



## Impacts of ground-level ozone on sugarcane production

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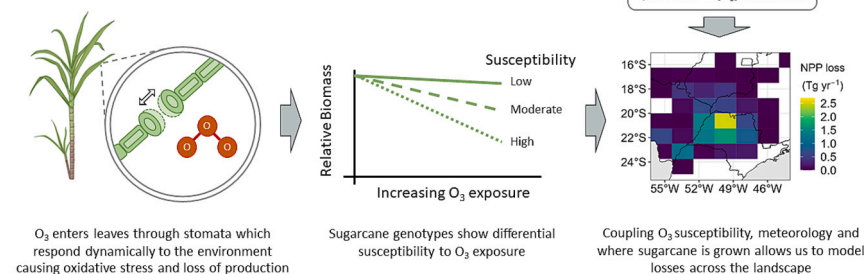
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### HIGHLIGHTS

- Sugarcane is a commodity crop grown across the (sub)tropics.
- Ozone exposure causes a reduction in productivity of sugarcane.
- Cultivars of sugarcane show different sensitivities to ozone.
- Spatial modelling shows variation in the risks of O<sub>3</sub> across south-central Brazil.
- Ozone poses a substantial production risk for the world's largest regional producer.

### GRAPHICAL ABSTRACT

#### Impacts of ozone (O<sub>3</sub>) on sugarcane production



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### ABSTRACT

Sugarcane is a vital commodity crop often grown in (sub)tropical regions which have been experiencing a recent deterioration in air quality. Unlike for other commodity crops, the risk of air pollution, specifically ozone (O<sub>3</sub>), to this C<sub>4</sub> crop has not yet been quantified. Yet, recent work has highlighted both the potential risks of O<sub>3</sub> to C<sub>4</sub> bioenergy crops, and the emergence of O<sub>3</sub> exposure across the tropics as a vital factor determining global food security. Given the large extent, and planned expansion of sugarcane production in places like Brazil to meet global demand for biofuels, there is a pressing need to characterize the risk of O<sub>3</sub> to the industry. In this study, we sought to a) derive sugarcane O<sub>3</sub> dose-response functions across a range of realistic O<sub>3</sub> exposure and b) model the implications of this across a globally important production area. We found a significant impact of O<sub>3</sub> on biomass allocation (especially to leaves) and production across a range of sugarcane genotypes, including two

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commercially relevant varieties (e.g. CTC4, Q240). Using these data, we calculated dose-response functions for sugarcane and combined them with hourly O<sub>3</sub> exposure across south-central Brazil derived from the UK Earth System Model (UKESM1) to simulate the current regional impact of O<sub>3</sub> on sugarcane production using a dynamic global vegetation model (JULES vn 5.6). We found that between 5.6 % and 18.3 % of total crop productivity is likely lost across the region due to the direct impacts of current O<sub>3</sub> exposure. However, impacts depended critically on the substantial differences in O<sub>3</sub> susceptibility observed among sugarcane genotypes and how these were implemented in the model. Our work highlights not only the urgent need to fully elucidate the impacts of O<sub>3</sub> in this important bioenergetic crop, but the potential implications air quality may have upon tropical food production more generally.

## 1. Introduction

### 1.1. Sugarcane

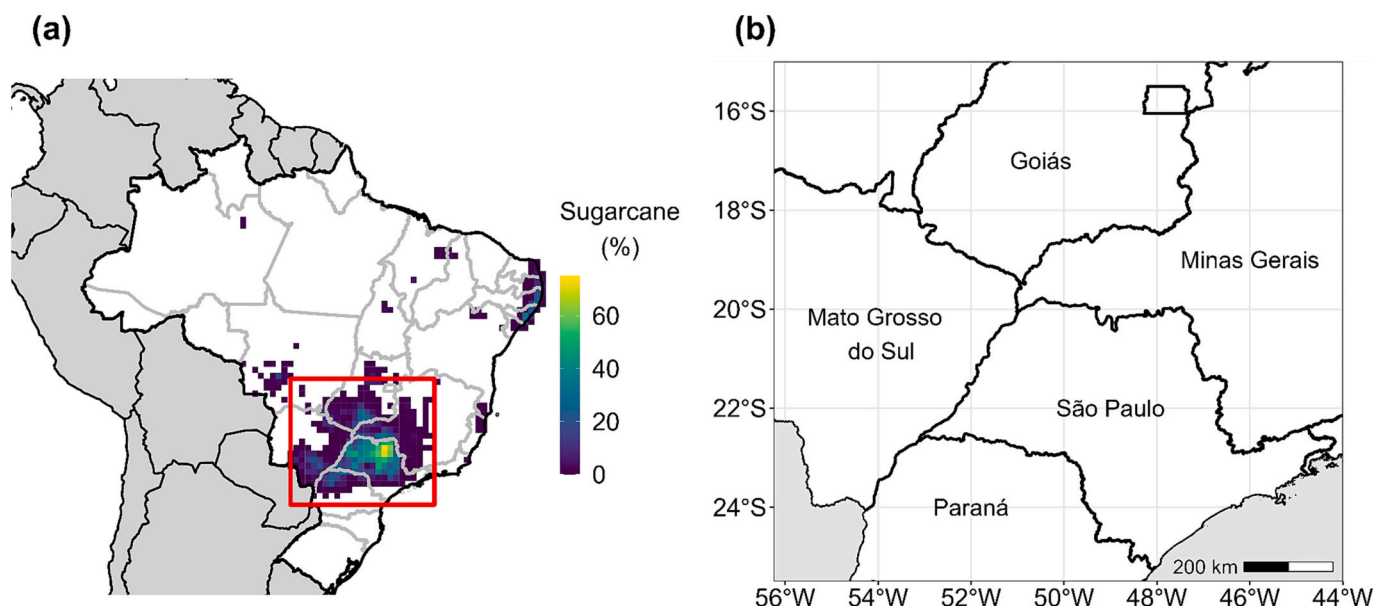
Sugarcane is the common name given to a diverse group of cultivated, sucrose-storing, tropical grasses that are an important food and commodity crop to many countries (Moore et al., 2014) and the source of ~80 % of world's sugar (FAOSTAT, 2021). Brazil is the world's largest producer of sugarcane, with approximately 99,706 km<sup>2</sup> given over to sugarcane production, resulting in 36.4 % of global output (i.e. 716 million tonnes), as compared to the second largest producer India at 405 million tonnes (20.5 % of global output) (FAOSTAT, 2021). The dominance of Brazil in sugarcane production has come about in part due to improved technologies and breeding programs, but in the main due to the rapid expansion in cultivated area over the last 20 years (Ogura et al., 2022), with production focused in south-central Brazil (Zheng et al., 2022). The state of São Paulo alone accounts for ~47 % of Brazil's total sugarcane production (Ogura et al., 2022) with the three next largest state producers (Goiás, Minas Gerais, and Mato Grosso do Sul) all in the south-central region (Fig. 1) and showing recent rapid expansion (Zalles et al., 2019). Brazil recognizes sugarcane as its third most valuable crop in terms of gross value (IBGE, 2023), supplying raw material for sugar, ethanol (biofuel), and direct energy production. Indeed sugarcane-derived products account for ~16.4 % of the Brazilian energy matrix (EPE, 2023). Sugarcane therefore contributes substantially to the bioeconomy of both Brazil and the state of São Paulo specifically (Ogura et al., 2022), and with increasing global demand for fossil fuel alternatives and the development of next generation biorefineries, a continued

expansion of this industry is likely in the coming years (Vandenbergh et al., 2022).

The recent rapid expansion of commodity crops such as sugarcane in Brazil has resulted in both the repurposing of existing C<sub>4</sub> grass dominated pasturelands and the conversion of natural vegetation, including both Amazon humid tropical forests and Cerrado dry tropical savannas (Ogura et al., 2022; Zalles et al., 2019). The resulting tension between the production of 'green fuels' at the cost of existing natural systems requires that steps are taken to maximise productivity of existing and planned sugarcane agricultural areas (Rossetto et al., 2022; Spera, 2017).

