## SUPPLEMENTARY MATERIAL

## 1. Location of MODIS pixels used for each site

Table S1. Location of MODIS MCD43A4.006 pixel used for each study site.

|  | Eddy covariance tower <br> location |  |  |  | MODIS pixel |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: |

## 2. Plant traits at the study sites

Table S2. Species sampled and their leaf traits.

| Site | Sampling <br> date <br> 2019 | Species | Leaves for site's LMA and $W_{c}$ community weighted means |  |  | Leaves with photosynthesis measurements |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | n | $\begin{gathered} \mathrm{LMA} \pm \mathrm{SD} \\ \mathrm{~g} \mathrm{~cm}^{-2} \\ \hline \end{gathered}$ | $\begin{gathered} \mathrm{W}_{\mathrm{c} \pm} \pm \mathrm{SD} \\ \mathrm{~g} \mathrm{~cm}^{-2} \end{gathered}$ | n | $\begin{gathered} \text { Aleaf } \pm \mathrm{SD} \\ \mu \mathrm{~mol} \mathrm{~m} \\ \mathrm{~m}^{-2} \mathrm{~s}^{-1} \end{gathered}$ | $\begin{gathered} \mathrm{LMA} \pm \mathrm{SD} \\ \mathrm{~g} \mathrm{~cm}^{-2} \\ \hline \end{gathered}$ |
| CSs | April 21 and 26 <br> April 21 and | Artemisia californica | 5* | $0.0103 \pm 0.0014$ | $0.0155 \pm 0.0011$ | 5 | $12 \pm 3.8$ | $0.0112 \pm 0.0005$ |
| CSs | $26$ <br> April 21 and | Eriogonum fasciculatum | 4* | $0.0150 \pm 0.0005$ | $0.0203 \pm 0.0012$ | 5 | $21.2 \pm 4$ | $0.0157 \pm 0.0019$ |
| CSs | $26$ <br> April 21 and | Leymus condensatus | 10 | $0.0086 \pm 0.0020$ | $0.0123 \pm 0.0021$ | 5 | $11 \pm 6.1$ | $0.0089 \pm 0.0012$ |
| CSs | $26$ <br> April 21 and | Malosma laurina | 33 | $0.0097 \pm 0.0019$ | $0.0142 \pm 0.0023$ | 5 | $23.5 \pm 9.8$ | $0.0108 \pm 0.0007$ |
| CSs | 26 | Salvia mellifera | 32 | $0.0086 \pm 0.0022$ | $0.0159 \pm 0.0027$ | 5 | $20.4 \pm 7.2$ | $0.0099 \pm 0.0014$ |
| Ds | May 1, 2, 11 | Cylindropuntia ramosissima | 7 | $0.1806 \pm 0.0721$ | $0.3776 \pm 0.0987$ | 0 |  |  |
| Ds | May 1, 2, 11 | Hymenoclea salsola | $10^{*}$ | $0.0111 \pm 0.0019$ | $0.0391 \pm 0.0079$ | 5 | $33.2 \pm 9.2$ | $0.0152 \pm 0.0039$ |
| Ds | May 1, 2, 11 | Hyptis emoryi | 9 | $0.0103 \pm 0.0018$ | $0.0178 \pm 0.0025$ | 0 |  |  |
| Ds | May 1, 2, 11 | Justicia californica | 20 | $0.0092 \pm 0.0034$ | $0.0220 \pm 0.0051$ | 5 | $26.4 \pm 6.9$ | $0.0125 \pm 0.001$ |
| Ds | May 1, 2, 11 | Larrea tridentata | 10* | $0.0134 \pm 0.0011$ | $0.0207 \pm 0.0014$ | 5 | $10.3 \pm 3.9$ | $0.0169 \pm 0.0021$ |
| Ds | May 1, 2, 11 | Parkinsonia florida | 10* | $0.0074 \pm 0.0005$ | $0.0151 \pm 0.0020$ | 5 | $21.6 \pm 2.8$ | $0.011 \pm 0.0012$ |
| Dc | May 9 and 10 | Erodium cicutarium | $5^{*}$ | $0.0050 \pm 0.0007$ | $0.0079 \pm 0.0014$ | 5 | $13.8 \pm 5.6$ | $0.0054 \pm 0.0002$ |
| Dc | May 9 and 10 | Eriogonum fasciculatum | 10* | $0.0119 \pm 0.0020$ | $0.0212 \pm 0.0020$ | 5 | $29.9 \pm 3.5$ | $0.023 \pm 0.0029$ |
| Dc | May 9 and 10 | Purshia tridentata | 10* | $0.0154 \pm 0.0011$ | $0.0248 \pm 0.0013$ | 5 | $18.2 \pm 4.6$ | $0.023 \pm 0.0029$ |
| Dc | May 9 and 10 | Stipa speciosa | $5^{*}$ | $0.0092 \pm 0.0006$ | $0.0122 \pm 0.0002$ | 5 | $10.7 \pm 2.1$ | $0.0064 \pm 0.0006$ |
| Dc | May 9 and 10 April 19, 24, | Yucca schidigera | 25 | $0.1225 \pm 0.0145$ | $0.1585 \pm 0.0164$ | 0 |  |  |
| G | 26 | Amsinkia menziesii | $10^{*}$ | $0.0059 \pm 0.0006$ | $0.0248 \pm 0.0017$ | 5 | $21.2 \pm 6.5$ | $0.0068 \pm 0.0006$ |
| G | April 19, 24, | Avena sp. | 24 | $0.0041 \pm 0.0010$ | $0.0142 \pm 0.0037$ | 5 | $22.4 \pm 5$ | $0.0042 \pm 0.0006$ |


