



# PAST, PRESENT AND FUTURE OF REDWOODS: A Redwood Ecology & Climate Symposium



Photo by Anthony Ambrose

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## Research Abstracts – Keynote Talks

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### Current and Future Trends in Coast Redwood Climate

Healy Hamilton, Senior Research Fellow, Marine Conservation Institute

Today, coast redwood forests occupy a relatively cool, moist, and narrow sliver of coastal California. This limited distribution suggests that coast redwood forests have a narrow envelope of climatic conditions within which they can compete. Global climate change may alter the range of conditions for temperature and moisture that support healthy coast redwood forests. As part of the RCCI, we used several approaches to identify areas of high climate stability and change across the range of coast redwoods, both now and into the future. We investigated contemporary patterns of significant climate change in the coast redwood region, and predicted where those changes may be greatest in the century ahead. Using 115 years of weather station data, we compared seasonal temperature and precipitation trends in recent decades against a 20<sup>th</sup> century baseline. We found significant changes in climate patterns are already occurring across the coast redwood range. Increases in spring minimum temperatures (fewer cold spring days) were identified as the most salient current trend in coast redwood forests. Total precipitation has not yet significantly changed, but remains highly variable year-to-year. The southernmost portion of redwood forest in Santa Cruz County is already experiencing minimum temperature increases in all four seasons, while redwood forests in Humboldt and Del Norte counties are currently demonstrating climate stability.

To investigate how climate might change throughout the coast redwoods range in the future, we used a climate analogue approach. We determined historical occurrences of anomalous weather years, including individual years of extreme heat, cold, rainfall, drought and combinations thereof. These past conditions were used as input into species distribution models, to map the distribution of coast redwood bioclimate under extreme conditions that may approach future climates. The results suggest patterns of stability and change similar to observed trends. Under a wide range of potential future climate conditions, almost the entire distribution of Humboldt and Del Norte county redwood forest climate remains stable. The most severe departures from normal conditions occur in Santa Cruz and Monterey regions of coast redwood range under futures that are warmer and drier. While we do not yet know how coast redwood forests and their many associated species will respond to these current or projected climate changes, our results suggest where the most severe changes are likely to occur. These findings can direct future field-based monitoring of coast redwood trees in these climate-risk zones. Citizen science participation to refine coast redwood distribution at fine spatial scales, particularly at the edges of their range, could significantly improve our understanding of the climate tolerance of coast redwoods.

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### Millennium-scale crossdating of *Sequoiadendron giganteum* and *Sequoia sempervirens*

Allyson L. Carroll, Humboldt State University

Through RCCI, we created tree ring records from both *Sequoiadendron giganteum* (SEGI) and *Sequoia sempervirens* (SESE) dating back over 1000 years. This dendrochronological study (*i.e.*, analysis of tree rings) created a baseline, or ruler, upon which researchers can study tree growth, climate, fire history, and even date archeological structures. Here we present and

compare crossdated chronologies for SEGI and SESE at sites spanning their native ranges in California using samples collected at multiple heights along trunks of standing trees. This work represented the most successful and comprehensive crossdating effort for SESE with 8 sites, 76 trees, 864 radii (*i.e.*, rings measured from the oldest wood to the youngest), and 250,530 growth rings encompassing the years 328 to 2012 AD. We also contributed to the catalog of work on SEGI that was originally started by the pioneer of dendrochronology A.E. Douglass in the early 1900s by adding chronologies from 5 sites, 44 trees, 602 radii, and 233,182 growth rings encompassing the years 474 to 2012 AD. SEGI yielded very obvious ring patterns, while SESE rings were more difficult to crossdate. For both species, site-level chronologies were created for crossdating and reflected reliable marker years (*i.e.*, consistent narrow or large rings or patterns thereof). SEGI site-level chronologies generally showed common synchrony while SESE site synchrony grouped along the north-south gradient. For SESE, common signals occurred within northern and southern chronologies with mid-range sites providing a bridge for crossdating. The southernmost SESE sites (Landels-Hill Big Creek Reserve and Big Basin) generally correlated more strongly with the SEGI sites than the northern SESE sites. Landels-Hill Big Creek Reserve captured the strongest drought signal of all sites. All SESE and SEGI chronologies shared some marker years that were often strong regional drought events such as the one reflected in the narrow ring width for 1924. Many other notable marker years (*e.g.*, fire signals at Landels-Hill and Montgomery Woods, 1739 growth reduction for northern SESE, 1580 missing ring for SEGI) emerged from this work and are related to past environmental and climatic events, underscoring the many applications of crossdated chronologies.