### 1.2. O<sub>3</sub> impacts on vegetation

At the Earth's surface, ozone (O<sub>3</sub>) is considered a major atmospheric pollutant, posing a risk to both human health and plant growth (Lelieveld et al., 2015; Mills et al., 2018a). Indeed the impact of O<sub>3</sub> on agronomic productivity has been appreciated since the early observation of 'oxidative stipple' in grape vines (Richards et al., 1958). The "O<sub>3</sub>-yield gap" recognized in many temperate and commodity crop species (Ainsworth, 2017; Mills et al., 2007) results in an estimated global productivity loss of 12.4 %, 7.1 %, 4.4 % and 6.1 % for soybean, wheat, rice and maize, respectively (Mills et al., 2018b). Recognition of the impacts of O<sub>3</sub> on both agronomic and natural systems, as well as the direct implications on human health, has resulted in air quality control treaties across many mid-latitude countries. Treaties such as the UNECE's Long-range Transboundary Air Pollution (LRTAP) Convention and the resulting Task Force on Hemispheric Transport of Air Pollution (TF



**Fig. 1.** Percentage of land cover dedicated to sugarcane production across Brazil in year 2020 (a) and modelling extent in south-central Brazil (b). Sugarcane coverage data from MapBiomass v6 ([www.plataforma.brasil.mapbiomass.org](http://www.plataforma.brasil.mapbiomass.org)) see Souza et al. (2020) for details, plotted here with a spatial resolution of 0.5°. Note map lines delineate general study area in South America and does not necessarily depict accepted national boundaries.

HTAP, [www.htap.org](http://www.htap.org)) that have helped to reduce O<sub>3</sub> or at least stabilize ambient O<sub>3</sub> levels across much of the global north (Doherty, 2015; Mills et al., 2018a).

### 1.3. Air quality and O<sub>3</sub> exposure in Brazil

Unlike countries in the global north, much of the (sub)tropics is seeing the continuing expansion of O<sub>3</sub> precursor emissions (especially NO<sub>x</sub>); from population growth, urbanization, biomass burning associated with land conversion, and the rise in large-scale agro-industry (Granier et al., 2000; Hewitt et al., 2009). When coupled to high temperatures and generally high levels of volatile organic compounds (VOCs) emitted by tropical vegetation, it is likely that O<sub>3</sub> responsiveness will emerge as a significant factor in limiting tropical crop yields and quality in the coming decades (Hayes et al., 2020; Tai et al., 2014).

In Brazil levels of O<sub>3</sub> are generally high, with large episodic events associated with vehicle exhaust in metropolitan areas (Schuch et al., 2019), biomass burning, and regional atmospheric drought (Targino et al., 2019). Historically, the controlled pre-harvest burning of sugarcane was itself responsible for considerable O<sub>3</sub> precursor emissions (Urban et al., 2016). However, this has been reduced by the industry's adoption of mechanization, in part due to efforts such as the "Green Ethanol" protocol, a voluntary standard established in response to São Paulo State Law number 11,241 banning the practice of pre-harvest burning (on lands with a low topographic gradient) by 2021 (ALESP, 2002). Yet, the waste sugarcane biomass (bagasse) is still often burnt for direct energy production in industrial plants which, given the trans-boundary movement of O<sub>3</sub> and its precursors (e.g. NO<sub>x</sub>), can still result in reduced air quality across rural areas (Squizzato et al., 2021).

Regions of poor air quality (including high O<sub>3</sub>) around south-central Brazil have been noted as having an impact on native vegetation (Moura et al., 2018a), and are projected to deteriorate further in the coming decades, both in response to continued precursor emissions from expanding megacities (Folberth et al., 2015), and due to changes in atmospheric chemistry in a warmer world (Brown et al., 2022). Given this trajectory, there is a need to consider how the magnitude of O<sub>3</sub> damage to sugarcane may vary across south-central Brazil in order to support a sustainable expansion of the industry (Rossetto et al., 2022).

### 1.4. Impact of O<sub>3</sub> on sugarcane

Sugarcane and related C<sub>4</sub> grass species have a biophysical CO<sub>2</sub> concentrating mechanism (Sage et al., 2014), resulting in high light, nitrogen, and water use efficiencies (i.e. carbon fixed per unit water lost) under warm-climatic conditions (Rao and Dixon, 2016). Given their inherent high water use efficiency, it is often presumed that C<sub>4</sub> grasses may avoid the deleterious effects of O<sub>3</sub> via stomatal exclusion; or that, by evolving effective antioxidant capacity to deal with climate induced oxidative stress, they may be well placed to deal with the reactive oxygen species formed by exposure to O<sub>3</sub> (Grantz and Vu, 2009). Certainly, in the early examination of C<sub>4</sub> grasses, it appeared that they showed only limited impacts of O<sub>3</sub> fumigation (Volin et al., 1998). However, from a recent broad examination of O<sub>3</sub> susceptibility in C<sub>4</sub> bioenergy crops by Li et al. (2022) it is clear that a high degree of variability exists, with initial studies indicating sugarcane itself to be highly susceptible to O<sub>3</sub> (Grantz and Vu, 2009; Grantz et al., 2012; Moura et al., 2018b). Yet despite this, it is notable that the only attempts to date to model the implications of sugarcane's O<sub>3</sub> susceptibility have been parameterized using the response of the C<sub>3</sub> crop cotton (*Gossypium hirsutum*) (Chuwah et al., 2015; Yi et al., 2018).

Commercial sugarcane cultivars are complex interspecific hybrids, primarily between *Saccharum officinarum*, known as the noble cane, and *S. spontaneum*, with contributions from *S. robustum*, *S. sinense*, *S. barberi*, and related grass genera such as *Miscanthus*, *Narenga*, and *Erianthus* (Moore et al., 2014). Domesticated sweet chewing cane *S. officinarum* was often hybridised with lines of *S. spontaneum* to confer reduced

susceptibility to abiotic and biotic stress, improved vegetative vigour, and to enhance traits beneficial to mechanized harvesting and processing. Although screening for O<sub>3</sub> tolerance is not currently a key trait in breeding programs, Grantz et al. (2012) demonstrated that the degree of photosynthetic inhibition seen under O<sub>3</sub> fumigation was inversely proportional to the contribution of *S. spontaneum* germplasm to the hybrid genome of four genetic lines. Similarly, it is well established that there are differences across modern sugarcane cultivars in traits, such as antioxidant capacity (Boaretto et al., 2014; Moura et al., 2018c) and intrinsic water use efficiency (Basnayake et al., 2015; Natarajan et al., 2021), that may well determine cultivar specific responses to O<sub>3</sub> (Wedow et al., 2021).

Given the current extent, and planned expansion of sugarcane production to meet global demand for 'green' biofuels across the world, there is a pressing need to characterize the potential risk of current O<sub>3</sub> exposure to the sugarcane industry. We therefore sought to: a) derive sugarcane O<sub>3</sub> dose-response functions across a full range of O<sub>3</sub> exposure; and b) model the implications of observed sugarcane susceptibility to O<sub>3</sub> across the globally important production area of south-central Brazil. Our findings will not only have direct implications on the world's primary sugarcane producing region, but will also inform global efforts to identify and address the O<sub>3</sub> yield gap under future climate and land use change scenarios.

## 2. Materials and methods

### 2.1. Experimental facility

All O<sub>3</sub> exposure studies were conducted at the joint University of Exeter (UoE) and James Cook University (JCU) TropOz research facility ([www.tropoz.org](http://www.tropoz.org)) located on the Nguma-bada campus of JCU in Cairns, Queensland, Australia. This unique tropical facility allows for the study of O<sub>3</sub> responses of plants grown under ambient tropical humid conditions, and consists of nine independently controlled and monitored Open Top Chambers (OTCs). Each chamber (internal volume 22.2 m<sup>3</sup>) was ventilated with charcoal filtered air at ~2 m<sup>3</sup> s<sup>-1</sup> using separate inline square centrifugal fans (ICQ560-VEE, Pacific Ventilation, Melbourne, VIC) augmented with O<sub>3</sub> generated on site and supplied to each chamber between 8:00 and 17:00. A different level of O<sub>3</sub> was applied to each individual chamber to achieve a semi-continuous gradient of nine different O<sub>3</sub> exposures (average chamber daytime concentrations ranged between 15 and 120 ppb, Table A1). This gradient-design provides a better approach than replicating a small number of exposure points, when seeking to identify a potentially nonlinear treatment response (Kreyling et al., 2018). Ozone concentrations in each chamber were monitored sequentially using an ultraviolet (UV) absorption O<sub>3</sub> analyser (Model 205, 2B technology, Boulder CO, USA) in air brought to a centralized service hub via a vacuum pump (Labport 840FT.18, KNF, Moreland West, VIC, Australia). A typical sampling sequence was achieved every 22 min allowing approximately three O<sub>3</sub> concentration readings per hour per chamber. Environmental variables including air temperature (T), vapour pressure deficit (VPD), and photosynthetically active radiation (PAR) were monitored using a single meteorological monitoring station (Campbell Scientific, Logan UT, USA) established in the central OTC and recording data averaged over every five minutes (Table A2).