|  | 26 |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| G | $\begin{aligned} & \text { April 19, 24, } \\ & 26 \end{aligned}$ | Brassica nigra | 21 | $0.0049 \pm 0.0009$ | $0.0190 \pm 0.0040$ | 5 | $30.2 \pm 3.4$ | $0.0053 \pm 0.0007$ |
| G | $\begin{aligned} & \text { April 19, 24, } \\ & 26 \end{aligned}$ | Bromus diandrus | 25 | $0.0039 \pm 0.0006$ | $0.0107 \pm 0.0016$ | 5 | $12 \pm 3.6$ | $0.0042 \pm 0.0005$ |
| G | $\begin{aligned} & \text { April 19, 24, } \\ & 26 \end{aligned}$ | Leymus condensatus | 0 |  |  | 2 | $26.3 \pm 1.3$ | $0.01 \pm 0.0015$ |
|  | April 19, 24, |  |  |  |  |  |  |  |
| G | 26 <br> April 19, 24, | Stipa pulchra | 6* | $0.0097 \pm 0.0031$ | $0.0190 \pm 0.0072$ | 5 | $13.1 \pm 4.1$ | $0.0059 \pm 0.0013$ |
| G | $26$ <br> July 18 and | Unidentified forb | 28 | $0.0048 \pm 0.0007$ | $0.0209 \pm 0.0039$ | 5 | $28.9 \pm 3.2$ | $0.0059 \pm 0.0012$ |
| PJw | $19$ <br> July 18 and | Bernardia incana | 8* | $0.0098 \pm 0.0016$ | $0.0134 \pm 0.0016$ | 4 | $5.7 \pm 1.9$ | $0.0106 \pm 0.0011$ |
| PJw | 19 | Juniperus californica | 3* | $0.0719 \pm 0.0079$ | $0.0099 \pm 0.0010$ | 3 | $4.3 \pm 0.2$ | $0.0719 \pm 0.0079$ |
| PJw | July 18 and 19 <br> July 18 and | Pinus monophylla | 36 | $0.0608 \pm 0.0125$ | $0.0632 \pm 0.0095$ | 5 | $1.2 \pm 0.7$ | $0.0679 \pm 0.0186$ |
| PJw | 19 | Quercus palmeri | 30 | $0.0180 \pm 0.0041$ | $0.0119 \pm 0.0042$ | 5 | $13.5 \pm 3.9$ | $0.0202 \pm 0.0015$ |
| Sierra <br> sites | TRY <br> database | Pinus contorta |  |  |  |  | 3.0 | 0.0227 |
| Sierra <br> sites | TRY <br> database | Pinus jeffreyi |  |  |  |  | 8.9 | 0.0240 |
| Sierra sites | TRY <br> database | Pinus monticola |  |  |  |  | 8.2 | 0.0200 |
| Sierra <br> sites | TRY <br> database | Pinus ponderosa |  |  |  |  | 6.3 | 0.0234 |

