

NUMERICAL DATA

Effect of elevated atmospheric CO₂ concentration on growth and physiology of wheat and sorghum under cadmium stress (2018)

Effect of [CO₂] and [Cd] treatments on leaf area, shoot and root DW (dry weight) of wheat and sorghum plants.

Shoot/Root ratio	RDW (g)	SDW (g)	SLA	LA (cm ²)	C _{Cd} (mg/kg)	[CO ₂]	Plant species
2.20b	1.62j	3.54f	228.2h	435.8g	0	Ambient	Wheat
1.94e	1.76b	3.42g	214.1l	383.2i	10		
2.00de	1.22m	2.44k	226.1i	343.7k	20		
1.66f	1.13n	1.90n	234.8f	284.1n	40		
2.43a	1.84h	4.50d	210.1m	506.3e	0	Elevated	
2.05cd	2.06f	4.23e	200.1n	442.1f	10		
2.11bc	1.40l	2.90i	216.7k	390.1h	20		
1.74f	1.26m	2.20l	231.9g	326.9l	40		
1.97de	2.75b	5.42b	214.8l	569.4c	0	Ambient	Sorghum
2.04cd	2.26d	4.63c	246.2d	551.6d	10		
1.42g	1.91g	2.72j	217.9k	300.7m	20		
1.25h	1.46k	1.83n	256.6c	241.2o	40		
2.02cde	3.04a	6.17a	237.7e	715.5a	0	Elevated	
2.20b	2.45c	5.40b	265.8b	693.9b	10		
1.50g	2.16e	3.22h	221.6j	361.2j	20		
1.30h	1.60j	2.03m	274.7a	285.7n	40		

*C_{Cd} : Cd concentration, LA: leaf area, SLA: specific leaf area, SDW: shoot dry weight, RDW: root dry weight in each column different letters show significant differences (P < 0.05) between treatments

For wheat and sorghum at both CO₂ levels, increases in Cd concentration ([Cd]) decreased all the indices of plant growth, i.e. leaf area, and shoot and root DW, whereas at the same [Cd], e[CO₂].

Source: <https://www.tandfonline.com/doi/full/10.1080/00103624.2018.1547388>

Atmospheric CO₂ enrichment effect on the Cu-tolerance of the C₄ cordgrassT *Spartinadensiflora* (2018)

Photosynthetic pigments concentrations (µg/g) and DES state in randomly selected, fully expanded penultimate leaves of *Spartinadensiflora* in response to treatment with a range of Cu concentrations at 400 and 700 ppm CO₂ after 30 d of treatment. Values represent mean ± SE, n = 5. Different letters indicate means that are significantly different from each other (LSD, P < 0.05).

[Cu] (mM)	[CO ₂] (ppm)	Chl a	Chl b	Phe a	b-carotene	Lutein	Neoxanthin	Violaxanthin	Zeaxanthin	DES
0	400	242.5±19.2 ^a	91.5±6.7 ^a	14.1±2.3 ^a	5.8±0.5 ^a	20.1±4.1 ^a	8.9±1.4 ^a	3.8±1.1 ^a	6.4±0.9 ^a	0.60±0.05 ^a
	700	200.4±17.4 ^a	82.6±5.1 ^a	15.5±2.5 ^a	5.6±0.4 ^a	17.8±2.4 ^a	7.1±1.2 ^a	4.5±0.9 ^a	5.9±1.1 ^a	0.52±0.04 ^a
15	400	162.6±6.5 ^b	51.4±1.9 ^b	5.7±0.2 ^b	4.5±0.2 ^a	15.3±0.8 ^a	7.1±0.4 ^a	3.6±0.2 ^a	4.7±0.3 ^a	0.42±0.01 ^a
	700	174.2±11.5 ^b	44.4±1.6 ^b	8.8±2.4 ^b	5.9±1.4 ^a	15.4±2.9 ^a	5.5±1.1 ^a	6.7±2.2 ^a	5.9±1.2 ^a	0.49±0.06 ^a
45	400	167.3±8.7 ^b	54.0±3.1 ^b	4.9±0.5 ^b	5.1±0.3 ^a	15.9±0.9 ^a	6.8±0.6 ^a	3.6±1.2 ^a	5.1±0.3 ^a	0.41±0.07 ^a
	700	166.5±7.4 ^b	58.3±4.5 ^b	7.8±1.2 ^b	5.1±0.4 ^a	15.3±1.8 ^a	6.6±0.2 ^a	5.4±1.3 ^a	5.5±0.4 ^a	0.51±0.04 ^a