### 2.2. Plant material

To develop O<sub>3</sub> dose-response functions for sugarcane, four *Saccharum* genetic lines were selected: *Saccharum officinarum* L. cv. Badila, a 'noble cane' often grown commercially in the early 20th Century and commonly used as the basis for breeding programs; *Saccharum spontaneum* cv. Mandalay, a clone used historically in Australian sugarcane breeding programs; and two commonly used commercial hybrid lines Q240 – which although being released in 2009 by Sugar Research

Australia (SRA) still dominates (~40 %) Australian (Queensland) sugarcane plantings, and CTC4 which represented around 10 % of 2019/20 plantings in the main producing areas of Brazil (Braga Junior et al., 2021).

Cane material held at the SRA germplasm collection (Meringa, Queensland, Australia) was supplied by SRA under licence in October 2021. This material was used to derive ~60 one-eye sets from each genetic line, which were treated with a systemic fungicide (Tilt 500 EC, Syngenta Australia, active constituent Propiconazole) and set to germinate under ambient shadehouse conditions at JCU's Environmental Research Complex ([www.jcu.edu.au/environmental-research-complex](http://www.jcu.edu.au/environmental-research-complex)). After one month an even cohort of sugarcane starts from each line were set out into individual 30-L pots containing a high organic matter potting mix augmented with a volcanic stone Quinkan to improve drainage. After establishment of the plants, all pots were maintained at close to field capacity with daily dripline irrigation, and fertilized every two weeks using Peters Professional Blossom Booster (10–13–16 + 1.2 Mg + TE, ICL Australia, Bella Vista, NSW, Australia).

For the determination of O<sub>3</sub> dose-response functions, four individuals of each genetic line were placed into each of the nine OTCs. Three of the varieties (Badila, Q240 and CTC4) were moved to the OTCs when ~10 cm tall on 13 November 2021 to begin the O<sub>3</sub> exposure experiment (duration 96 days). However, given initial slow growth, Mandalay was delayed and placed into the OTC on 28 March 2022 to begin its 100-day exposure. As a result of this delay different environmental conditions were experienced during the two experimental runs (Table A2); however, in both cases plants experienced full-sunlight and were not subjected to drought. Any residual differences in environmental conditions experienced between the two experimental runs should be accounted for in the calculation of O<sub>3</sub> flux dose-response functions (see Section 2.3).

### 2.3. Characterizing O<sub>3</sub> dose-response functions of sugarcane biomass

At the conclusion of the experiment, plants were harvested and oven dried (70 °C) until constant mass to determine biomass production. Biomass was partitioned into: leaf (all photosynthetic material distal from leaf ligules), stalks (including cane and leaf sheath), and roots. At this stage, the most recently emerged leaf after ligule separation (L + 1) from each plant was collected separately to determine individual leaf area, dry biomass and thereby leaf mass per area (LMA).

In addition to relating changes in biomass to the concentration-based O<sub>3</sub> exposure metric AOT40 (ppm.h), we estimated accumulated O<sub>3</sub> flux into leaves using the Deposition of O<sub>3</sub> for Stomatal Exchange (DO<sub>3</sub>SE) model v3.1. The use of DO<sub>3</sub>SE to calculate the Phytotoxic Ozone Dose (POD<sub>y</sub>, mmol m<sup>-2</sup>, above a threshold *y* in nmol O<sub>3</sub> m<sup>-2</sup> projected leaf area (PLA) s<sup>-1</sup>) provides for an O<sub>3</sub> metric that accounts for differences in stomatal conductance as a result of species-level traits and environmentally dynamic conditions. Using flux-based metrics allows for the comparison of O<sub>3</sub> susceptibilities collected under diverse environmental conditions and facilitates the linking of observed susceptibility to dynamic vegetation models. Dynamic models which integrate vegetation responses to multiple environmental factors such as soil moisture and temperature to determine O<sub>3</sub> flux (Emberson, 2020; Pleijel et al., 2022).

We modelled stomatal conductance in DO<sub>3</sub>SE using an empirically derived model (Jarvis, 1976; Emberson et al., 2000). In the weeks prior to harvest, stomatal conductance to water vapour (g<sub>s</sub>) was measured on both the abaxial and adaxial surfaces of L + 1 leaves of all plants from CTC4, Q240 and Badila using a handheld porometer (SC1, Decagon Devices, Pullman WA, USA). As porometer data could not be collected on Mandalay, given its narrow leaf blade, and the fact we observed no significant decline in g<sub>s</sub> of sunlit (PAR > 1500 μmol m<sup>-2</sup> s<sup>-1</sup>) L + 1 leaves as measured using the porometer across O<sub>3</sub> exposure, we also collected data from the control chamber plants of all genotypes using a portable photosynthesis analyser (LI-6400XT, LiCOR Biosciences, Lincoln NE, USA). This leaf-level gas-exchange data comprised survey

measurements collected every three minutes for ~24 h per leaf using a buffer volume and with the LI-6400xt tracking ambient PAR and temperatures. All g<sub>s</sub> data were converted to Relative Stomatal Conductance (RSC) before fitting DO<sub>3</sub>SE Jarvis parameters (Tables 1 and A3) using the method described by Hayes et al. (2020a), with stomatal conductance of O<sub>3</sub> (g<sub>o3</sub>) = 0.663 × g<sub>s</sub>. The ability of the DO<sub>3</sub>SE model to represent g<sub>s</sub> in sugarcane was verified by comparing modelled values with those measured using the LI-6400xt (Fig. A1). All genotypes showed a strong positive correlation between observed and modelled daylight (i.e. 5:30 to 19:00) data with an R<sup>2</sup> ranging from 0.59 in CTC4 to 0.83 in Mandalay.

The O<sub>3</sub> flux-based metric POD<sub>y</sub> was calculated for each genotype-chamber combination using the DO<sub>3</sub>SE model calibrated using gas exchange data (Table 1) and *y*-values 0 to 8 nmol m<sup>-2</sup> s<sup>-1</sup>. The *y*-value represents an instantaneous O<sub>3</sub> flux below which no damage due to O<sub>3</sub> is assumed to occur, and is considered reflective of plants' resistance to O<sub>3</sub> (Agathokleous and Saitanis, 2020). We selected two thresholds (i.e. 0 and 2 nmol m<sup>-2</sup> s<sup>-1</sup>) to use in the regional simulations (see Section 2.4); the use of POD<sub>0</sub> did not presuppose the same inherent resistance across sugarcane genotypes (Agathokleous et al., 2019), whereas POD<sub>2</sub> allowed for comparison with previously published O<sub>3</sub> dose-response functions (Moura et al., 2018b). Dose-response functions were calculated for the relative biomass decline using a maximum biomass estimated as the *y*-intercept of a linear regression between average total biomass (*n* = 4) of each chamber (*n* = 9) and the calculated POD<sub>y</sub>.

### 2.4. Modelling impacts of O<sub>3</sub> exposure in south-central Brazil

To model the potential risks of current O<sub>3</sub> levels to sugarcane production across south-central Brazil, we used the Joint UK Land Environment Simulator (JULES) (Best et al., 2011; Clark et al., 2011) v5.6. JULES is a land surface model used to study soil-vegetation-atmosphere interactions in which the land is divided into gridcells and vegetation is represented by up to 13 plant functional types (PFTs). Outputs are given as an average for each gridcell assuming homogeneity within the gridcell, and each PFT represents the average behaviour of that vegetation type based on observations (Harper et al., 2016). Sugarcane is a C<sub>4</sub> crop so here we focused only on the C<sub>4</sub> PFT, with modifications made to better represent sugarcane (see Section 2.4.1). Each gridcell can contain a mixture of PFTs and non-vegetation cover, and we prescribed the fraction of the C<sub>4</sub> PFT in each gridcell to match the observed distribution of sugarcane across south-central Brazil (see Section 2.4.5).

