

Fig A – Blume et al.

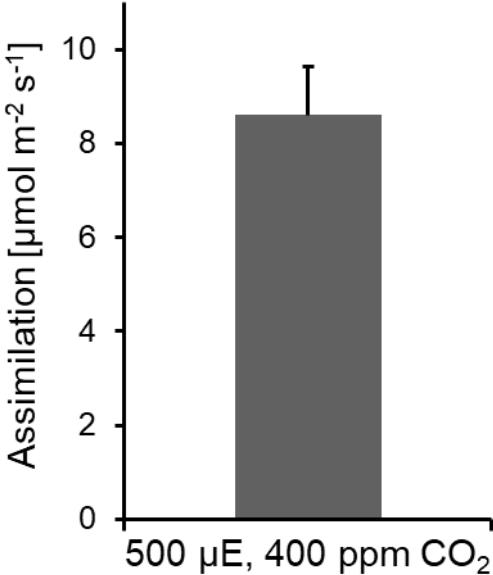


Fig A. CO₂ assimilation of 5-week-old *Arabidopsis* plants. Net CO₂ assimilation was recorded under ambient CO₂ conditions (400 ppm) and light saturating conditions (500 μE).

Fig B – Blume et al.

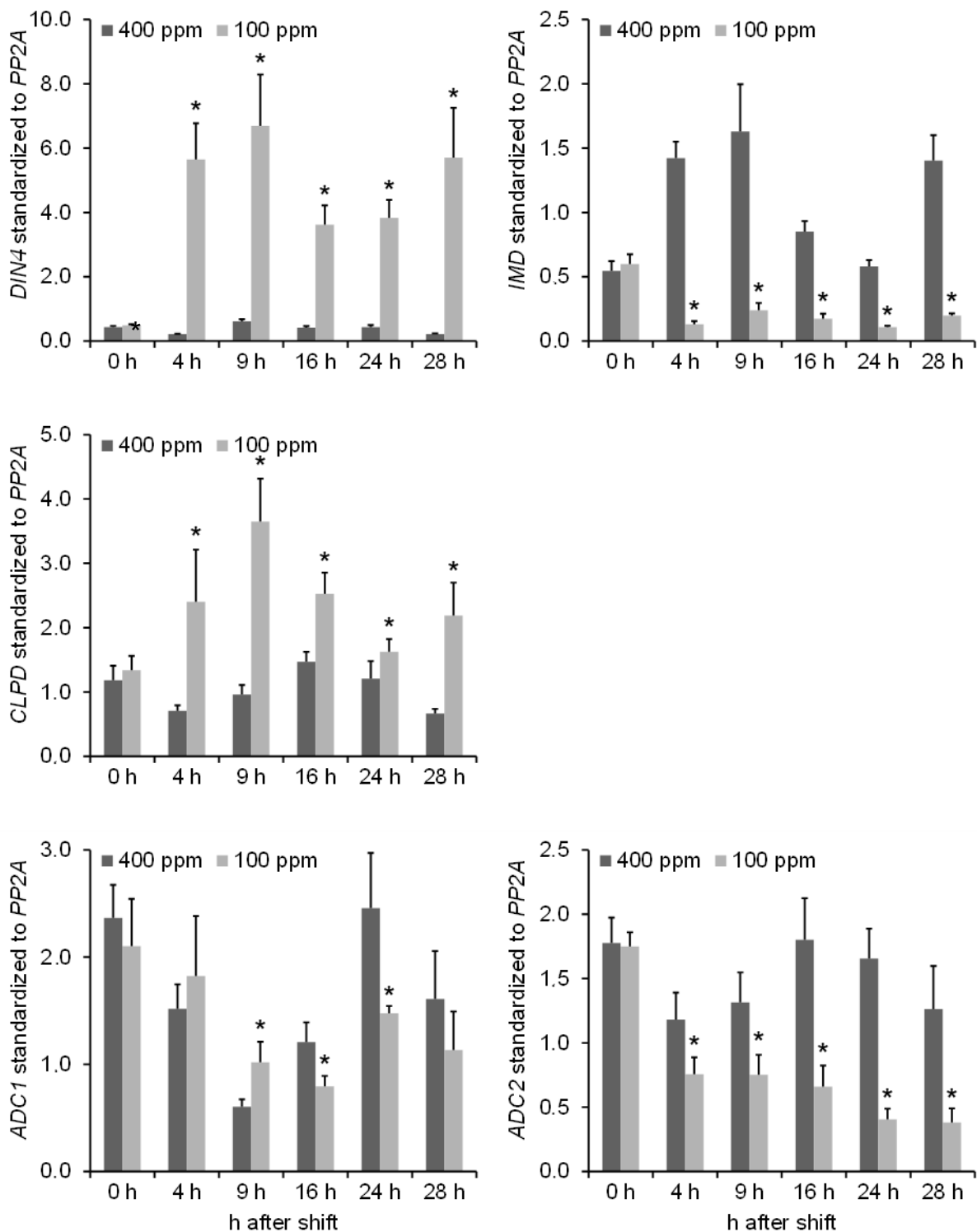


Fig B. Transcript levels of selected genes of leucine synthesis (*IMD*) and degradation (*DIN4*), protein degradation (*CLPD*), and polyamine synthesis (*ADC1*, *2*) at 400 ppm and 100 ppm CO₂ standardized to *PP2A*. Data are the mean of five biological replicates \pm SD. Significance was tested according to the two-tailed Student's t-test; values were changed to log₂ ratios to allow this test (* $p < 0.05$) *DIN4*, Dark-induced 4; *IMD*, Isopropylmalate dehydrogenase; *CLPD*, caseinolytic protease D; *ADC*, arginine decarboxylase.

Fig C – Blume et al.

