense rain regime the effect of urea on N uptake decreased in 2018 and was  
399 enhanced in 2019 under irrigation regime. The LNG treatment showed the highest N content in both  
400 years, which is consistent with findings by Senna et al. (2020).

401 Our results showed that maize production can be increased by soil improvements resulting from legume  
402 residue and gypsum in both water regimes. Since legume residue contributed relatively little to N uptake  
403 in the restricted irrigation regime, the 35% difference in grain yield between the N and LNG treatments  
404 in 2019 is due mainly to soil improvement caused by soil management (Moura et al. 2020). This result  
405 confirms that the combination of legume residues with gypsum and urea application can be a good  
406 strategy to maintain crop productivity in cohesive soil and drought conditions.

## 407 **Conclusions**

408

409 Our results showed the possibility of improving crop management in conditions of cohesive tropical soil  
410 with an irregular water regime. These improvements were primarily apparent in certain key areas,  
411 including soil organic matter, water holding capacity, penetration resistance and calcium content. This  
412 underscores the importance of soil management for the adaptation of tropical agroecosystems to climate  
413 change. The improvements in soil attributes resulted in positive effects in physiological parameters, with  
414 a reduction in plant stress reflected mainly in the performance index ( $PI_{abs}$ ). These changes resulted in  
415 increased biomass and maize grain yield, even in a restricted irrigation regime. However, it is worth  
416 highlighting the significant contribution of added nitrogen to decreasing plant stress, increasing soil  
417 organic carbon, enhancing physiological parameters, and increasing biomass production. Therefore, the  
418 inclusion of nitrogen alongside gypsum and legume residue should be part of new research on climate  
419 change adaptation strategies for tropical agriculture.

420

421 **Conflict of Interest:** The authors declare that they have no conflict of interest.

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Ecophysiological response of maize to soil management in the Amazonian periphery in the context of climate change adaptation

Regional Environmental Change

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**Table 1.** Soil organic carbon (SOC) fractions: particulate organic carbon (POC), mineral associated-organic carbon (MOC) and total organic carbon (TOC).

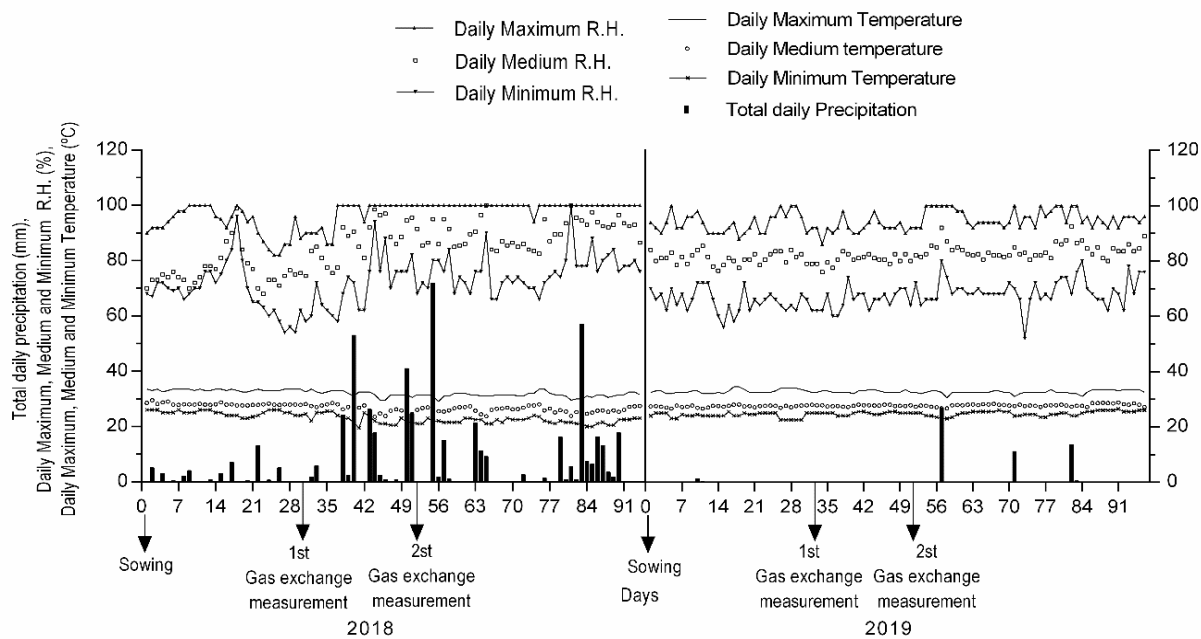
SOC fractions (g/kg)	Treatments						
	C	N	G	NG	LG	L	LNG
POC	1.13 b	1.55 ab	1.83 a	1.86 a	1.71 ab	1.81 a	2.00 a
MOC	5.76 b	6.13 b	5.64 b	6.94 b	9.23 a	9.33 a	10.62 a
TOC	6.91 e	7.68 d	7.80 d	8.76 c	10.99 b	11.09 b	12.53 a