This modelling framework incorporates continuous and spatially explicit environmental information (e.g. meteorological conditions and [O<sub>3</sub>]) and calculates vegetation responses and fluxes in each gridcell, taking into account soil properties and vegetation processes, including photosynthesis, respiration, and carbon partitioning for each PFT. JULES outputs information at the gridcell scale, rather than at the individual plant scale, and therefore differs from some detailed crop models by not including management decisions such as harvesting and ratooning. However, the model has substantial complexity including the representation of vegetation responses to atmospheric composition, e.g. CO<sub>2</sub> (Huntingford et al., 2013), aerosols (Mercado et al., 2009; Rap et al., 2018), and O<sub>3</sub> (Leung et al., 2022), as well as interaction with other abiotic factors such as temperature (Huntingford et al., 2017), drought (Harper et al., 2021), and changes in nutrient cycling (Huntingford et al., 2022). The model is regularly updated to incorporate new process understanding and additional observations, and the specific model set-up used here is described in more detail in Section 2.4.1.

#### 2.4.1. Details on JULES environment and parameterization

Sugarcane was represented in JULES by using the C<sub>4</sub> PFT. However, to better represent carbon fixation, respiration and stomatal conductance of sugarcane, we used PFT parameters adapted by Vianna et al. (2022). The work by Vianna et al. (2022) used field data from 11 sites across Brazil representing four cultivars (RB867515, IACSP95-5000,

**Table 1**

Genotype specific stomatal conductance parameterization utilized in DO<sub>3</sub>SE to calculate Phytotoxic Ozone Dose (POD).  $g_{O_3\text{-max}}$  is the maximum stomatal conductance to O<sub>3</sub>,  $f_{\text{min}}$  is the fraction of  $g_{O_3\text{-max}}$  at minimum stomatal conductance ( $g_{O_3\text{-min}}$ ).  $L_d$  the effective leaf blade width,  $f_{\text{temp}}$ ,  $f_{\text{PAR}}$ ,  $f_{\text{VPD}}$  are the functions of  $g_s$  response to air temperature (T, °C), photosynthetically active radiation at the leaf surface (PAR,  $\mu\text{mol m}^{-2} \text{s}^{-1}$ ), and vapour pressure deficit (VPD, kPa) respectively. Data derived from leaf-level gas exchange using Li-6400XT.

Genotype	$g_{O_3\text{-max}}$ ( $\text{mmol m}^{-2} \text{s}^{-1}$ )	$L_d$ (m)	$f_{\text{min}}$ (fraction)	$f_{\text{PAR}}$ (unitless)	$f_{\text{VPD}}$		$f_{\text{temp}}$		
					VPD <sub>min</sub> (kPa)	VPD <sub>max</sub> (kPa)	T <sub>min</sub> (°C)	T <sub>opt</sub> (°C)	T <sub>max</sub> (°C)
Badila	185	0.05	0.06	0.005	3.5	6.0	21	34	47
CTC4	172	0.05	0.06	0.008	3.8	5.8	24	37	50
Mandalay	245	0.01	0.06	0.004	1.67	6.64	16	31	46
Q240	153	0.05	0.06	0.010	3.5	7.0	20	35	50
IACSP94-2094 <sup>a</sup>	363	0.05	0.06	0.0014	1.93	5.40	15	32	46
IACSP95-5000 <sup>a</sup>	342	0.05	0.06	0.0015	1.60	6.82	13	32	46

<sup>a</sup> Data from Moura et al. (2018b).

RB72454, and CTC14) to tune JULES-Crop for its application to sugarcane. Although this modelling framework does not account for genotype explicitly in the selection of fitting parameters (Table A4), when evaluated it was able to reproduce observed GPP ( $\text{kg C m}^{-2} \text{yr}^{-1}$ ) with an  $r^2 = 0.78$  and an accuracy metric ( $d$ ) = 0.92, representing a RMSE of 6.75  $\text{Mg ha}^{-1}$  of stalk dry matter (Vianna et al., 2022).

Stomatal conductance ( $g_s$ ) within JULES is calculated using the Medlyn stomatal conductance model (Medlyn et al., 2011), parameterized using  $g_l$  from Oliver et al. (2022) ( $g_l = 1.62$  for C<sub>4</sub> plants). Photosynthetic rates were calculated using the Collatz photosynthesis model, and O<sub>3</sub> flux calculated as per Oliver et al. (2018). See Appendix A: Supplementary Information for further details. The Collatz model in JULES uses the photosynthesis parameter  $V_{\text{max}}$ , calibrated by Vianna et al. (2022) based on the top leaf nitrogen concentration (Table A4).

#### 2.4.2. Ozone data

Given a paucity of directly measured O<sub>3</sub> data across much of south-central Brazil (especially in rural sugarcane growing regions), the O<sub>3</sub> exposure data used herein were simulated using the Earth System Model UKESM1 for the period 2000 to 2015. Modelled [O<sub>3</sub>] data are available at a monthly resolution representing 0 to 40 m above orography, and at a horizontal resolution of 1.25° latitude by 1.875° longitude. For our work, the monthly mean surface [O<sub>3</sub>] was adapted to reflect both the diel cycle and daily variation observed in the region. Specifically, we projected the same daily variation and diel cycle seen at 53 point-locations in the state of São Paulo in 2018 (CETESB, 2022) to every modelled grid cell (see Appendix A: Supplementary Information, and Figs. A2 to A5 for details). This provided the realistic and biologically relevant [O<sub>3</sub>] climatology required to drive our model, and notably resulted in an ~40 % increase in annualized flux of O<sub>3</sub> as compared to the use of monthly mean concentrations as given by UKESM1 (Fig. A6).

#### 2.4.3. Calibrating JULES for sugarcane susceptibility to O<sub>3</sub>

The O<sub>3</sub> damage scheme employed here in JULES is the same as that in Sitch et al. (2007) (see Eqs. (1), (2), and (3)) and works by modifying net photosynthesis ( $A_{\text{net}}$ ) and stomatal conductance ( $g_s$ ) by an O<sub>3</sub> damage factor ( $F$ ). With  $F$  defined (Eq. (1)) by a sensitivity parameter ( $\alpha$ ) and the flux of O<sub>3</sub> above a threshold ( $y$ ) so that  $A$  decreases linearly (Eq. (2)) as O<sub>3</sub> flux increases above the threshold and the rate of decrease depends on the sensitivity parameter. The decrease in  $A$  affects the Net Primary productivity (NPP), and the model assumes that a) O<sub>3</sub> damage is instantaneous at the point of uptake and b) results in a coordinated reduction in  $g_s$  (Eq. (3)).

$$F = 1 - \alpha \times (\text{Flux } O_3 > y) \quad (1)$$

$$A_{\text{mod}} = A_{\text{net}} \times F \quad (2)$$

$$g_{\text{mod}} = g_s \times F \quad (3)$$

We derived  $\alpha$  within the damage scheme using two thresholds of  $y = 0$ , and 2  $\text{nmol m}^{-2} \text{s}^{-1}$ . To do this we calculated the reduction in modelled NPP compared to a simulation with no O<sub>3</sub> damage for each grid cell over a yearlong run of JULES, using a first approximation for  $\alpha$ . Iterative adjustment of  $\alpha$  was then carried out until the relative NPP loss modelled in each grid cell due to the O<sub>3</sub> flux matched the dose-response functions observed in this study (i.e. Mandalay, combined commercial, and Badila, see Results and Table 3). We also tuned  $\alpha$  to fit the observed O<sub>3</sub> dose-response function for two highly susceptible sugarcane varieties (IACSP94-2094 and IACSP95-5000) published by Moura et al. (2018b). This resulted in the development of eight different  $\alpha$  functions across four different susceptibilities (e.g. low, moderate, high, and very high) and two thresholds; and was aimed at characterizing the full range of potential O<sub>3</sub> impacts on sugarcane production (Table 3).