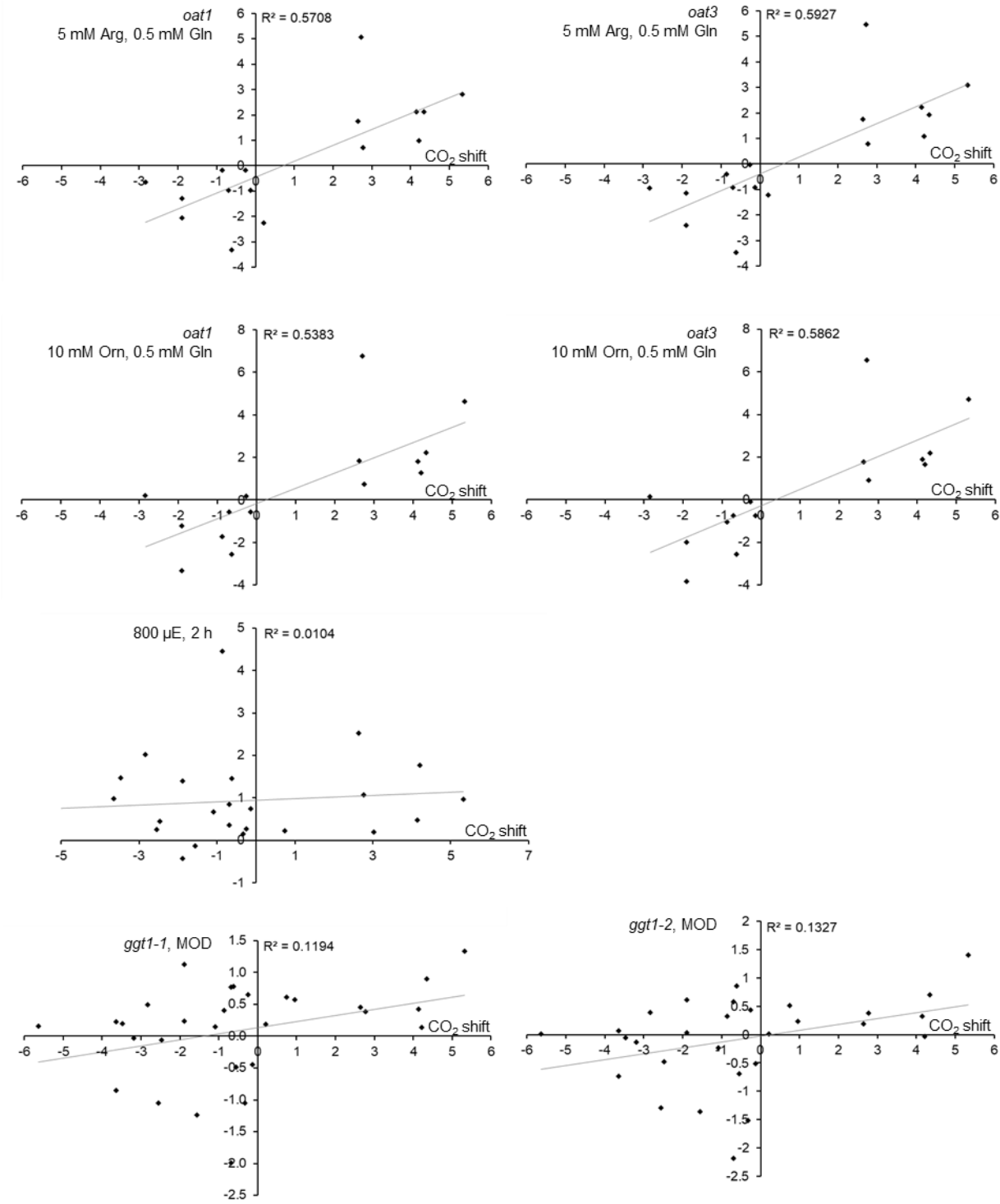


Fig C. Correlation of log₂ metabolite changes between the low CO₂ shift and in response to different stresses/different *Arabidopsis* mutants. The fold changes are given in Fig 6.

Fig D – Blume et al.

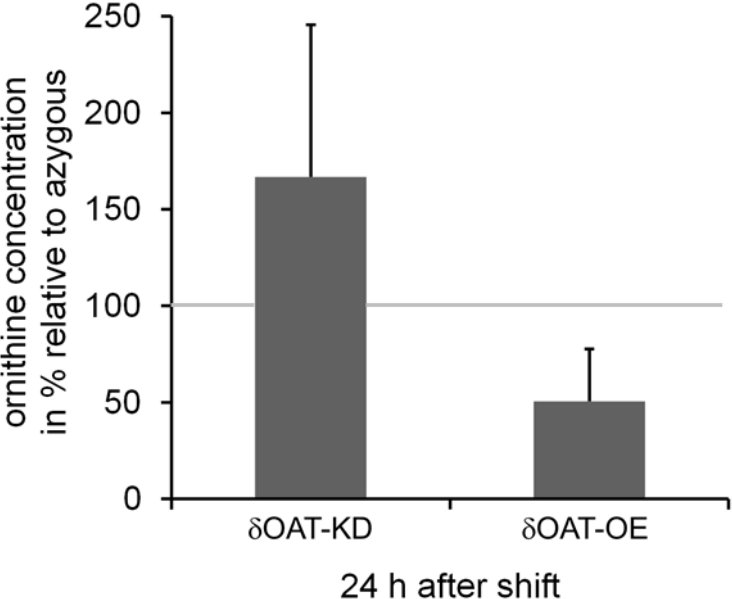


Fig D. Concentration of ornithine in knockdown (KD) and overexpression (OE) lines of δOAT after a shift from 400 ppm to 100 ppm CO₂ relative to azygous plants. Plant material was harvested 24 h after shifting the plants to low CO₂ concentrations. Data are the mean of three (KD) or two (OE) biological replicates \pm SD. Grey line: wild type δOAT level.