**Table 2.** Exchange base cations ( $\text{mmol}_c \text{m}^{-3}$ ) and sum of base cation (SBC).

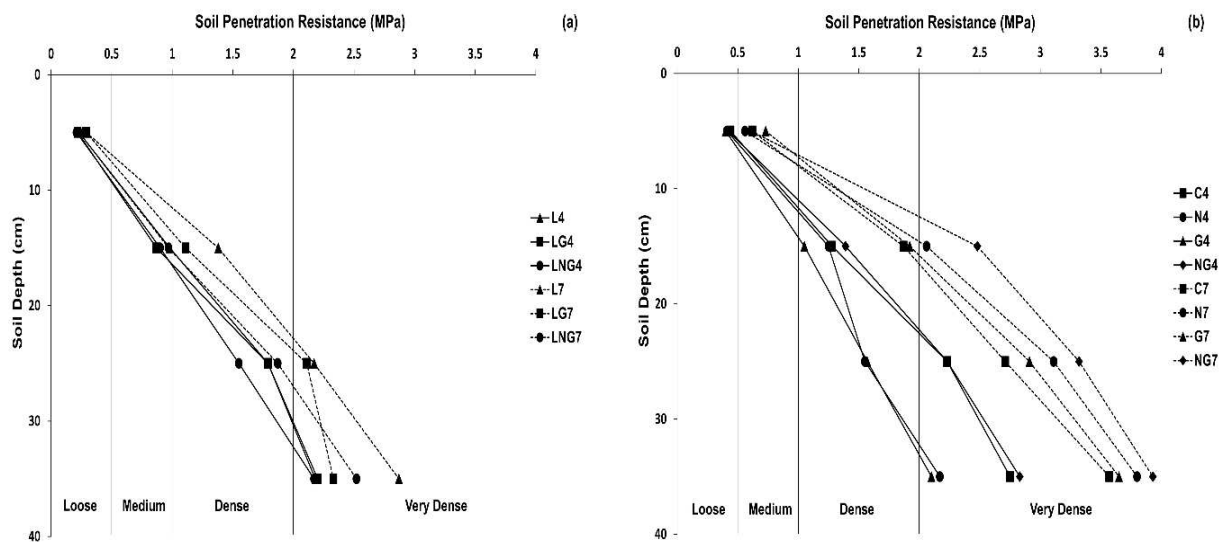
Cations content ( $\text{mmol}_c \text{m}^{-3}$ )	Treatments						
	C	N	G	L	NG	LG	LNG
	0 - 20 cm						
Ca	5.46 c	5.51 c	9.75 b	11.92 a	13.13 a	12.73 a	14.49 a
Mg	2.22 bc	1.46 c	1.08 d	3.59 a	1.01 d	2.41 b	2.16 bc
K	0.44 ab	0.37 b	0.42 ab	0.48 a	0.37 b	0.44 ab	0.46 a
SBC	8.25 c	7.34 c	11.24 b	16.00 a	14.50 a	15.57 a	17.10 a
	20 - 40 cm						
Ca	2.83 c	2.75 c	8.27 a	4.67 b	8.47 a	7.82 a	7.83 a
Mg	0.86 b	0.79 bc	0.54 bc	1.45 a	0.48 c	0.86 b	0.76 bc
K	0.53 a	0.32 c	0.60 a	0.50 ab	0.37 b	0.50 ab	0.35 bc
SBC	4.62 c	3.85 c	9.74 ab	6.62 b	9.86 a	9.57 ab	9.34 ab