#### 2.4.4. JULES model runs

To develop spatially explicit risk maps of potential O<sub>3</sub> impacts on sugarcane production, JULES was used to calculate annual yields with or without consideration of O<sub>3</sub> susceptibility for a 10-year period (after a 20-year model spin up) across south-central Brazil. Meteorological forcing at 6-h intervals for the years 2005 to 2015 was taken from CRUJRA v2.1 reanalysis (Harris et al., 2020; Kobayashi et al., 2015) at a horizontal resolution of 1.25° latitude by 1.875° longitude, whilst O<sub>3</sub> data was supplied as a yearly climatology with hourly variation as described above and in Appendix A Supplementary Information. Spatially explicit model outputs using  $\alpha$  calibrated at four levels of susceptibility (i.e. low, moderate, high and very high) and two thresholds (i.e. 0 and 2  $\text{nmol m}^{-2} \text{s}^{-1}$ ) were compared to the control model output (i.e. with no O<sub>3</sub> damage) to calculate both proportional decline (% of control) and absolute impacts (reduction in NPP;  $\text{kg C m}^{-2} \text{yr}^{-1}$ ).

#### 2.4.5. Scaling of risk to impacts on sugarcane production and conversion to yield

To convert the modelled risk of O<sub>3</sub> across the landscape to potential reductions in regional sugarcane production, we scaled JULES model outputs by the fractional cover of sugarcane found in each grid cell in the year 2020 (Fig. 1). Sugarcane coverage was derived from MapBiomass v6 ([www.plataforma.brasil.mapbiomass.org](http://www.plataforma.brasil.mapbiomass.org)) (see Souza et al. (2020) for details) and resampled at a spatial resolution (i.e. 1.25° latitude by 1.875° longitude) commensurate with a JULES run. Furthermore, to estimate yield losses of fresh cane ( $\text{Mg ha}^{-1}$ ) we converted from JULES output NPP ( $\text{kg C m}^{-2} \text{yr}^{-1}$ ) employing standard assumptions informed by crop modelling of mature sugarcane (Dias and Sentelhas, 2018; Marin et al., 2016; Marin et al., 2015; Souza et al., 2020). Specifically this included assuming fixed carbon content of biomass of 50 %, above ground biomass representing 80 % of total biomass, a harvest index (proportion cane of total above ground biomass production) of 70 %, and a constant dry matter content of 25 %.

2.5. Statistical analysis

All statistical analyses were conducted in R v 4.2.0 (R Core Team, 2022). Our determination of O<sub>3</sub> dose-response functions utilized a gradient design (Kreyling et al., 2018) in which individual chambers represented an un-replicated point on the gradient of O<sub>3</sub> exposure. A linear regression between chamber averaged data for individual plants (n = 4) and O<sub>3</sub> metric (i.e. AOT40, or POD<sub>y</sub>) was translated to genotype specific O<sub>3</sub> dose-response functions (i.e. changes in relative biomass) using a maximum biomass estimated as the y-intercept of a linear regression between total biomass and the O<sub>3</sub> metric. When considering morphological impacts of O<sub>3</sub> exposure on sugarcane traits, we fitted a linear mixed effects model to predict, for example LMA, with increasing POD<sub>y</sub> and included ‘genotype’ as a random effect.

3. Results

3.1. O<sub>3</sub> dose-response function of sugarcane

Sugarcane genotypes grown as part of this study showed significant differences in growth over the ~100 day experimental period (Table 2, Fig. 2a). The two commercial varieties (i.e. CTC4, Q240) produced substantially more biomass than either the *S. officinarum* cv. Badila or *S. spontaneum* cv. Mandalay, with Mandalay having the lowest final biomass under low [O<sub>3</sub>]. All genotypes tested generally showed a decline in biomass with increasing O<sub>3</sub> exposure whether exposure was expressed in terms of [O<sub>3</sub>] (i.e. AOT40) or O<sub>3</sub> flux (i.e. POD<sub>0</sub> or POD<sub>2</sub>). The response of relative biomass to the metric AOT40 ranged from -0.0015 in Mandalay to -0.0017, -0.0025 and -0.0069 in Q240, CTC4 and Badila respectively. In considering susceptibility to O<sub>3</sub> flux, the genotype-specific O<sub>3</sub> dose-response functions also varied (Table 2, Figs. 2b and A7) from the least susceptible Mandalay, through Q240 and CTC4, to the most susceptible genotype Badila. The pattern of O<sub>3</sub> susceptibility across genotypes was the same when using either no O<sub>3</sub> flux threshold (i.e. POD<sub>0</sub>) or a threshold of 2 nmol m<sup>-2</sup> s<sup>-1</sup> (i.e. POD<sub>2</sub>). Generally we estimated a more negative slope (greater susceptibility) within a genotype when using an increased threshold (Fig. A7). However, in most species the fit was not significantly different between model estimates of relative biomass decline when using a threshold of 0 or 6 nmol m<sup>-2</sup> s<sup>-1</sup>, as indicated by an overlap in confidence intervals. Only in Badila did we see an apparent significant increase in susceptibility when using a greater threshold, possibly as a result of the non-linear response in biomass response observed in this genotype (Fig. 2a).

3.2. Morphological impacts of O<sub>3</sub> on sugarcane

Across all sugarcane genotypes, increasing O<sub>3</sub> flux had a statistically significant (p < 0.01) and negative impact (beta = -0.15, 95 % CI [-0.24, -0.06], t<sub>(32)</sub> = -3.28) on the LMA of the most recently emerged leaf (i.e. L + 1) (Fig. A8). Similarly, across all genotypes there was a highly significant (p < 0.001) and positive impact of O<sub>3</sub> flux on carbon partitioning with a significant impact of O<sub>3</sub> on the ratio of above ground

Table 2

Genotype specific O<sub>3</sub> dose-response functions (±1 S.E.) of sugarcane grown for ~100 days in open top chambers under a range of [O<sub>3</sub>] (Fig. 2), using two thresholds for calculating Phytotoxic Ozone Dose (0 and 2 nmol m<sup>-2</sup> s<sup>-1</sup>). Relative biomass calculated after setting y-intercept to 1 at POD<sub>y</sub> = 0.

POD	Genotype	Total biomass (g)			Relative biomass			p-Value
		Intercept	Slope	S.E.	Slope	S.E.	95% CI	
POD <sub>0</sub>	Badila	388	± 22	-3.91	± 0.73	-0.0101	± 0.0571	<0.01
	CTC4	435	± 14	-1.90	± 0.50	-0.0044	± 0.0011	<0.01
	Mandalay	264	± 11	-0.35	± 0.42	-0.0013	± 0.0016	ns
	Q240	531	± 22	-1.63	± 0.83	-0.0031	± 0.0016	<0.1
POD <sub>2</sub>	Badila	363	± 17	-3.99	± 0.73	-0.0109	± 0.0020	<0.001
	CTC4	423	± 11	-1.96	± 0.51	-0.0046	± 0.0012	<0.01
	Mandalay	263	± 10	-0.33	± 0.40	-0.0013	± 0.0015	ns
	Q240	521	± 17	-1.68	± 0.86	-0.0032	± 0.0016	<0.1

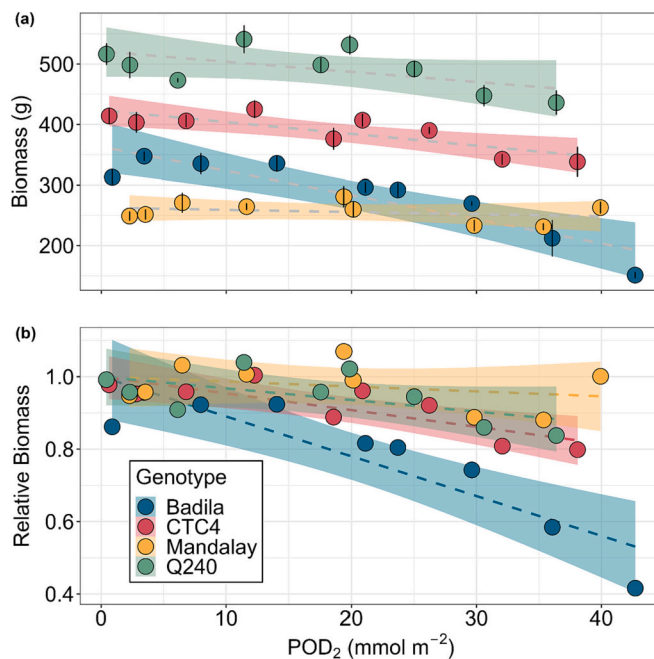


Fig. 2. Ozone dose-response functions of four genetic lines of sugarcane. Values represent average (n = 4 ± 1.S.E.) change in biomass (a) and relative biomass (b) as compared to phytotoxic O<sub>3</sub> dose (POD) above a threshold of 2 nmol m<sup>-2</sup> s<sup>-1</sup>.