De-Epoxidation State (DES)

$$DES = \frac{[Antheraxantin] + [Zeaxanthin]}{[Violaxanthin] + [Antheraxantin] + [Zeaxanthin]}$$

Chl a, Chl b and Phe a concentrations decreased in presence of Cu in similar degree in both atmospheric CO₂ concentrations treatment (Two-way ANOVA: Cu, p < 0.05), but without statistical differences between Cu treatments. Contrarily the concentrations of each specific carotenoids and DES state did not vary with Cu and CO₂ concentrations treatments.

Source: <https://www.sciencedirect.com/science/article/pii/S0176161717302808>

Atmospheric CO₂ enrichment effect on the Cu-tolerance of the C₄ cordgrass *Spartinadenisiflora* (2018)

Total Cu, Ca, K, Mg and P concentrations for leaves and roots of *Spartinadenisiflora* in response to treatment with a range of Cu concentrations at 400 and 700 ppm CO₂ for 30d.

Tissue	[Cu] (mM)	[CO ₂] (ppm)	Cu (mg Kg ⁻¹)	Ca (mg g ⁻¹)	K (mg g ⁻¹)	Mg (mg g ⁻¹)	P (mg g ⁻¹)
Leaves	0	400	4.9± 1.0 ^a	3.4± 0.3 ^a	25.4± 0.2 ^a	3.3± 0.2 ^a	2.6± 0.1 ^a
		700	5.4± 0.2 ^a	3.1± 0.2 ^a	24.3± 0.3 ^a	2.9± 0.1 ^a	2.9± 0.1 ^a
	15	400	179.6± 2.0 ^b	3.1± 0.2 ^a	17.8± 0.4 ^b	2.9± 0.1 ^a	1.6± 0.2 ^b
		700	248.9± 6.3 ^{bc}	3.1± 0.2 ^a	19.2± 0.4 ^b	2.7± 0.1 ^a	1.9± 0.2 ^b
	45	400	737.5± 4.0 ^d	3.6± 0.1 ^a	22.1± 0.6 ^{ab}	3.4± 0.4 ^a	2.6± 0.2 ^a
		700	291.4± 4.9 ^c	2.9± 0.4 ^a	19.6± 1.1 ^b	3.0± 0.2 ^a	2.1± 0.1 ^{ab}
Roots	0	400	7.6± 0.5 ^a	2.4± 0.1 ^a	10.5± 0.3 ^a	1.4± 0.1 ^a	1.9± 0.2 ^a
		700	8.2± 1.4 ^a	2.2± 0.1 ^a	15.3± 1.0 ^a	1.7± 0.3 ^a	2.7± 0.1 ^b
	15	400	715.8± 11.0 ^b	1.9± 0.1 ^b	7.8± 0.5 ^b	1.1± 0.2 ^a	1.3± 0.2 ^a
		700	581.3± 9.0 ^c	1.6± 0.2 ^b	6.0± 0.3 ^b	1.0± 0.2 ^a	1.0± 0.5 ^a
	45	400	588.5± 3.0 ^c	1.5± 0.2 ^b	7.3± 0.4 ^b	1.2± 0.2 ^a	1.5± 0.2 ^a
		700	627.7± 12.0 ^c	1.5± 0.3 ^b	8.6± 0.5 ^a	1.1± 0.2 ^a	1.8± 0.4 ^a

Values represent mean ± SE, n = 6. Different values indicate means that are significantly different from each other (Two-way ANOVA, p < 0.05)

Source: <https://www.sciencedirect.com/science/article/pii/S0176161717302808>

Effect of elevated atmospheric CO₂ concentration on growth and leaf litter decomposition of *Quercus acutissima* and *Fraxinus rhynchophylla* (2017)

Comparison of the growth parameters of *Q. acutissima* and *F. rhynchophylla* in the ambient air (380 ppm) and elevated CO₂ (700 ppm) chambers.