**Table 3.** Gravimetric soil moisture (g/g) 7 days without water in 2019.

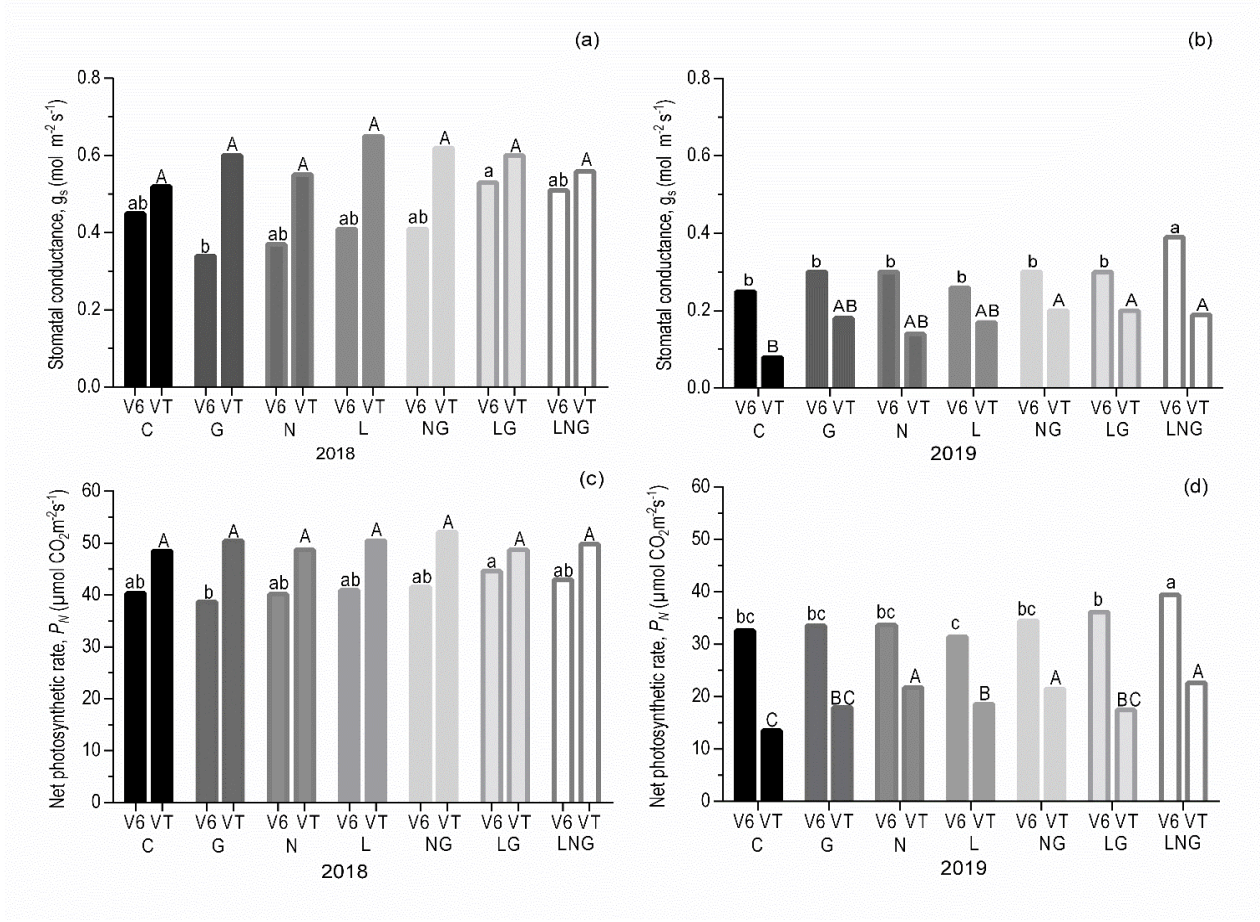
Layer (cm)	Treatments						
	C	N	G	L	NG	LG	LNG
0 - 10	0.022 b	0.021 b	0.023 b	0.052 a	0.026 b	0.051 a	0.044 a
11 - 20	0.024 c	0.026 c	0.041 bc	0.056 ab	0.042 b	0.059 ab	0.069 a
21 - 30	0.052 ab	0.041 b	0.068 ab	0.098 a	0.051 ab	0.057 ab	0.087 a
31 - 40	0.051 bc	0.043 c	0.040 c	0.089 a	0.045 c	0.073 b	0.086 a