Table 3

JULES O<sub>3</sub> sensitivity parameter (α) calibrated for four sugarcane scenarios.

Genotype susceptibility	JULES O <sub>3</sub> sensitivity parameter (α)	
	POD <sub>0</sub>	POD <sub>2</sub>
Low <sup>a</sup>	0.04	0.04
Moderate <sup>b</sup>	0.1	0.1
High <sup>c</sup>	0.25	0.28
Very high <sup>d</sup>	0.9	0.75

<sup>a</sup> Calibrated using *S. spontaneum* cv. Mandalay (this study).

<sup>b</sup> Calibrated using average of CTC4 and Q240 (this study).

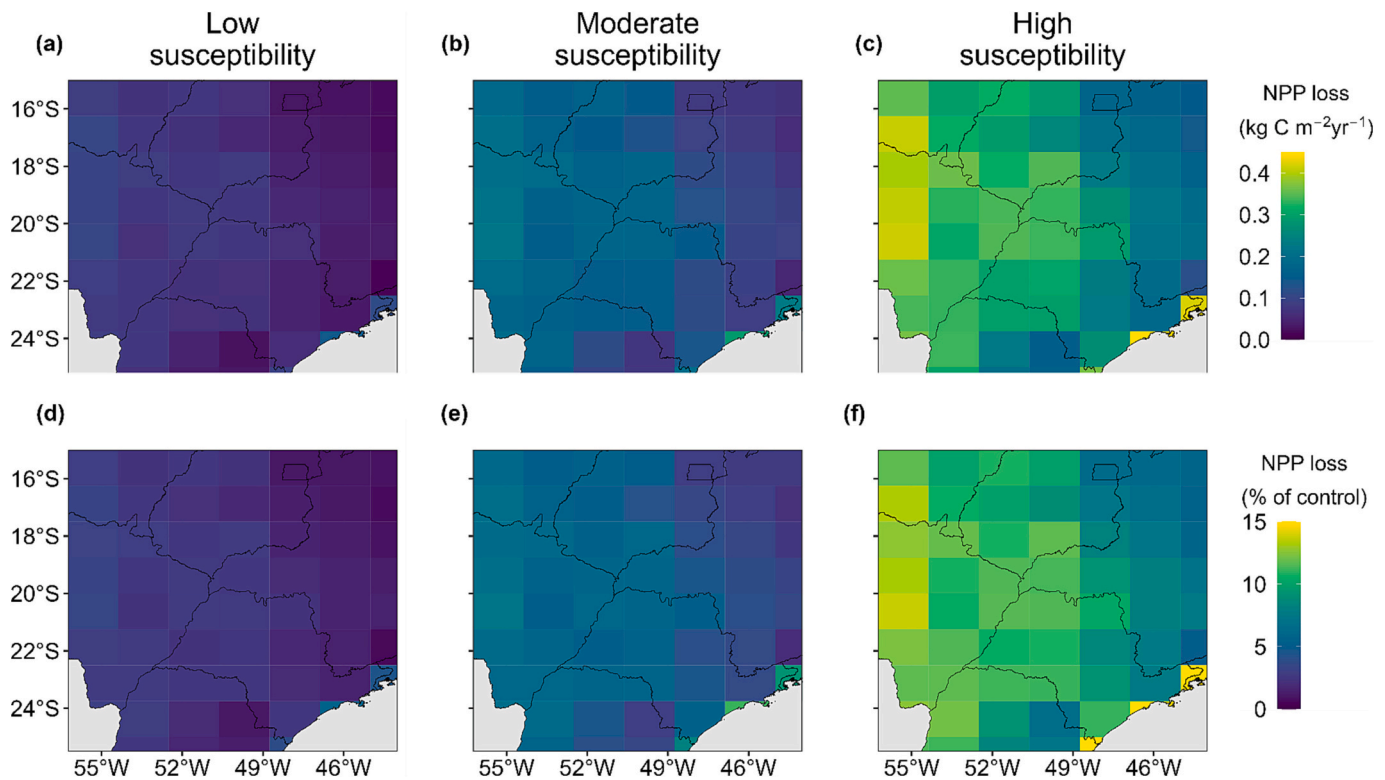
<sup>c</sup> Calibrated using *S. officinarum* cv. Badila (this study).

<sup>d</sup> Calibrated using average of IACSP94-2094 and IACSP95-5000 (Moura et al., 2018b).

to below ground biomass (beta = 0.03, 95 % CI [0.02, 0.04], t<sub>(32)</sub> = 7.75, Fig. A9).

3.3. Modelling risk of O<sub>3</sub> across south-central Brazil

Our model results highlighted the potential risk of O<sub>3</sub> to sugarcane NPP and, therefore, total production across south-central Brazil (Figs. 3 and A10). All dose-response functions tested showed current O<sub>3</sub>

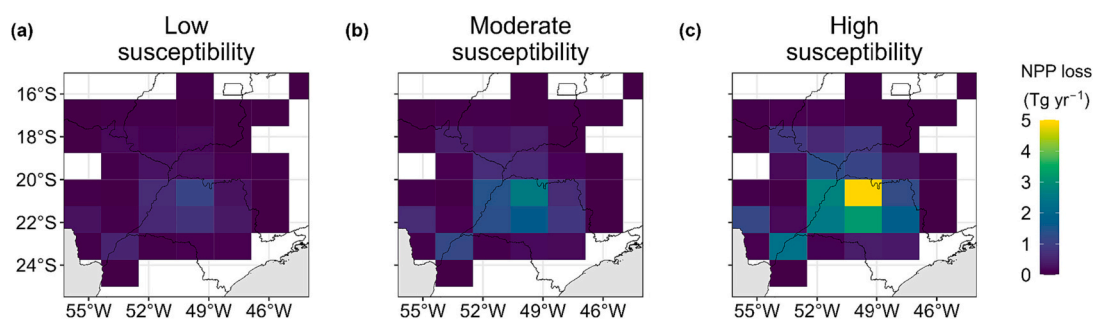


**Fig. 3.** Modelled risk of present day (i.e. 2010 to 2015)  $[O_3]$  on potential sugarcane production across south-central Brazil, expressed as absolute reduction in NPP ( $kg\ C\ m^{-2}\ yr^{-1}$ ) and % loss of control NPP (i.e. no  $O_3$ ). Model calibrated against the range of observed  $O_3$  susceptibility seen in this study examining biomass decline against  $POD_2$ , with low susceptibility (a, d) representing *S. spontaneum* cv. Mandalay, moderate susceptibility (b, e) representing two commercial sugarcane cultivars (CTC4 and Q240) and high susceptibility (c, f) representing *S. officinarum* cv. Badila. Note for clarity cells exceeding upper limit of scale in sub-figures (c) and (f) set to upper limits of  $0.45\ kg\ C\ m^{-2}\ yr^{-1}$  and 15 % respectively.

exposure would likely lead to a reduction in sugarcane production across the region. The magnitude of this risk was impacted by spatial variation in  $O_3$  exposure, stomatal conductance, and which dose-response function (i.e. low-, moderate- or high- susceptibility) and threshold value (i.e.  $POD_0$  or  $POD_2$ ) was implemented. Examining the range of potential risk using our moderate susceptibility function (representative of two commercial varieties Q240 and CTC4), we saw the risk to NPP across the region ranging from 0.05 to 0.29 (average 0.14)  $kg\ C\ m^{-2}\ yr^{-1}$  (Fig. 3a-c) representing between 2.0 and 11.2 % (average 5.1 %) of control model NPP (Fig. 3d-f) when using a threshold of  $2\ nmol\ m^{-2}\ s^{-1}$ . The risk of NPP loss increased to between 0.36 and 0.62 (average 0.50)  $kg\ C\ m^{-2}\ yr^{-1}$  (i.e. 14.7 to 23.6 % (average 17.8 %) of control model NPP) when using no threshold (Fig. A10). For additional context, we also modelled the potential risks on production using previously published data from two other commercial varieties grown in Italy, shown to be

highly susceptible to  $O_3$  (i.e. IACSP94-2094 and IACSP95-5000). Although the comparison of data derived from OTC's and free air  $O_3$  enrichment should be made with caution (Montes et al., 2022), using the observed dose-response for these varieties, current  $O_3$  exposure in south-central Brazil (Fig. A11) would be expected to reduce NPP by on average  $0.42\ kg\ C\ m^{-2}\ yr^{-1}$  (i.e. 14.9 % of control NPP) using  $POD_2$  or  $1.69\ kg\ C\ m^{-2}\ yr^{-1}$  (i.e. 61 % of control NPP) using  $POD_0$  (Fig. A11).