Plant growth parameters	<i>Q. acutissima</i>			<i>F. rhynchophylla</i>		
	Ambient air	Elevated CO ₂	<i>p</i>	Ambient air	Elevated CO ₂	<i>p</i>
Total dry weight (g)	14.77±2.06	15.35±3.59	0.892	29.5±4.60	28.9±1.71	0.914
S/R	0.51±0.009	0.35±0.031	0.003	1.16±0.06	0.98±0.16	0.307
LWR	0.16±0.002	0.14±0.004	0.001	0.11±0.02	0.13±0.008	0.028
Thickness (mm)	0.09±0.002	0.11±0.003	0.002	0.13±0.002	0.15±0.01	<0.001
Leaf area (cm ²)	34.90±2.30	32.23±3.14	0.518	34.74±2.48	24.23±2.79	0.030
SLA (cm ² g ⁻¹)	235.1±8.79	176.7±5.93	0.001	241.1±5.96	181.9±2.19	<0.001

Values are means ± SE.

* Shoot - root ratio (g g⁻¹) = total shoot weight (g) / total root weight (g)

* Leaf weight ratio (g g⁻¹) = total leaf weight (g)/ total plant weight (g)

* Specific leaf area (cm² g⁻¹) = total leaf area (cm²)/ total leaf weight

The growth of *Q. acutissima* and *F. rhynchophylla* did not statistically differ between the ambient air and elevated CO₂ chambers. However, the shoot/root (S/R) ratio differed between the two species. The S/R ratio of *Q. acutissima* was significantly lower in elevated CO₂ chamber. Leaf growth conspicuously differed between the conditions. The thickness of the leaf blade was significantly higher (22% for *Q. acutissima* and 15% for *F. rhynchophylla*) in the elevated CO₂ chamber. The leaf area of *F. rhynchophylla* litter was significantly lower in elevated CO₂ chamber. And the specific leaf area (SLA) of the leaf litter of *Q. acutissima* and *F. rhynchophylla* were significantly lower at the higher CO₂ concentration.

Source: <https://journals.plos.org/plosone/article/file?id=10.1371/journal.pone.0171197&type=printable>

Effects of elevated atmospheric CO₂ on tree mycorrhiza associations (2016)

Species	Growing conditions	%Info	Total	Other effects on mycorrhizae	Root response
<i>Betula alleghaniensis</i>	Mesocosms (700 l l ⁻¹), growing in competition with <i>Betula papyrifera</i>	↑			Mass ↑, length ↑
<i>Betula papyrifera</i>	Mesocosms (700 l l ⁻¹), growing in competition with <i>Betula alleghaniensis</i>	↑	↑		Mass =↑, length =
<i>Betula papyrifera</i>	Pots in GC (700 l l ⁻¹)			Altered morphotype assemblages, extraradical hyphal length	Mass ↑
<i>Betula pendula</i>	OTC (700 l l ⁻¹), no fertilizer	↑		Altered species composition	Mass
<i>Liriodendron tulipifera</i>	24 weeks, pots in GC (•150 and •300 μ l ⁻¹)	=			Mass
<i>Liriodendron tulipifera</i>	OTC (•150 and •300 μ l ⁻¹)	=			Mass ↑
<i>Pinus caribaea</i>	49 weeks, pots in GC (660 l l ⁻¹)	=			
<i>Pinus echinata</i>	41 weeks, pots in GC	=		Signif. # in % Inf. at 34	Mass ↑
	(double CO#), no fertilizer			weeks, not at @nal harvest ↑	
<i>Pinus echinata</i>	24 weeks, pots in GC (double CO#)	=		Signif. In % Inf. at 6 weeks, not at @nal harvest	Mass