**Fig. 1** Precipitation, daily maximum, mean and minimum temperature and relative humidity at the study site, 2018–2019

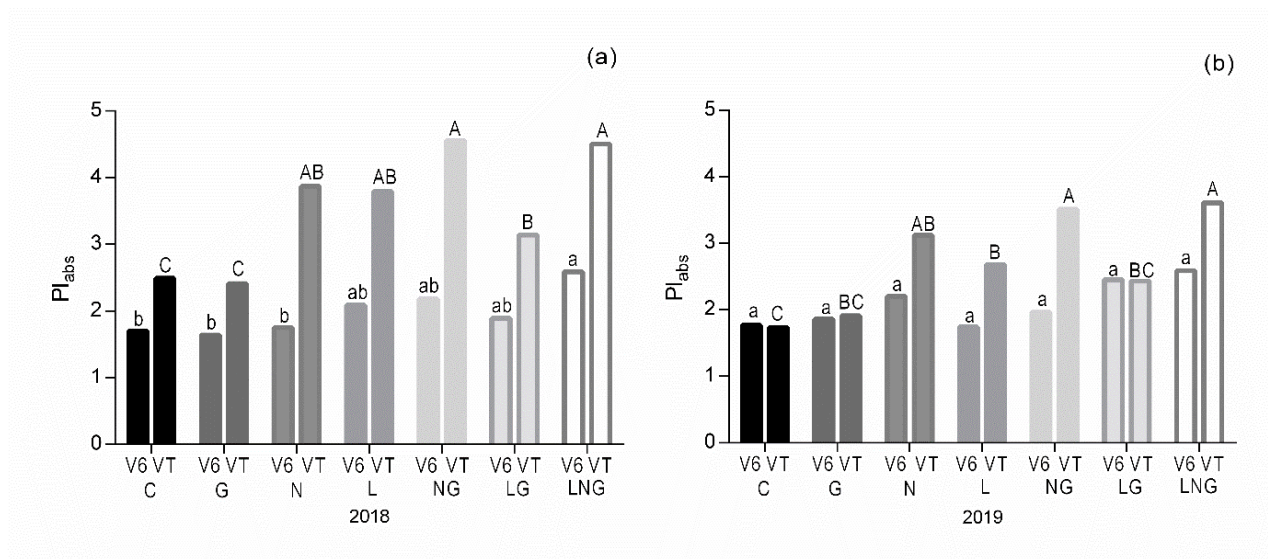


**Fig. 2** Response of different treatments with legume residue (a) and without legume residue (b) in soil penetration resistance (MPa). C: control; G: gypsum; N: nitrogen; L: legume residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum; 4: four days without water; 7: seven days without water. Vertical bars mean the critical levels by Hazelton & Murphy (2007)