In converting spatially explicit maps of the  $O_3$  risk across the landscape to consider the likely impacts on regional production, we took into account where sugarcane production currently occurs. Specifically, we took the fractional cover of sugarcane found across the region in 2020 (Figs. 4 and A12). Depending upon the  $O_3$  susceptibility assumed (i.e. low, moderate or high), we predicted NPP of total sugarcane across the region to be reduced by between 2.5 and 11 % when using  $POD_2$ . However, when examining only the commercial cultivars (i.e. moderate



**Fig. 4.** Predicted total production losses due to present day  $O_3$  damage in sugarcane across south-central Brazil, assuming sugarcane growing locations as per the year 2020 (Souza et al., 2020). Model calibrated using three susceptibilities to  $O_3$  using the flux metric  $POD_2$ . Control model NPP total equals  $243\ Tg\ yr^{-1}$  with predicted total losses in (a) low susceptibility representing *S. spontaneum* cv. Mandalay of 2.5 %, (b) moderate susceptibility representing two commercial sugarcane cultivars (i.e. CTC4 and Q240) of 5.6 %, and (c) high susceptibility representing *S. officinarum* cv. Badila of 11 %.

susceptibility) the likely reduction in NPP increased from 5.6 to 18.3 % of the control scenario NPP when considering  $POD_0$  instead of  $POD_2$ .

#### 4. Discussion

Our results demonstrate that experimental  $O_3$  exposure has a significant and substantial impact on the morphology and biomass accumulation of the globally important  $C_4$  bioenergy crop sugarcane. The decline in relative biomass seen with increasing AOT40 averaged  $-0.21$  % per ppm.h in the two commercial varieties (i.e. CTC4 and Q240) tested making them comparable with other moderately ' $O_3$  sensitive' crops (Mills et al., 2007) and generally greater than the cotton parameterization previously used in assessment of  $O_3$  impacts on sugarcane production, i.e.  $-0.1495$  % per ppm.h used by Yi et al. (2018) and Chuwah et al. (2015). When making our calculations we assumed a linear decrease in yield with increase in  $O_3$  (concentration or flux), as this is in line with the methodology currently used (CLRTAP et al., 2017). For the varieties Mandalay, CTC4 and Q240 a linear response is appropriate, whereas for Badila the response may be sigmoidal over the range of  $O_3$  exposure used. As this was consistent across the range of flux-thresholds tested, we retained the use of a linear function.

When considering the more biologically relevant metric of  $O_3$  flux under prevailing experimental conditions (i.e.  $POD_y$ ) we still see a substantial difference in susceptibility of genotypes tested (Table 2, Fig. A7), highlighting the likely role that fundamental plant traits play in shaping observed  $O_3$  responses. In our novel dataset, we observed a trend in  $O_3$  susceptibility across genotypes similar to the decline of net photosynthesis reported by Grantz et al. (2012), with values ranging from *S. spontaneum* (low susceptibility) through commercial hybrids (moderate susceptibility) to *S. officinarum* (high susceptibility). In comparing the  $O_3$  susceptibility reported here to the only comparable  $O_3$ -flux data available for sugarcane, an average dose-response at  $POD_2$  of  $-0.03645$  seen in above ground biomass of genotypes IACSP95-5000 and IACSP94-2094 (Moura et al., 2018b), we found all genotypes tested here to be generally less susceptible (Table 2). Using the 'very high' susceptibility observed in Moura et al. (2018b) to parameterize JULES resulted in a substantial increase in modelled risk to production across south central Brazil (c.f. Figs. 3 and A11). Although this is likely due to the difference in cultivar susceptibility often seen in  $C_4$  crops (Li et al., 2022), differences between the experimental design and analysis i.e. repeated-treatments in Moura et al. (2018b), and gradient-designs (this study), as well as fundamental differences between responses seen in OTC and free air  $O_3$  enrichment studies (Montes et al., 2022) likely play a role. Further work to identify drivers of  $O_3$  susceptibility in relevant production genotypes is needed to assess the accuracy of any future model parameterizations.

When using our observed  $O_3$  susceptibility to parameterize the dynamic vegetation model JULES, we found that current  $[O_3]$  in south-central Brazil is likely contributing to a substantial yield gap (i.e. difference between potential and observed yields) seen in sugarcane across the region. Our control model with sugarcane not exposed to  $O_3$  predicted potential NPP of up to  $3.1 \text{ kg C m}^{-2} \text{ yr}^{-1}$ , equating to a yield of  $\sim 138 \text{ Mg ha}^{-1}$ . The potential risk of current  $O_3$  on production of commercial cultivars (moderate susceptibility in this study) across south-central Brazil equated to a yield gap of between  $2.2$  and  $13.0 \text{ Mg ha}^{-1}$  (average  $6.3 \text{ Mg ha}^{-1}$ ) when using  $POD_2$ , and between  $16.1$  and  $27.8 \text{ Mg ha}^{-1}$  (average  $22.4 \text{ Mg ha}^{-1}$ ) using  $POD_0$ . This resulted in a prediction of water-limited yield ( $Y_w$ ), while accounting for  $O_3$  impacts, in a range comparable to that derived from calibrated crop modelling estimates at between  $93.5$  and  $132 \text{ Mg ha}^{-1}$  (average  $109 \text{ Mg ha}^{-1}$ ) in south-central Brazil (Dias and Sentelhas, 2018). Calibrated crop models do not currently account for the impacts of  $O_3$ , however by their nature they implicitly account for  $O_3$  damage in the data used to parameterize them. While observed yields in the region are currently much lower than even water-limited potential at between  $38.5$  and  $90.4 \text{ Mg ha}^{-1}$  (average

$78 \text{ Mg ha}^{-1}$ ) (Dias and Sentelhas, 2018), our finding suggests that  $O_3$  is likely contributing to the substantial yield gap observed in Brazil (Dias and Sentelhas, 2018; Marin et al., 2016) and that, even if improvements to growing conditions and management practices are made, a yield gap will likely remain. Indeed, the adoption of improved irrigation (currently only a small fraction of total production area) may lead to a greater  $O_3$  penalty via the removal of stomatal exclusion of  $O_3$  through reduced  $g_s$  during periods of drought (Gao et al., 2017; Harmens et al., 2019), which are often associated with poor air quality (Targino et al., 2019).

Given the range of genetic material used in modern sugarcane breeding programs, there is the potential for selecting tolerant commercial cultivars for regions of particularly high  $O_3$  risk. However, there is also the more general need to consider  $O_3$  sensitivity in the screening of new cultivars to avoid inadvertently developing 'high yield' varieties that are more susceptible to  $O_3$  (Osborne et al., 2016). We suggest that cultivar  $O_3$ -susceptibility, the possible counter-intuitive impacts of irrigation, and predictions of future air quality across Brazil all need to be considered to ensure the sustainable expansion of the industry (Rossetto et al., 2022) within Sugarcane Agroecological Zoning (da Silva et al., 2021).