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### **Separating effects of tree size and age on trunk growth in California redwoods**

Stephen C. Sillett, Humboldt State University

*Sequoia sempervirens* (SESE) and *Sequoiadendron giganteum* (SEGI) are the tallest and largest trees with incredible capacity to sequester carbon in decay-resistant wood and fire-resistant bark, yet little is known about how crown structure, growth rates, and allocation of aboveground wood production change as they age. We climbed, mapped, and sampled trees representing the full size and age ranges of both species within 16 RCCI plots and nearby old-growth forests. A total of 137 trees were mapped, which involved intensive measurements of main trunks and appendages. A 3-D model based on the complete set of measurements was used to error-check each tree's data prior to final calculations of all aboveground quantities. From nearly 20 thousand mapped branches, we dissected 257 spanning the range of heights and diameters to develop predictive equations. Applying these to the complete inventory of branches yielded accurate whole-tree estimates of leaves as well as bark, cambium, sapwood, and heartwood on branches. A total of 1462 increment cores (*i.e.*, thin cylindrical samples) encompassing 458213 annual rings were collected at multiple heights on trunks to estimate minimum tree ages and to quantify bark, cambium, sapwood, heartwood, and annual growth of trunks. To be comprehensive, we measured numerous superlative trees, including the tallest known of each species (SESE = 115.72 m, SEGI = 96.29 m). The largest SESE was at least 1450 years old and had 424 Mg dry mass, 1103 m<sup>3</sup> total volume, 5467 m<sup>2</sup> cambium area, 9549 m<sup>2</sup> leaf area, and 1.12 billion leaves. A 2520-year-old tree became the oldest SESE known (by 300 years). The largest SEGI was at least 3240 years old and had 550 Mg dry mass, 1512 m<sup>3</sup> total volume, 5978 m<sup>2</sup> cambium area, 6726 m<sup>2</sup> leaf area, and 1.94 billion leaves. The 137-tree dataset enabled us to develop equations for accurately predicting aboveground quantities of any redwood in an old-growth forest using simple measurements obtainable from

the ground (e.g., trunk diameter at breast height, crown volume), which will be critical for future research and long-term monitoring. Annual ring width declined with age in both species, averaging 1 mm in SESE by 1700 years and SEGI by 1000 years. Despite declining ring widths, annual rates of main trunk wood volume growth increased with size through old age in both species. The fastest growing tree was a 108.6-m-tall, 1180-year-old SESE whose main trunk produced 1.61 m<sup>3</sup> of wood annually during the last decade. Rates of heartwood production also increased with age in both species, averaging 90 % of the annual wood volume increment in SESE by 1100 years and SEGI by 1600 years. Mass growth of SESE and SEGI was comparable for the first 650 years when aboveground dry mass of both species averaged 50 Mg, but by 1000 and 2000 years an average SEGI had accumulated only 89 and 75 % of an average SESE's dry mass (106 and 348 Mg, respectively). In both species, tree size and crown structure were much stronger determinants of trunk growth than age *per se*, and trees retained high annual rates of carbon sequestration through old age. Regardless of tree age, annual rates of wood production increased during the last century in both species, and the pace of increase was unprecedented in our tree-ring record. Only one tree of each species (both relatively small) exhibited a significant decline in wood production during the last century, whereas declining growth following disturbance (e.g., fire) was more frequent in prior centuries. Aside from the obvious effect of size (*i.e.*, larger trees have more leaves to photosynthesize), causes for increasing rates of wood production during the last century may reflect climate change or other anthropogenic effects (e.g., fire suppression).

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## Redwood seedling responses to drought

Anthony Ambrose, U.C. Berkeley

Increasing drought severity and duration associated with climate change may significantly impact coast redwood and giant sequoia trees in the coming decades, especially young seedlings given their small stature and limited rooting depth. As part of the Redwoods and Climate Change Initiative, we studied the impact of drought on coast redwood and giant sequoia seedlings originating from seed stocks obtained from different geographic locations throughout each species' range (north, central, and south provenances). We imposed a 6-week period of severe soil water deficit followed by a 2-week period of re-watering in a controlled greenhouse experiment in order to assess how the two species, and different sampled provenances of those species, respond to drought. The single drought event had a significant effect on seedling growth and physiology, but most water-stressed seedlings of both species largely recovered their physiological functioning by the end of the 2-week recovery period. Importantly, the experiment revealed that seedlings of the two species exhibited contrasting drought-response strategies. Compared to coast redwood, giant sequoia seedlings showed higher leaf photosynthesis and transpiration rates, greater control over plant water status, and a lower degree of stem water transport impairment due to drought, as well as lower relative growth rates and lower growth sensitivity to drought. Drought-stressed coast redwood seedlings had 40% lower average relative growth rates compared to well-watered control seedlings over the entire experiment, while drought-stressed giant sequoia seedlings had 13% lower average relative growth rates compared to control seedlings. Drought-stressed coast redwood seedlings had 80% lower average photosynthesis rates and 88% lower average transpiration rates than control seedlings at the time of maximum drought, compared to 61% lower average photosynthesis rates and 85% lower average transpiration rates in drought-stressed versus control giant sequoia seedlings. Drought had a substantially greater effect on plant water status in coast redwoods