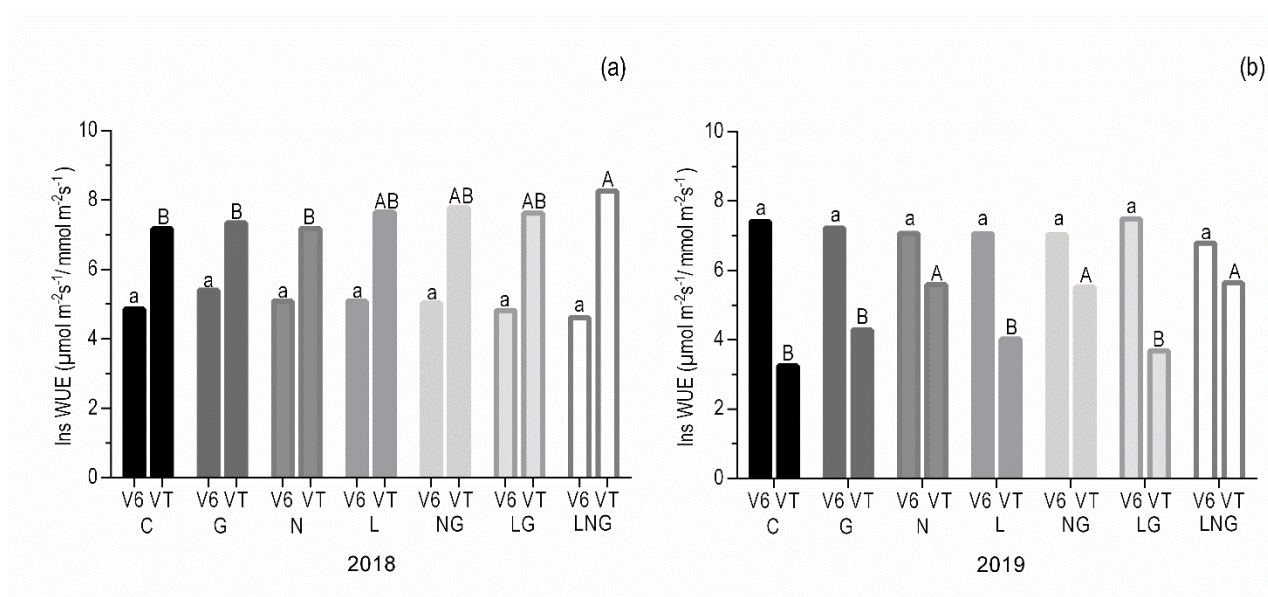


**Fig. 3** Stomatal conductance,  $g_s$  ( $\text{mol m}^{-2} \text{s}^{-1}$ ) in 2018 (a) and in 2019 (b) and net photosynthetic rate,  $P_N$  ( $\mu\text{mol m}^{-2} \text{s}^{-1}$ ) in 2018 (c) and 2019 (d). Notes: Different lower-case letters at the V6 stage and different capital letters at the VT stage in each graph indicate significant differences at the 5% level per Tukey's HSD test. C: control; G: gypsum; N: nitrogen; L: legume residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum

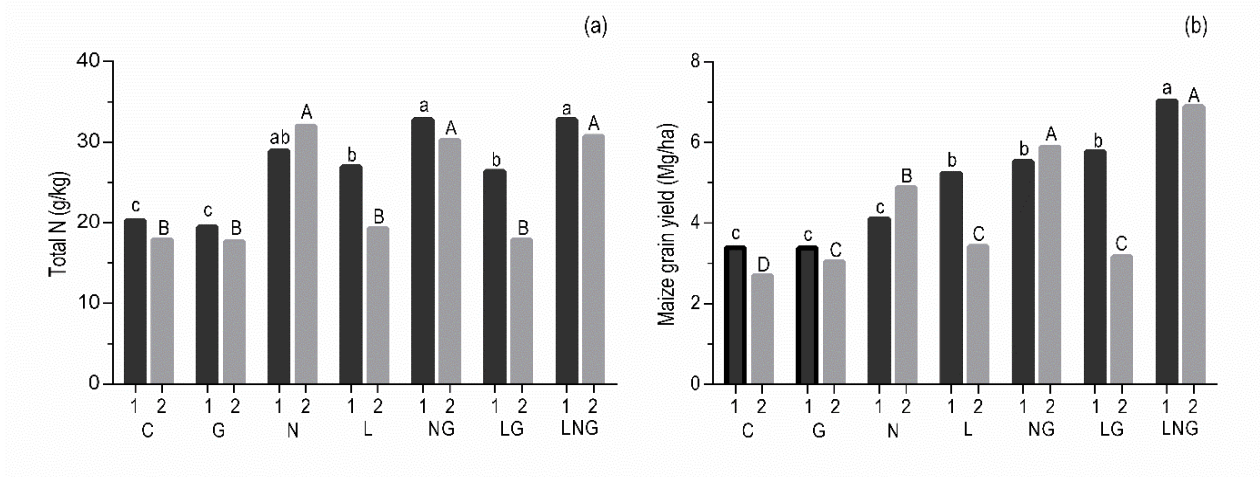




**Fig. 4** Performance Index ( $PI_{abs}$ ) in 2018 (a) and 2019 (b). Notes: Different lower-case letters at the V6 stage and different capital letters at the VT stage in each graph indicate significant differences at the 5% level per Tukey's HSD test. C: control; G: gypsum; N: nitrogen; L: legume residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum



**Fig. 5** Instantaneous leaf water use efficiency (Ins WUE) in 2018 (a) and 2019 (b). Notes: Different lower-case letters at the V6 stage and different capital letters at the VT stage in each graph indicate significant differences at the 5% level per Tukey's HSD test. C: control; G: gypsum; N: nitrogen; L: legume residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum



**Fig. 6** Effects of different treatments on total N ( $\text{g kg}^{-1}$ ) in 2018 (=1) and 2019 (=2) (a) and maize grain yield (GY) in 2018 (1) and 2019 (2). Notes: Different letters in each graph indicate significant differences at the 5% level per the Tukey's HSD test. C: control; G: gypsum; N: nitrogen; L: legume residue; NG: nitrogen+gypsum; LG: legume residue+gypsum; LNG: legume residue+nitrogen+gypsum

# **ANEXO**

## Springer Journal – Regional Environmental Change

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<http://physicsweb.org/articles/news/11/6/16/1>. Accessed 26 June 2007