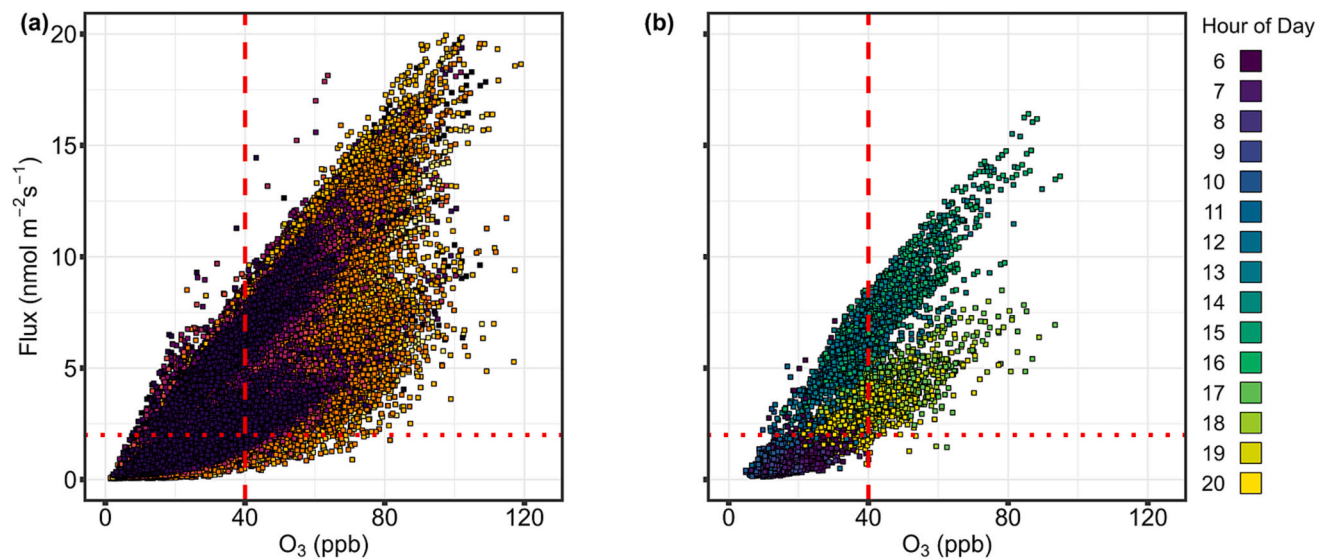
The advantages of a flux based approach to assessing  $O_3$  susceptibility have been well established (Plejel et al., 2022) and has been used here to translate observed susceptibility in sugarcane to a landscape-scale model that has an ability to predict dynamic  $g_s$ . In plotting the relationship between ambient  $O_3$  concentration and modelled stomatal flux of  $O_3$  (Fig. 5) across south-central Brazil we show how stomatal exclusion – the limitation of  $O_3$  flux due to stomatal closure – results in a broad range of  $O_3$  fluxes calculated for any given level of  $[O_3]$  exposure. This avoidance of  $O_3$  via stomatal exclusion would not be reflected in concentration metrics such as AOT40 or SUM60 used to characterize  $O_3$  exposure.

When calculating  $POD_y$ , the use of a threshold value (e.g. 1, 2 or  $6 \text{ nmol m}^{-2} \text{ s}^{-1}$ ) is often expounded to account for the plant's intrinsic antioxidant capacity or indeed the commonly observed hormetic response of plants to increasing  $[O_3]$  (Agathokleous et al., 2019). Across our control model run, the consideration of an instantaneous threshold of  $2 \text{ nmol m}^{-2} \text{ s}^{-1}$  (equating to  $[O_3] \sim 10 \text{ ppb}$  under non limiting  $g_s$ ) reduces the sum of accumulated flux by  $\sim 43$  % as compared to not having a threshold. The inclusion of a flux threshold acts to shorten the daily period over which  $O_3$  damage occurs (Fig. 5b). Lower  $g_s$  in the morning and late afternoon, combined with lower  $[O_3]$  during these periods, leads to  $O_3$  fluxes that fall below the  $2 \text{ nmol m}^{-2} \text{ s}^{-1}$ , despite midday  $O_3$  fluxes that are often above this threshold. The influence of this threshold value can also be compounded over time when using dynamic vegetation models as is shown when modelling annualized production losses. Specifically, when considering no threshold ( $POD_0$ , Fig. A12), yield losses were predicted to be approximately three times larger in south-central Brazil compared to when using  $POD_2$  (Fig. 4). Care is therefore needed to ensure the biological relevance of metrics applied to ensure an accurate representation of potential  $O_3$  damage.

The  $O_3$  damage scheme used in global DGVM's such as JULES is by necessity simplistic, representing productivity decreases via an instantaneous reduction in both  $A_{net}$  and concomitant  $g_s$ . It therefore does not account for the potential decoupling of photosynthetic capacity and  $g_s$  under high  $[O_3]$  (Cernusak et al., 2021; Li et al., 2021), the observed acceleration of leaf senescence under high  $[O_3]$  (Gielen et al., 2007), nor the cumulative impacts of progressive and increasing root biomass decline under high  $[O_3]$  (Fig. A9, see also Grantz and Vu (2009)). For perennial crops such as sugarcane, often grown with multiple ratoon cycles, it is reasonable to assume a cumulative change in partitioning might translate into further reductions in productivity over, for example, a 5-year cycle.

The integration of more biologically realistic  $O_3$  damage schemes into dedicated crop models (i.e. DSSAT/CANEGRO, APSIM-Sugarcane) has previously been identified as a bottleneck to the development of





**Fig. 5.** Daytime (06:00 to 20:00) hourly  $O_3$  exposure and calculated stomatal fluxes of  $O_3$  into sugarcane modelled using JULES across south-central Brazil. Model outputs represent only control scenario for clarity (given negative feedback of  $O_3$  damage on  $g_s$  in current  $O_3$  scheme as used in JULES). (a) shows all terrestrial grid cells with grid cell identity represented by different colors, in (b) results of a single grid cell ( $-20.625, 310.3125$ , chosen as it contains the highest fraction of land cover dedicated to sugarcane production in south-central Brazil) with hour represented by color. Thresholds of fluxes equal to  $2 \text{ nmol m}^{-2} \text{ s}^{-1}$  (dotted red line) and concentration equal to 40 ppb (dashed red line) indicated.

accurate predictions on future  $O_3$  yield impacts (Emberson et al., 2018). However, despite the extensive resources marshalled through international efforts such as the AgMIP program (<https://agmip.org/4309-2/>) we are only now beginning to see the inclusion of  $O_3$  damage in this fashion into the modelling of highly studied temperate crops such as winter wheat (Feng et al., 2022). Our results highlight the likely impact of current  $O_3$  on sugarcane production, as well as the need to account for genotypic variation in functional traits that can determine  $O_3$  susceptibility (Wedow et al., 2021). The development of modelling-frameworks that can account for this variation (both between cultivars and during ontogenetic development), and translate impacts of NPP decline into changes in sugarcane quality (i.e. sucrose and fibre content) may provide more accurate estimates of the economic costs of  $O_3$ . This will also allow insights into how potential  $O_3$  mediation strategies, such as farm management practice and the use of new cultivar selection, can be used to close the  $O_3$  yield gap (Emberson et al., 2018).

#### CRedit authorship contribution statement

**Alexander W. Cheesman:** Conceptualization, Methodology, Formal analysis, Investigation, Writing – original draft, Visualization, Supervision, Project administration, Funding acquisition. **Flossie Brown:** Methodology, Investigation, Formal analysis, Writing – review & editing. **Nahid Farha:** Investigation. **Thais M. Rosan:** Resources. **Gerd Folberth:** Writing – review & editing, Funding acquisition. **Felicity Hayes:** Conceptualization, Writing – review & editing, Funding acquisition. **Barbara B. Moura:** Resources, Writing – review & editing, Funding acquisition. **Elena Paoletti:** Resources, Writing – review & editing, Funding acquisition. **Yasutomo Hoshika:** Resources, Writing – review & editing, Funding acquisition. **Colin P. Osborne:** Writing – review & editing, Funding acquisition. **Lucas A. Cernusak:** Resources, Writing – review & editing, Funding acquisition. **Rafael V. Ribeiro:** Conceptualization, Writing – review & editing, Funding acquisition. **Stephen Sitch:** Conceptualization, Writing – review & editing, Supervision, Funding acquisition.

#### Declaration of competing interest

The authors declare that they have no known competing financial

interests or personal relationships that could have appeared to influence the work reported in this paper.

#### Data availability

Data will be made available on request.

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#### Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.scitotenv.2023.166817>.

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