## The influence of atmospheric carbon dioxide concentration on the carbon isotope composition of plant tissues

BRIAN A. SCHUBERT<sup>1\*</sup> AND A. HOPE JAHREN<sup>2</sup>

<sup>1</sup>University of Louisiana at Lafayette, School of Geosciences, Lafayette, LA 70504

(\*correspondence: schubert@louisiana.edu)

<sup>2</sup>University of Hawaii at Manoa, School of Ocean and Earth Science and Technology, Honolulu, HI 96822 (jahren@hawaii.edu)

Many environmental controls influence the ratio of <sup>12</sup>C:<sup>13</sup>C fixed within plant tissue. However, how the concentration of atmospheric carbon dioxide, a raw material for photosynthesis that affects many aspects of plant biology, affects the net isotopic discrimination between plant tissue and atmospheric CO2 has remained uncertain. Here we present a relationship quantifying how atmospheric  $CO_2$  concentration ( $pCO_2$ ) affects plant carbon isotope fractionation that is based on chamber and field studies conducted on a great diversity of C<sub>3</sub> plants. This relationship reconciles the wide range of fractionation factors previously reported and provides the framework for applying the  $pCO_2$  effect to periods of changing  $pCO_2$  level. Our analysis provides a minimum value for the fractionation due to catalysis by RuBisCO, which has implications for reconstructing plant water-use efficiency across any interval of  $pCO_2$  change. Our work also helps to explain why terrestrial substrates commonly show a larger carbon isotope excursion than marine substrates during intervals of rapid changes in the global carbon cycle, and can be used to quantify the background and maximum  $pCO_2$  levels for these events. We conclude that the effect of changing  $pCO_2$  level on plant carbon isotope fractionation must be accounted for when analysing  $\delta^{1\bar{3}}C$  records from terrestrial substrates across any interval of  $pCO_2$  change.