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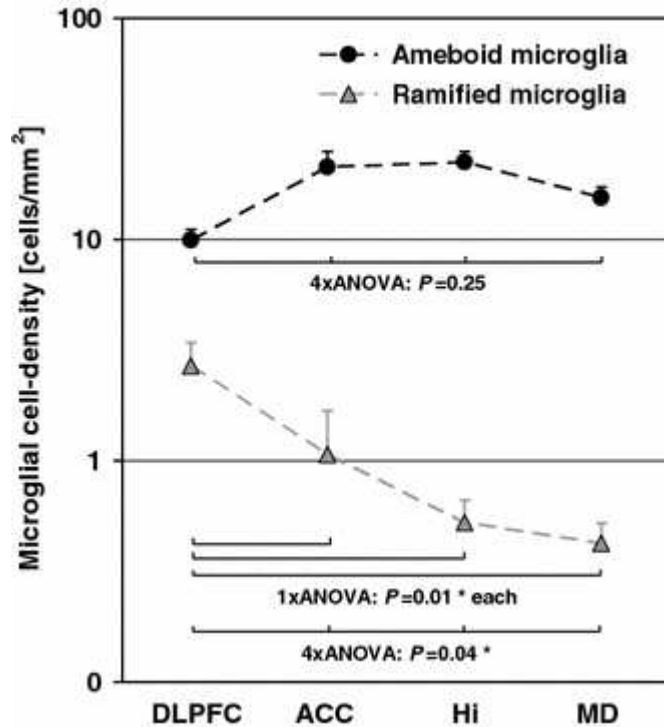
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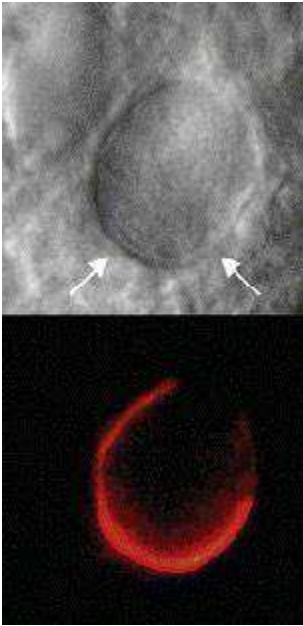
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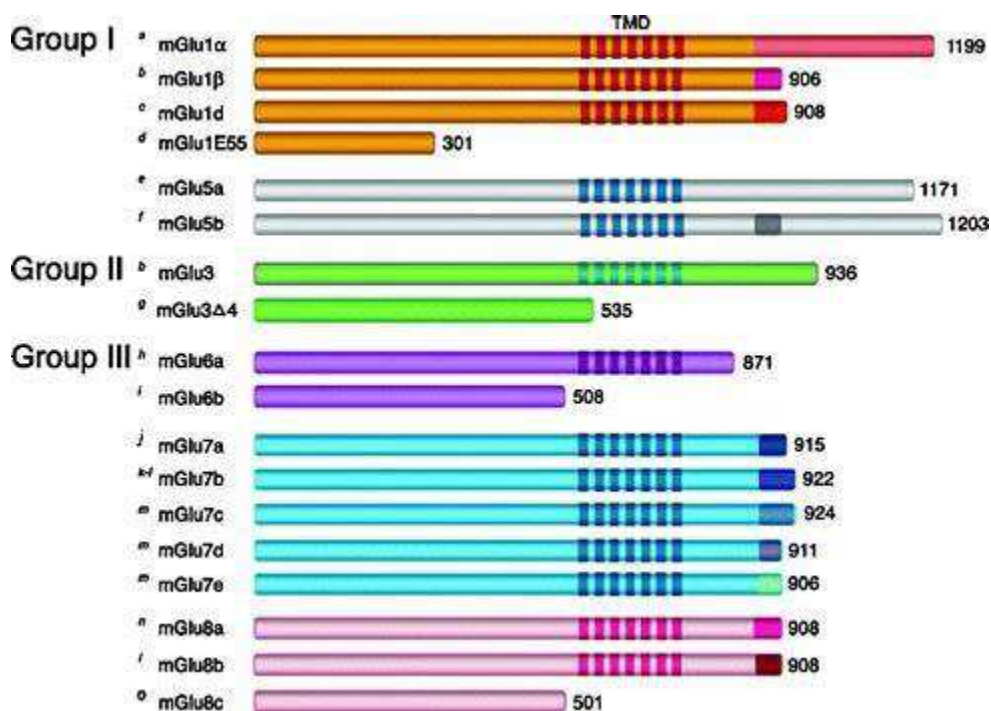


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