than giant sequoias, with 298% lower average daytime shoot water potential in drought-stressed coast redwood seedlings compared to controls at the time of maximum drought versus 157% lower average daytime shoot water potentials in drought-stressed giant sequoia seedlings compared to controls. The greater water stress levels experienced in coast redwoods was associated with significantly greater levels of stem embolism (i.e., air bubbles in water-conducting wood cells which prevent further movement of water in the affected cells) compared to giant sequoia seedlings. The two species showed contrasting whole-plant biomass allocation patterns independent of drought-effects, with a greater proportion of total plant biomass allocated to roots in giant sequoia and a greater proportion of total plant biomass allocated to stems in coast redwood. Notably, there were only minor response differences among the different provenances we tested, indicating that seed source location may not substantially influence drought effects in either species. Both species appear to have a threshold response to drought where seedling growth and physiology become significantly impaired as soil moisture levels decline to less than about 10-15% and plant water potentials decline to less than about -1.5 MPa. These results suggest that despite important species-level differences, both coast redwood and giant sequoia seedlings may experience significant levels of water stress and as a result reduced physiological performance and growth under drier and warmer conditions projected for California in the near future.

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## **Chemical signals of climate and physiology in redwoods**

Todd Dawson, U.C. Berkeley

It is well established that the chemical components (e.g., carbon and oxygen) that comprise the leaves and wood of any tree provide signals about tree function and about climate. Through the Redwoods and Climate Change Initiative, we used chemical signals in the leaves and wood of redwoods to extend what was learned from our measurements of physiology and tree ring sizes to explore how redwood trees have responded to environmental variation and change in both space and time. Analyses of leaf stable carbon isotope ratios for both coast redwood and giant sequoia from throughout their geographical ranges show species-level differences and a marked gradient with tree height but only minor differences among populations within each species. The vertical gradient is best explained by tree response to changes in both microenvironment and physiology that are known to change with tree height. In contrast, leaf stable oxygen isotope ratios for both species showed no clear trend with tree height but very clear and marked differences between populations with giant sequoia displaying a generally stronger inferred leaf-level response to the higher summer evaporative conditions present in the Sierra Nevada Mountains as compared to the coast. Both species showed population level differences in oxygen isotope ratios with the driest and warmest sites most distinct from the others. Detailed analyses of the stable carbon and oxygen isotope ratios in individual (annual) tree rings were also used to explore how climate and tree response to climate was recorded for both redwood species. Wood processed from two periods in a single year (spring and summer) and for each year over a 21-year continuous record where climate data for each location was recorded revealed unique (local) climatic conditions and tree responses to climate for each species and population. Isotope variation was most pronounced in the tree ring record and corresponded to (1) drought (e.g. 1976/1977), (2) above-average temperatures, (3) growing season evaporative conditions, and (4) above-average precipitation (e.g., 1982-1985). Isotope variation also appeared to be influenced by tree size and microsite conditions (e.g., streamside versus upslope locations). Having established the climate-physiology-isotope associations now allows us to develop a robust calibration for looking at periods when and places where climate records do

not exist. One very strong relationship across both species was how carbon and oxygen isotopes co-varied; based on a physiological model and real calibrations we can infer that during dry and also warm periods trees appear to first down-regulate their water use and secondly their carbon fixation and that high evaporative conditions drive some of the most marked changes in both variables. These new isotope data are permitting us to reconstruct the responses of redwoods to past and current environments like no other set of information can. Moreover, our analyses can be extended with the use of models to provide well-grounded predictions of redwood responses to future change as well. This information could also be useful for guiding future efforts to conserve and protect redwoods under the novel changes they have been experiencing since human impacts have increased.

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### **Establishment of long-term reference plots in old forests of both California redwood species, including a detailed mass, carbon, and structural analysis of coast redwood.**