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## 3. Details about the relationships between vegetation indices and gross primary production

The exploratory analysis of the data for each site indicate that linear regressions adequately describes relationship between Vegetation Indices (VIs) and Gross Primary Production (GPP) at the studied sites (see Figure S1-S3). For the relationship with NDVI, only two sites (grassland and coastal sage shrubland) showed some degree of curvature at the highest values of NDVI, although the linear regression explained $71-85 \%$ of GPP's variation (the highest $r^{2}$ ). The relationship for EVI and NIRv was linear in all the sites.


Figure S1. Relationship between NDVI and midday gross primary production for each site.


Figure S2. Relationship between EVI and midday gross primary production for each site.


Figure S3. Relationship between EVI and midday gross primary production for each site.

Detailed information about the coefficients of the linear regressions between VIs and GPP for each site and across all sites is shown in Table S3. The observations in most of the sites do not homogeneously sample the their VI range, each site having more observations towards its lower end than towards the higher end (see Figures S1-S3). In addition, there are large differences in the number of observations available between sites, each sites having 5 to 12 years of observations. In order to minimize the impact of these issues on the results of our regression analyses, we performed our analysis as follows: for each site we 1) split the VI range of each site in 10 equally sized bins and took 10 random samples from each bin, 2) run the linear regression for each site with these more balanced samples and store the results, 3) then we run a linear regression for the samples taken for all the sites to estimate the across-sites relationships and store the results, 5) repeat all previous steps 200 times and average the obtained regression coefficients. Besides minimizing effects of unequal sampling within and across sites, this procedure allows using the entire dataset, and prevents that $P$ values become meaningless with very large sample numbers. EVI and NIR ${ }_{v}$ have stronger within-site relationships in 9 of 10 sites, narrower ranges in its intercept and slopes, and stronger across-sites relationships.

We performed a similar procedure to provide an overall statistical measure that allowed us to compare the differences on within-site relationships across VIs. For this, we fitted the data to a statistical model evaluating the shared effect of the vegetation index across sites (equivalent to the across-site relationship), and the effect of site differences in the intercept (site effect) and slopes (interaction of site and VI): $=\beta_{0}+\beta_{1} \cdot V I+\beta_{2} \cdot$ site $+\beta_{3} \cdot V I \cdot$ site $+e$, where $\beta$ are the statistical coefficients for each term. For a given VI, the percent of the overall GPP variance explained by site differences in intercepts and slopes (site + site-VI interaction effects) provides a measure of how different the relationships are between sites. This analysis was implemented on the same samples described above, and the results of the 200 runs were averaged. According to this analysis, site differences in intercepts and slopes in NDVI explain more than twice overall GPP variation ( 23.6 \% variance explained) than in EVI ( $9.9 \%$ ) and NIRv ( $8.4 \%$ ). These results, together with the narrower ranges in intercepts and slopes, support quantitatively that site/biome relationships for EVI and NIR ${ }_{\mathrm{v}}$ are more convergent than for NDVI.