<i>Pinus palustris</i>	Pots in OTC (720 l l-")	↑	No changes in morphotype assemblages, effect larger at low N and adequate water	Length
<i>Pinus ponderosa</i>	OTC (•175 and •350 l l-")		↑ Extraradical fungal hyphae↑,, Mycorrhizal turnover	Area density ↑
<i>Pinus ponderosa</i>	Pots in GC (700 l l-")	↑	↑ Density ↑	
<i>Pinus radiata</i>	49 weeks, pots in GC (660 l l-")	=		
<i>Pinus strobus</i>	Pots in GC (700 l l-")	↑	Altered morphotype assemblages	Mass ↑
<i>Pinus syl.estrus</i>	Pots in GC (700 l l-")	↑	Altered morphotype assemblages	↑
<i>Pinus syl.estrus</i>	120 d, pots in GC (double CO#)	=	No effect on total fungal mass	
<i>Pinus taeda</i>	Open bottom pots in OTC (700 l l-")	=		↑
<i>Populus tremuloides</i>	Open bottom pots in OTC (700 l l-")	=	Extraradical mycorrhizal hyphal	Mass
length # under N-poor conditions, \$ under N-rich conditions				
<i>Populus hybrids</i>	2 years, OTC (-350 l l-")	AM =		Mass ↑
<i>Quercus alba</i>	24 weeks, pots in GC (double CO#)		Increased↑ mycorrhizal infection before increase in root mass, alterations in species abundance	Mass ↑
<i>Quercus alba</i>	OTC (•150 and •300 l l-")	↑		Mass ↑
<i>Tsuga canadensis</i>	Pots in GC (700 l l-")	EM = AM ↑		Mass ↑

Experimental (growing) conditions, response of the percentage of mycorrhizal infection (%Inf), of total amount of mycorrhizae, as well as the root response are reported for each study. EM, ectomycorrhizae; AM, arbuscular mycorrhizae; -, no significant changes ($P \geq 0.05$); , significantly enhanced ($P < 0.05$).

Source: R. Ceulemans et al. (2016), Effects of CO₂ Enrichment on Trees and Forests: Lessons to be Learned in View of Future Ecosystem Studies, Annals of Botany

Effects of elevated CO₂ on tree litter decomposition (2016)

Tree species and tissue	Plant growth conditions	Decomposition conditions	Response of litter quality	Response of decomposition
<i>Acer pseudoplatanus</i> , senesced leaves	1 season, 600 μ l-", pots in solar domes	8 months, chambers	C / N , lignin / N	=
<i>Betula pubescens</i> , senesced leaves	1 season, 600 μ l-", pots in solar domes	5 months, chambers	C / N , lignin / N	↓
<i>Betula pubescens</i> , live roots <2 mm	1 season, 600 μ l-", pots in solar domes:	3 months, Chambers with soil		=
<i>Castanea sati.a</i> , senesced leaves	non-fertilized 2 years, 700 ll l-", pots in GC	6 months, chambers: Incomplete decomposer community complex decomposer community	C / N C / N C / N , lignin = C / N , lignin =	↓
<i>Fraxinus excelsior</i> , senesced leaves	1 season, 600 ll l-", pots in solar domes		C / N , lignin / N	
<i>Liriodendron tulipifera</i> , senesced leaves	1 season, •300 ll l-", pots in GC, exposed to ozone		N , lignin	↓
<i>Liriodendron tulipifera</i> , senesced leaves	2 years, •150 and -300 ll l-", OTC		C / N , lignin / N	↓
<i>Picea sitchensis</i> , senesced needles	1 season, 600 ll l-", pots in solar domes		C / N , lignin / N	=
<i>Picea sitchensis</i> , live roots !2 mm	1 season, 600 ll l-", pots in solar domes: 3 months, chambers with soil fertilized non-fertilized	3 months, chambers with soil	C / N C / N	=

Tree species and tissue, fumigation conditions and duration, decomposition conditions and duration, and changes in litter chemistry and decomposition rates are reported for every study. OTC, Open top chambers; GC, growth chambers; -, no significant changes ($P \geq 0.05$); #, significantly enhanced ($P \leq 0.05$); \$, significantly decreased ($P \leq 0.05$).

Source: R. Ceulemans et al. (2016), Effects of CO₂ Enrichment on Trees and Forests: Lessons to be Learned in View of Future Ecosystem Studies, Annals of Botany