Robert Van Pelt, Humboldt State University

The largest single effort to monitor old-forest responses to a changing environment throughout the ranges of both *Sequoia sempervirens* (SESE) and *Sequoiadendron giganteum* (SEGI) began as a Save the Redwoods League-funded project in 2009. At the core of this unprecedented leaf-to-landscape effort was the establishment of 16, 1-hectare (2.5-acre) plots distributed throughout the species' latitudinal ranges in which we took detailed measurements on forest structure and composition as a baseline to track changes over time. In each plot we conducted an inventory of all aboveground plant material by species, live and dead, at a resolution never before attempted. This involved measuring a tagged population of more than 6000 trees, a subsample for shrubs of more than 5200 individuals, and more than 1300 vegetation plots for estimates of seedlings and herbaceous vegetation. For the needed resolution on the two redwood species, more than 100 trees of all sizes and vigor were climbed and mapped to develop relationships for the stand-level quantification of biomass (leaf, bark, sapwood, heartwood, dead wood) and surface areas (leaf, bark, cambium). So far for the 11 SESE plots, similar biomass relationships have been developed for each species of tree, shrub, fern, herb, and woody debris that occurs in these forests. In addition, massive accumulations of dead wood, particularly in rain forests of Del Norte and Humboldt Counties, forced us to adopt a new classification system for coarse woody debris. We found that above-ground biomass in the northernmost SESE plots was the highest ever recorded, and the dead wood component was 2–3 times what had been previously reported in these or other forests. Stand basal area for live SESE exceeded 400 m<sup>2</sup> ha<sup>-1</sup> in one plot, and stand totals of 300–400 m<sup>2</sup> ha<sup>-1</sup> occurred in six others. When living vegetation was combined with standing dead and downed logs, the aboveground portion represented oven dry masses of 3500–5000 Mg ha<sup>-1</sup> (Mg ha<sup>-1</sup> = metric tons per hectare). Up to 25% of the total biomass was dead wood in the form of snags and logs. The largest component masses in SESE forests were wood and bark of live trees and the decaying wood of dead trees. Carbon content varied only slightly among these components (*i.e.*, 45–53%), so roughly half of the total forest mass was carbon, and by far the bulk of this was bound up in decay-resistant heartwood. Five of the northern plots had > 25 Mg ha<sup>-1</sup> of leaf dry mass, which is a global record, as is the accompanying 15–17 Leaf Area Index (m<sup>2</sup> of projected leaf area per m<sup>2</sup> of ground area). Long-term changes in these forests will be detected by annual surveys of growth and mortality, remeasurement twice per decade, the establishment of smaller plots in other landscape positions, and the linking of the ground-based research to LiDAR and other remote sensing techniques to allow detecting landscape-level change.

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## Research Abstracts – Poster Presentations

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### Poster 1

#### Inter-annual climate-radial growth relationships and regional synchrony of *Sequoia sempervirens* and *Sequoiadendron giganteum* tree-ring series

Allyson L. Carroll, Stephen C. Sillett, and Russell D. Kramer, Humboldt State University

Through RCCI we investigated inter-annual climate sensitivities and regional synchrony of crossdated chronologies of *Sequoia sempervirens* (SESE) and *Sequoiadendron giganteum* (SEGI) from sites across their native ranges. We described relationships between high-frequency radial growth variation and monthly climate variables of Palmer Drought Severity Index (PDSI), precipitation, maximum temperature, and minimum temperature for the period from 1895 to present using correlation and principal components analyses. For SESE, summer soil moisture (PDSI) correlated with radial growth, and this relationship extended into the spring months for southern sites. Also, southern sites were more limited by maximum temperatures than northern sites. In the case of SEGI, we confirmed the findings of Hughes *et al.* (1989, 1992) that rather than tracking a strong signal for temperature or precipitation, SEGI's smallest rings recorded extreme events. In addition to seeing this relationship expressed in summer PDSI correlations, we showed that June temperature also captured the association of drought with SEGI radial growth. The strongest relationship among all sites was between growing season soil moisture and growth at the southernmost SESE site, Landels-Hill Big Creek Reserve. In fact, SESE growth at Landels-Hill showed a strong relationship with state and regional drought. Examination of relationships between the high-frequency tree ring chronologies revealed that SEGI chronologies were highly associated with each other while the SESE chronologies separated more distinctly into northern and southern signals. While fog data were sparse and complex, inter-annual growth for northern SESE showed a negative correlation with summer "airport fog" (*i.e.*, 400 m ceiling height, Johnstone & Dawson 2010), which we considered a surrogate for cloudiness in low elevation forests. SESE growth in low elevation forests increased with decreasing summer cloudiness (see RCCI, Sillett *et al.*, "Annual rates of trunk wood production in old-growth redwood forests since 1750). The relationship between summer cloudiness and inter-annual growth generally weakened towards the southern end of the SESE range.

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### Poster 2

#### Annual rates of trunk wood production in old-growth redwood forests since 1750

Stephen C. Sillett, Robert Van Pelt, Russell D. Kramer, and Allyson L Carroll, Humboldt State University

Old-growth forests dominated by *Sequoia sempervirens* (SESE) and *Sequoiadendron giganteum* (SEGI) contain world-record biomass and store large quantities of carbon in trees with decay-resistant wood and fire-resistant bark, yet little is known about how annual growth rates vary in response to climate. As a first step toward understanding this relationship, we quantified wood production in SESE and SEGI trees from 16 RCCI plots and nearby old-growth forests. Increment cores (*i.e.*, thin cylindrical wood samples) were collected at multiple heights