Table S3. Results of linear regression analyses for the temporal (single-site) and spatial (across-sites) relationships between vegetation indices and gross primary production.

| Site | NDVI |  |  | EVI |  |  | NIR ${ }_{v}$ |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $r^{2}$ <br> (P value) | Intercept | Std slope* | $\mathrm{r}^{2}$ <br> (P value) | Intercept | Std slope* | $r^{2}$ <br> (P value) | Intercept | Std slope* |
| Coastal sage shrubland | $\begin{gathered} 0.74 \\ (<0.001) \end{gathered}$ | -21.8 | 11.5 | $\begin{gathered} 0.80 \\ (<0.001) \end{gathered}$ | -11.2 | 7.4 | $\begin{gathered} 0.80 \\ (<0.001) \end{gathered}$ | -8.0 | 7.1 |
| Desert chaparral | $\begin{gathered} 0.43 \\ (<0.001) \end{gathered}$ | -5.3 | 6.4 | $\begin{gathered} 0.62 \\ (<0.001) \end{gathered}$ | -6.6 | 5.7 | $\begin{gathered} 0.62 \\ (<0.001) \end{gathered}$ | -4.6 | 5.3 |
| Desert shrubland | $\begin{gathered} 0.52 \\ (<0.001) \end{gathered}$ | -6.5 | 12.2 | $\begin{gathered} 0.64 \\ (<0.001) \end{gathered}$ | -6.5 | 8.8 | $\begin{gathered} 0.61 \\ (<0.001) \end{gathered}$ | -4.9 | 10.0 |
| Grassland | $\begin{gathered} 0.83 \\ (<0.001) \end{gathered}$ | -16.1 | 9.6 | $\begin{gathered} 0.87 \\ (<0.001) \end{gathered}$ | -7.5 | 5.4 | $\begin{gathered} 0.88 \\ (<0.001) \end{gathered}$ | -4.7 | 4.8 |
| Oak-pine forest | $\begin{gathered} 0.02 \\ (0.401) \end{gathered}$ | 5.3 | 1.1 | $\begin{gathered} 0.06 \\ (0.066) \end{gathered}$ | 4.9 | 1.3 | $\begin{gathered} 0.08 \\ (0.022) \end{gathered}$ | 5.3 | 1.4 |
| Oak-pine woodland | $\begin{gathered} 0.61 \\ (<0.001) \end{gathered}$ | -10.0 | 7.0 | $\begin{gathered} 0.61 \\ (<0.001) \end{gathered}$ | -6.1 | 4.7 | $\begin{gathered} 0.61 \\ (<0.001) \end{gathered}$ | -3.4 | 4.2 |
| Pinyon-juniper woodland | $\begin{gathered} 0.39 \\ (<0.001) \end{gathered}$ | -7.0 | 7.1 | $\begin{gathered} 0.67 \\ (<0.001) \end{gathered}$ | -9.4 | 7.0 | $\begin{gathered} 0.72 \\ (<0.001) \end{gathered}$ | -8.9 | 8.1 |
| Ponderosa pine forest | $\begin{gathered} 0.50 \\ (<0.001) \end{gathered}$ | -25.8 | 10.5 | $\begin{gathered} 0.72 \\ (<0.001) \end{gathered}$ | -15.1 | 8.0 | $\begin{gathered} 0.73 \\ (<0.001) \end{gathered}$ | -10.9 | 7.7 |
| Sierran mixed conifer forest | $\begin{gathered} 0.08 \\ (0.049) \end{gathered}$ | -0.2 | 2.5 | $\begin{gathered} 0.33 \\ (<0.001) \end{gathered}$ | -7.9 | 4.7 | $\begin{gathered} 0.33 \\ (<0.001) \end{gathered}$ | -4.1 | 4.2 |
| Subalpine forest | $\begin{gathered} 0.02 \\ (0.455) \\ \hline \end{gathered}$ | 7.4 | -0.6 | $\begin{gathered} 0.29 \\ (<0.001) \\ \hline \end{gathered}$ | -4.0 | 4.1 | $\begin{gathered} 0.25 \\ (<0.001) \\ \hline \end{gathered}$ | -1.6 | 4.2 |
| Site minimum | 0.02 | -25.8 | -0.6 | 0.06 | -15.1 | 1.3 | 0.08 | -10.9 | 1.4 |
| Site maximum | 0.83 | 7.4 | 12.2 | 0.87 | 4.9 | 8.8 | 0.88 | 5.3 | 10.0 |
| Across-site relationship | $\begin{gathered} 0.47 \\ (<0.001) \end{gathered}$ | -3.5 | 4.3 | $\begin{gathered} 0.68 \\ (<0.001) \end{gathered}$ | -5.3 | 4.7 | $\begin{gathered} 0.71 \\ (<0.001) \end{gathered}$ | -3.3 | 4.6 |