on main trunks of 106 trees, and dendrochronology was used to crossdate annual rings, whose widths were measured to the nearest micron. The mapped taper of each tree's trunk was combined with predicted bark radii and measured ring widths to determine wood radii backwards in time and to calculate wood volumes for each year. Wood volume growth was determined by subtracting preceding-year volumes from subsequent volumes. Each tree's growth history was taken as far back as crossdated rings allowed. The longest whole-trunk records extended back to 1179 and 781 AD for SESE and SEGI, respectively. By 1750 AD, only 50 trees remained in the dataset, and replication was too low at most sites to continue further. Because rates of wood volume growth increased through old age in both species, we derived a size-detrended rate of wood production that did not increase merely because trunks enlarged over time. Despite a high degree of inter-annual variation in wood production corresponding to high-frequency climatic fluctuations, several longer-term trends were evident. Following a period of relatively rapid growth in the 1940s, growth was depressed in the 1950s and 1960s, surged in the 1970s, and has remained high in the 4 northernmost SESE forest plots (Jedediah Smith Redwoods State Park, Prairie Creek Redwoods State Park, Redwood National Park, Humboldt Redwoods State Park). Wood production since the surge in 3 of these forests (all but Prairie Creek RSP) was higher than at any time on record. No such growth surge was evident in SESE forests farther south except Samuel P. Taylor SP, though a trend of increasing growth during the 20<sup>th</sup> century was evident in all but the southernmost SESE forest (Landels-Hill Big Creek Reserve). Compared to SESE, there was less inter-annual variation in SEGI growth. A trend of increasing wood production during the 20<sup>th</sup> century was evident in all SEGI forests except Whitaker Forest, which was partially logged starting about 1870. Growth at Whitaker Forest surged following logging, returning to pre-logging rates by about 1910. Two other SEGI forests exhibited less pronounced growth surges, Mountain Home State Forest in the 1870s and Freeman Creek Grove in the 1880s. Overall, wood production of SESE forests was consistently higher than that of SEGI forests, except for the 1880s and late 1950s. Temporary release from competition following partial logging likely contributed to unusually high rates of SEGI wood production in the late 19<sup>th</sup> century. Heavy logging and slash burning of northern SESE forests might have contributed to unusually low rates of growth in the mid-20<sup>th</sup> century by obscuring sunlight. Gradually increasing rates of SEGI wood production during the 20<sup>th</sup> century could reflect extended growing seasons accompanying global warming. Declining cloud cover during the late 20<sup>th</sup> century probably contributed to the growth surge in northern SESE forests by increasing light availability. These possibilities and others (e.g., CO<sub>2</sub> fertilization) warrant further investigation.

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### **Poster 3**

#### **Structural development of redwood branches and its effects on wood growth**

Russell D. Kramer and Stephen C. Sillett, Humboldt State University

The heaviest forests in the world are those dominated by *Sequoia sempervirens* (SESE) and *Sequoiadendron giganteum* (SEGI). Branch contributions to above-ground mass growth are initially 10% and increase to 35% in the largest trees. Recent measurements of all branches from SESE and SEGI trees through the RCCI provide a new source of knowledge for within-tree carbon dynamics and branch structure. This study adds a finer scale of resolution to what we know about wood production and is important because branches represent a significant portion of redwood above-ground biomass. We dissected 31 branches from 8 SESE and 7 SEGI trees to identify the structural determinants of branch wood volume growth and develop a conceptual



model of branch development. Branch heights ranged from 15.6–104.5 m, diameters from 3.4–13.6 cm, ages from 11–258 years, and wood production from 3.4–80.3 cm<sup>3</sup> per year. The ratio of energetically expensive cambium to energy-supplying leaf area did not change with size or age but increased with height and light availability. The ratio of expensive heartwood depositional area to supplying leaf area increased with both size and age. Branch size, light availability, species, and heartwood area explained 87.5% of the variation in wood volume growth. After accounting for the positive effects of size and light, wood volume growth declined with both heartwood depositional area and age, but the age effect was trivial in comparison. These results suggest that age-related declines in wood volume growth may be caused by expansion of the heartwood sink, not age *per se*. The oldest branch (258 years) was also 57% heartwood by volume and produced the least wood relative to its size of any branch. Additionally, the negative heartwood effect is twice as strong in SEGI. Cone production correlated with reduced wood volume growth in SEGI branches after accounting for size and light availability even though the cones are long-lived and photosynthetic. The most fecund (i.e. had the most cones) SEGI branch had 48 % of its mass and 26 % of its photosynthetic area in cones, and 1/3 SEGI branches had > 30% of their mass in green cones. A reduction in growth with heavy reproductive output implies branches are not limited by their ability to support cones, but rather by their ability to provide for them. Thus, branch wood production in both species may be limited by competing internal carbohydrate sinks. We developed a conceptual model for branch development, based on a feedback system, where branch growth and structure interact to produce an array of appendage types, including limbs bearing reiterated trunks.

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#### **Poster 4**

#### **Establishing baseline reference conditions for old-growth redwood forests.**

Robert Van Pelt<sup>1</sup>, Stephen C. Sillett<sup>1</sup>, and Bill Kruse<sup>2</sup>

<sup>1</sup>Humboldt State University, <sup>2</sup>Kruse Imaging

The largest single effort to monitor old-forest responses to a changing environment throughout the ranges of *Sequoia sempervirens* (SESE) and *Sequoiadendron giganteum* (SEGI) began as a Save the Redwoods League-funded project in 2009. At the core of this unprecedented effort are 16 plots distributed throughout the ranges of both species in which we measured above-ground biomass, forest structure, and species composition. Eleven plots span the range of SESE from within a few kilometers of Oregon to south of Big Sur (670 km), and 5 plots span the more limited range of SEGI from Calaveras to the easternmost and one of the southernmost SEGI groves – Freeman Creek (279 km). Within each plot, a wide variety of techniques were used to obtain accurate stand-level estimates of biomass (leaf, bark, sapwood, heartwood, dead wood) and surface areas (leaf, bark, cambium) for each species of tree, shrub, fern, herb, and woody debris. This poster highlights the various methods we used to achieve this goal. Plots were all one hectare, scaled to be 10:1, or 316.23 x 31.623 m. Within the entire plot, complete inventories were made of all live trees > 5 cm DBH, all dead trees > 5 cm and taller than 0.5 m, and all downed wood > 30 cm diameter. Each live tree was identified to species, mapped for its X–Y position within the plot, and marked with an aluminum tag and nail (rain forests) or stainless steel tag and nail (all other plots) at 1.4 m or often higher with very large trees or tree clusters. Measurements of height, height to crown base, 4–8 crown radii, diameter at the nail, and nail height were collected for each tree. Trees not yet tagged, shrubs, ferns, herbaceous vegetation, and fine woody material was subsampled, most often using the

316 m plot centerline as a transect. The large and often irregular shapes of tree bases or tree clusters required detailed mapping to get the aboveground resolution needed for this study. One of a few different mapping methods were employed for all non-round tree bases, depending on circumstances. Tree mapping was done first, so that hard copy maps could be error-checked in the field and updated while performing the various plot-level tasks. Maps were error-checked a minimum of three times by the time plot installation was completed. Converting the relative plot coordinate system into real-Earth coordinates was a multi-stage process involving the comparisons of three data sets collected both from ground work and LiDAR-based estimates. The first was a comparison of a detailed DEM (Digital Elevation Model) prepared from ground sampling using lasers to the LiDAR-generated DEM. The second comparison involved using tree crown maps based on the stem map and measured crown radii with the LiDAR CHM (Canopy Height Model). The third comparison used mapped log positions with any large diameter logs detected while creating the LiDAR DEM. Once these three datasets had been linked, a minimizing algorithm was used to lock in the four plot corners (final error was 0.2–0.4 m).

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## **Poster 5**

### **Redwoods, Lasers, Optics, and Bytes: Computational Remote Sensing for the RCCI From Trees to the Landscape**

Bill Kruse, Kruse Imaging

How has remote sensing currently contributed to the Redwoods and Climate Change Initiative? Further, how can we scale the detailed ground truth measurements made by the Initiative's scientists in the long-term reference and smaller satellite plots to the forest and ultimately the landscape? LiDAR, multi-spectral, and hyper-spectral remotely sensed data sets contain a wealth of quantitative information about coast redwoods, their forests, and their topographic environment. Computational remote sensing is being used to mine this data for useful information about the RCCI's signature trees, plots, and the surrounding landscape. LiDAR is used to measure forests from an aircraft. It consists of a pulsed, near-infrared laser scanner that is able to measure the location of reflections from each laser pulse with the help of GPS and inertial navigation. Each reflected point contributes to a point cloud that can be analyzed to measure tree and forest 3D structure. Multi-spectral data is like 2D color digital photos but with less than ten additional spectral bands for analysis. General categories of vegetation and materials can be identified at relatively high ground resolution. Hyper-spectral data also produces 2D imagery but provides many more spectral bands for analysis. This enables more detailed differentiation of vegetation species and materials categories than multi-spectral data can but at the cost of lower ground resolution and higher processing requirements. Used together, these three complimentary remote sensing data resources work together to improve the analysis of our forest's health and change.

RCCI's Phase I utilized LiDAR data, containing 3D forest structure information, to support work in 12 of the 16 long-term reference plots. This data was processed using custom developed workflows, to identify location, height, and crown volume of individual trees and used to create high resolution ground, canopy surface, and canopy height raster maps at sub-meter geographic accuracy. All of the 12 long-term reference plot locations were selected and finalized with the help of the newly created LiDAR canopy height maps. An ArcGIS geodatabase of all newly created LiDAR derived data layers and selected supporting GIS layers was assembled for each LiDAR supported RCCI Phase I plot to provide a permanent geographic data reference archive. RCCI's Phase II has access to a selection of optical data sets that can be utilized to compliment the LiDAR derived geographic data layers. Color, near infrared and

hyperspectral imagery with 128 bands of spectral information, where available, can be used with LiDAR primarily to differentiate and identify vegetation types in the plots and surrounding areas. New LiDAR processing software being developed can also be used to automatically isolate individual trees and subcanopy vegetation from the LiDAR point clouds, allowing the identification of specific crown structures which are essential to accurate stand to landscape scaling. These derived information products provide a scalable plot-to-landscape context to the Initiative's active multi-disciplinary research programs. Calibration and validation of the computed products is provided by the detailed hands-on measurements in signature trees, the existing large old-growth reference plots, as well as the increasingly important smaller and more widely distributed satellite plots.