* Slopes are not directly comparable when contrasting relationships from different vegetation indices because NDVI, EVI and NIRv have very different ranges of possible values. To make slopes comparable among relationships from different vegetation indices, we also regressed GPP against centered
(on the mean across all sites) and scaled (to one standard deviation across all sites) vegetation indices data. We called the slopes obtained this way as
"standardized slopes" (or Std. slopes). In particular, these standardized slopes allows an appropriate assessment of which vegetation index has more
consistent (similar) slopes among sites. The scaling and centering is not needed for a comparing intercepts because are always in the same units (the same units of gross primary production, $\mu \mathrm{mol}$.


## 4. Analysis with Thornley (2002) and PROSAIL models

The GPP modeled with similar parametrization as in Figure 3b (see Appendix B) but including sitespecific LAI and Aleaf closely follows the eddy covariance GPP measurements (Figure S4).


Figure S4. Comparison of observed and modelled midday gross primary production (GPP) at the study sites. Observed GPP is the mean $\pm 1$ standard deviation of the cloudless eddy covariance observations taken under similar conditions (including similar vegetation indices). Modeled GPP is based on Thornley model and the in-situ observations of LAI and Aleaf.

The vegetation indices modeled from PROSAIL parametrized as in Figure 6 (see Appendix B), but with site-specific observations on soil reflectance (Figure S5), LAI and LMA, capture well the observed variation in vegetation indices at the study sites around the field sampling dates (Figure S6). The observed data for Figure 56 correspond to nine Landsat 8 pixels centered on the $90 x 90$ sampling area at the study sites, obtained during clear conditions around the field sampling dates. The Landsat 8 surface reflectance data was obtained from the U.S. Landsat Analysis Ready Data (ARD) product CU_LC08, downloaded from https://lpdaacsvc.cr.usgs.gov/appeears/ and was filtered for cloud and aerosol conditions.


Figure S5. Soil reflectance spectra from the eddy covariance sites used in this study. We measured the soil reflectance at the five sites, and obtained supplemental observations from three of the other sites from Serbin et al ("NASA HyspIRI Airborne Campaign Leaf and Canopy Spectra and Leaf Traits", available at https://ecosis.org). Our study sites span a range of soil reflectance, with bright, sandy granitic soils at the desert shrubland, desert chaparral and pinyon-juniper woodland sites, and dark soils with higher organic content at the coastal sage shrubland, grassland, mixed conifer forest, and oak-pine woodland sites.

Site

Coastal sage shrubland

- Desert shrubland
- Desert chaparral

Grassland
Pinyon-juniper woodland




Figure S6. Comparison of observed vegetation indices from Landsat 8 with those modelled with PROSAIL parametrized with site-specific information of LAI, LMA and soil reflectance. Landsat 8 data were.

## 5. Leaf chlorophyll data from plant species from Southern California and Sierra Nevada

The leaf chlorophyll content per leaf area ( Cab ) information used in analyses was based on existing datasets of leaf reflectance from five of our eddy covariance sites (dataset from Serbin et al). This data set contains measurements of shrub and tree species, including broadleaf and needleleaf, and either deciduous or evergreen, as well as grasses. We estimated $C_{a b}$ from these measurements using the red edge chlorophyll index following Gitelson and Solovchenko (2017). The Cab range found in this dataset (9$58 \mu \mathrm{~g} \mathrm{~cm}^{-2}$, Figure S7) was used for the global sensitivity analysis.