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## **Poster 6**

### **Monitoring Microclimate in the Redwoods**

Wendy Baxter, Anthony Ambrose, Chris Wong, Cameron Williams, Rikke Næsborg, and Todd Dawson, UC Berkeley

As part of the project goal to provide detailed climate information for the Redwood and Climate Change Initiative plots, microclimate monitoring systems were installed at the treetop and ground level at three coast redwood (*Sequoia sempervirens*) and three giant sequoia (*Sequoiadendron giganteum*) study sites. The information supports more detailed research conducted at specific sites that goes beyond what can be obtained from regional scale satellite or weather data. Measurements are continuously recorded every 30 minutes on key climatic variables, enabling not only species and site-level comparisons but also treetop and ground-level characterizations. Treetop environmental conditions experienced throughout the range of both species are driven mainly by latitude, distance from coast, elevation, and microsite topography. For the coast redwood sites, the first year of data highlight the importance of the summer coastal fog influence at Landels Hill-Big Creek Reserve (LHBCR) and Jedediah Smith Redwoods State Park (JSRSP) in moderating temperatures and vapor pressure deficits (VPDs). Montgomery Woods State Natural Reserve (MWSNR), which is farther inland, experiences higher summer temperatures and VPDs, but during the autumn and winter experiences conditions more similar to those at JSRSP. As the autumn sets in, LHBCR becomes the warmest site but has relatively stable temperatures throughout the year, as does JSRSP. As the northernmost site, JSRSP receives the most precipitation. However MWSNR consistently has the highest soil moisture, presumably due to an elevated water table resulting from an ancient landslide downstream of the site. The range of temperatures and VPDs experienced at the giant sequoia sites is greater than that at the coast redwood sites. However, all three giant sequoia sites follow very similar seasonal patterns, unlike the coast redwood sites. Calaveras Big Trees State Park (CBTSP) is consistently the warmest and experiences the highest VPDs. Although it is the northernmost giant sequoia site, its lower elevation is most likely responsible for these trends. The difference between treetop and ground conditions is greatest at CBTSP, with the treetop experiencing higher temperatures and VPDs. Freeman Creek Grove (FCG) generally receives less precipitation, which is also reflected by lower soil moisture. CBTSP received the most precipitation during the first year of measurements. In the context of climate change, increasing temperatures, decreasing fog and reduced snowpack may play important roles in seedling establishment and overall water-status of mature trees. Continued monitoring will facilitate characterization of inter-annual and long-term trends in environmental conditions, further clarify site-level differences, and may be important in detecting future climatic changes

at each site. These data have been essential to the physiological and isotopic research conducted at these sites, but have many other potential future applications and will be available to other researchers and land management agencies. A possible expansion of the network is being considered.

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### Within-crown and whole-tree water use patterns in giant sequoia trees

Anthony Ambrose<sup>1</sup>, Stephen Sillet<sup>2</sup>, Todd Dawson<sup>1</sup>, Stephen Burgess<sup>3</sup>, George Koch<sup>4</sup>, Robert Van Pelt<sup>2</sup>, Wendy Baxter<sup>1</sup>, Cameron Williams<sup>1</sup>, Rikke Næsborg<sup>1</sup>, Marie Antoine<sup>2</sup>, Jim Spickler<sup>2</sup>, Russell Kramer<sup>2</sup>, and Chris Wong<sup>1</sup>

<sup>1</sup>U.C. Berkeley, <sup>2</sup>Humbolt State University, <sup>3</sup>University of Western Australia, <sup>4</sup>Northern Arizona University

Giant sequoia (*Sequoiadendron giganteum*) trees are the largest and among the oldest living organisms on Earth, with many individuals possessing massive and complex crowns. Spatial and temporal variability in climatic factors interact with changes in tree structure and water status throughout individual crowns to determine water and carbon exchange rates at leaf, branch, and whole-tree scales. We measured sap flow rates throughout the crowns of 3 large *Sequoiadendron* trees growing at UC Berkeley's Whitaker Forest Research Station to characterize within-crown and whole-tree water-use patterns and to examine the relative influence of structural and environmental factors in driving these patterns. Sap flow was measured in individual branches located on opposite sides of each tree crown in lower, middle, and treetop locations, and at the tree base, live crown base, and treetop in the main trunk of each tree. Branch xylem water potential (i.e., water status) and environmental variables such as vapor pressure deficit and solar radiation patterns were also characterized at each sapflow measurement location to account for the effects of highly variable environmental conditions throughout each crown. Each tree was also crown-mapped and a sub-set of branches dissected to provide detailed tree structural information, including branch and trunk sapwood and leaf area. Environmental conditions at different crown positions in each tree were highly variable. Diurnal patterns of solar radiation, atmospheric vapor pressure deficit (VPD) and xylem water potential varied considerably among branches and reflected the interacting influence of both height and the daily trajectory of solar radiation exposure at each branch. In general, branches on the eastern side of each tree experienced greater solar radiation levels, higher VPD, and lower water potentials in the first half of the day, while branches on the western side of each tree experienced these conditions in the second half of the day. Sap flow in all branches, expressed as mean stomatal conductance rates, also showed considerable variability throughout the day and followed similar diurnal patterns as environmental conditions. Despite increasing levels of water stress with increasing height, branches in the upper crown generally exhibited greater stomatal conductance rates than branches in the lower crown for much of the day, suggesting that foliage in upper and outer crown positions is responsible for the majority of water use and consequently carbon gain at the whole-tree scale. Branch stomatal conductance rates were significantly correlated with solar radiation, VPD, and xylem water potential at all crown locations, although the strength of the correlations increased with height ( $r^2 = 0.28-0.50$  for solar radiation,  $0.25-0.39$  for VPD, and  $0.14-0.51$  for water potential). We estimate that sapflow through the base of the main trunk of large individual *Sequoiadendron* trees can exceed

2,000 liters of water per day during the summer, substantially more than any other tree species reported in the literature. Whole-tree sapflow rates also appeared to be influenced by the spatial distribution of trees and the consequent effect on solar radiation exposure of individual tree crowns as well as the vertical distribution of leaf area. These findings suggest that entire *Sequoiadendron* groves require enormous quantities of water, and raises concerns that tree and forest structure and function might be compromised with increasing drought conditions predicted for the Sierra Nevada Mountains in the future.

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## Glossary

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**Azimuthal**—referring to the direction or orientation of an object relative to north, usually measured in degrees

**Annual ring**—One year's growth of wood produced by the cambium and measured for thickness

**Bark**—Consists of two layers produced by cambium: a dead outer protective layer and a living inner layer that transports sugars from leaves to roots

**Basal area**—the area of ground occupied by the cross-section of tree trunks and stems

**Biomass**—organic material comprising a plant body

**Cambium**—Living cell layer beneath bark that produces bark from its outer surface and wood from its inner surface

**CHM**—canopy height model

**Crossdating**—the precise dating of annual growth rings based on the common patterns of ring widths over a population of trees

**Dendrochronology**—the dating and study of annual rings in trees

**DBH**—diameter at breast height; a standardized measurement of tree trunk width taken at 1.37m above ground (or 4.5 feet)

**DEM**—digital elevation model

**Embolism**—development of air bubbles within the water-conducting xylem cells of a plant, thus preventing further water movement within those cells

**Growth ring**—a layer of wood produced during a tree's growing season

**Heartwood**—Completely dead layer of wood beneath sapwood that contains toxic chemicals resistant to decay

**Inter-annual (high-frequency)**—occurring between years (from one year to the next) as opposed to a long-term trend (low-frequency)

**LAI (leaf area index)**—the one-sided leaf area per unit ground

**LiDAR (Light Detection and Ranging)**—a type of remotely sensed spatial data

**Marker years**—consistently narrow or large growth rings, or patterns thereof, that are the basis for crossdating

**Photosynthesis**—process in which light energy from the sun is used to convert CO<sub>2</sub> from the air into organic compounds in a plant

**PDSI (Palmer Drought Severity Index)**—a widely used index reflecting drought severity, or soil moisture

**Provenance**—geographic origin or source location of plant seeds

**Radii**—a series of growth rings spanning from the oldest rings to the youngest, capturing a cross-sectional view of tree growth

**Sap flow**— the movement of water through a plant

**Sapwood**—Partially living layer of wood beneath cambium that transports water from roots to leaves

**SEGI**—abbreviation for *Sequoiadendron giganteum* (giant sequoia)

**SESE**—abbreviation for *Sequoia sempervirens* (coast redwood)

**Snag**—a standing dead tree

**Stable carbon isotope ratio**— the ratio of naturally occurring carbon-12 to carbon -13 atoms within a compound

**Stable oxygen isotope ratio**—the ratio of naturally occurring oxygen-16 to oxygen-18 atoms within a compound

**Transpiration**—water loss from plant leaves due to evaporation through tiny pores in the leaf called stomata

**VPD**—vapor pressure deficit

**Water deficit**—reduction in the amount of soil water available for uptake by plant roots

**Water potential**—chemical potential of water in soils, plants, and the atmosphere

**Xylem**— the woody cellular system of higher plants responsible for conduction of water and dissolved minerals as well as support and food storage