Our analyses indicated that precise variation of $\mathrm{C}_{\text {ab }}$ across sites was not needed in our models to capture the observed variation in vegetation indices of the sites (see Figure 6b). For this reason, in some of the modeling analyses (Figure 6b-d) in our study we assumed a fixed Cab value of $30 \mu \mathrm{~g} \mathrm{~cm}$ close to the mean from these data (Figure S7).


Figure S7. Histogram of leaf chlorophyll content per leaf area in species found in five of the eddy covariance sites. Solid line corresponds to the mean.

## 6. Correlations between vegetation indices: results from PROSAIL

We analyzed the correlations between vegetation indices from modeling results with PROSAIL. This allowed us to assess whether the correlations found between vegetation indices were specific to the conditions at our study sites, the specific satellite product used, or a more generalizable pattern supported by radiative transfer theory (Figure S8). We parametrized PROSAIL to model surface reflectance from all possible combinations from varying each parameter 5 steps over the same ranges used in the global sensitivity analysis (see Supplementary Material 1), and varying mean leaf angle over $10-80{ }^{\circ}$. This generated more than 78,000 simulations from the resulting combinations. The results show similar patterns to those obtained from the correlations in satellite data (Figure 7). This confirms that the strong similarity between EVI and NIR $_{v}$ holds over a diverse combination of environmental and vegetation conditions.


Figure S8. Correlation analysis between vegetation indices, using reflectance data modeled with PROSAIL.

## 7. Compilation of field measurements of mean canopy leaf angles

We compiled field measurements of mean canopy leaf angles from the literature to aid us in the selection of a range of values that well represent the variation found in this parameter across ecosystems. This dataset gathered 531 records from 48 publications, and included data from single-species and multispecies canopies from most major biomes (dataset available at Hinojo-Hinojo and Goulden, 2020). Only data from field-grown plants were included. When studies presented data on seasonal variation on mean canopy leaf angles from the same species/locations, the data were averaged and considered as a single record. The records were originally measured using several techniques, including clinometers and protractor measurements of individual leaves throughout the canopy, leveled photographs and digital measurement of individual leaves on image processing software, canopy hemispheric photographs, canopy analyzers (e.g. LAI-2000 and LAI-2200), and LIDAR. We excluded the data from BOREAS from our analysis (see Hinojo-Hinojo and Goulden, 2020). In our global sensitivity analyses, we used the range $10-90 \%$ percentile range of the dataset, which was 25-630 (Figure S9).


Figure S9. Histogram of the compiled dataset of mean canopy leaf angle

## REFERENCES

Gitelson, A., Solovchenko, A., 2017. Generic Algorithms for Estimating Foliar Pigment Content. Geophys. Res. Lett. 44, 9293-9298.

Thornley, J.H.M., 2002. Instantaneous canopy photosynthesis: Analytical expressions for sun and shade leaves based on exponential light decay down the canopy and an acclimated non-rectangular hyperbola for leaf photosynthesis. Ann. Bot. 89, 451-458.

Hinojo-Hinojo, C, Goulden, M.L. 2020. A dataset of published measurements of canopy leaf inclination angles, UC Irvine, Dataset, https://doi.org/10.7280/D1T97H


[^0]:    Sites are CSs: coastal sage shrubland, Ds: desert shrubland, G: grassland, Dc: Desert chaparral, PJw: pinyon-juniper woodland. Measured traits are LMA: leaf mass per area, $\mathrm{W}_{c}$ : leaf water content, Aleaf: leaf photosynthesis. *Small-leaved species where LMA and $\mathrm{W}_{\mathrm{c}}$ trait measurements were made on compound samples of several leaves. Each compound sample may contain tens to several tens of leaves/leaflets.

