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Abstract Groundwater is an integral component of the water cycle, and it also influences the carbon cycle by supplying moisture to ecosystems. However, the extent and determinants of groundwater-vegetation interactions are poorly understood at the global scale. Using several high-resolution data products, we show that the spatial patterns of ecosystem gross primary productivity and groundwater table depth are correlated during at least one season in more than two thirds of the global vegetated area. Positive relationships, i.e., larger productivity under shallower groundwater table, predominates in moisture-limited dry to mesic conditions with herbaceous and shrub vegetation. Negative relationships, i.e., larger productivity under deeper groundwater, predominates in humid climates with forests, possibly indicating a drawdown of groundwater table due to substantial ecosystem water use. Interestingly, these opposite groundwater-vegetation interactions are primarily associated with differences in vegetation than with climate and surface characteristics. These findings put forth the first evidence, and a need for better representation, of extensive and non-negligible groundwater-vegetation interactions at the global scale.

1. Introduction

Groundwater is a critical source of soil moisture supply for evapotranspiration [Lautz, 2008; Martinec et al., 2009; Niu et al., 2007; Yeh and Famiglietti, 2009] and its components [Cramer et al., 1999; Maxwell and Condon, 2016] when radiation energy is abundant and precipitation is limited. Several studies have shown that under such moisture-limited conditions, this additional moisture supply also enables vegetation to limit the loss of productivity [Gou et al., 2015], especially in water-limited ecosystems [Baldocchi et al., 2010; Barbeta et al., 2015]. This access to and potential dependence on below-ground moisture supply from groundwater can even shape local-scale vegetation distribution and species segregation [Orellana et al., 2012; Silvertown et al., 2015], and it has been speculated that groundwater may also be a key driver of ecological patterns globally [Fan, 2015]. Unfortunately, efforts to link such patterns have been limited to the local scale in experimental studies and regional scale in remote-sensing or model-based studies (see Eamus et al. [2015] for an in-depth review). At the global scale, we still lack any knowledge on the relevance, extent, timing, and conditions leading to the groundwater-vegetation interactions.

The global-scale efforts to link groundwater and vegetation patterns have been impeded by a dearth of higher resolution data sets and models that consider both groundwater and vegetation processes [Bierkens et al., 2015]. As the collection of field observation data is impractical for regional- and global-scale studies, remote sensing data are suitable to identify groundwater-dependent ecosystems over a large spatial scale [Eamus et al., 2015, Pérez Hoyos et al., 2016]. The remote sensing data have successfully been used to link
terrestrial water storage variations (total storage including groundwater) and vegetation activities on a regional scale [A et al., 2015; McGrath et al., 2012; Yang et al., 2014]. Due to recent advances in Earth observation networks and modeling approaches, we can now investigate the relationship between fine-grained spatial patterns of groundwater and vegetation productivity in the global context as well.

In this study, we exploit relatively high spatial resolution (~10 km) global data sets of model-based groundwater table depth (WTD) together with gross primary productivity (GPP) derived from satellite remote sensing, FLUXNET observations, and multiple machine learning methods to identify where, when, and how spatial patterns of GPP and WTD are related globally. We then assess how these spatial patterns are associated with vegetation, land surface, and climate characteristics.

2. Data and Methods

The GPP data, at ~10 km spatial and 8 daily temporal resolution, were obtained from FLUXCOM [FLUXCOM, 2017; Jung et al., 2017; Tramontana et al., 2016]. Using nine machine learning algorithms, two observed GPP variants from two flux partitioning methods [Lasslop et al., 2010; Reichstein et al., 2005] were upcaled from 224 FLUXNET sites [Baldocchi et al., 2001] to the global scale. The machine learning algorithms were initialized trained to site-level observations of the explanatory climate and land surface variables. To capture variabilities in vegetation greenness and land surface temperature with reliable imprint of changes associated with groundwater, the variables from high-resolution satellite remote sensing data were fed into the extensive variable selection analysis [Jung and Zscheischler, 2013]. The machine learning algorithms and their training and a thorough cross-validation of the data are presented in detail in Tramontana et al. [2016]. To obtain the global GPP, the trained and validated machine learning algorithms were forced with global gridded satellite data of selected explanatory variables, at 10 km spatial and 8 daily temporal resolution, to obtain the GPP for the period of 2001–2012. In total, 18 variants (two observed GPP variants and nine algorithms) of global GPP time series were produced, and the ensemble median was used in this study.

Compared to other global GPP products, the median GPP from FLUXCOM is advantageous because it combines the strengths of multiple global satellite-based observations with site-level observations using several methods. The GPPs estimated only by a subset of algorithms used in FLUXCOM have shown better performance than the GPP from Moderate Resolution Imaging Spectroradiometer (MODIS) against site-level observations [Ichii et al., 2017; Tramontana et al., 2015; Yang et al., 2007]. At the global scale, the Model Tree Ensembles GPP product [Beer et al., 2010; Jung et al., 2011], also estimated by a subset of algorithms used in FLUXCOM, compares well with modern satellite observation of Sun-induced fluorescence [Frankenberg et al., 2013], and it has been used extensively to benchmark global land surface models [Anav et al., 2013; Bonan et al., 2012; Piao et al., 2013].

The WTD data were obtained from an observation-constrained global-scale groundwater model simulation that considers the lateral groundwater flow [de Graaf et al., 2015]. Mean seasonal climatology of WTD is available at ~10 km spatial and monthly temporal resolutions. Forced by climate-driven recharge, land surface, and aquifer parameters, their groundwater model distributes the groundwater recharge laterally based on topography with coasts and river channels as boundary conditions. The WTD simulation was validated against >1.6 million site-level WTD observations. See section S1 in the supporting information for details.

To characterize the spatial covariation of GPP and WTD, we calculated Spearman’s rank correlation coefficient within 11,134 river basins [Lehner et al., 2008] smaller than ~100,000 km² globally (see section S2). For a given basin, the correlation coefficient was calculated between two (or more for partial correlation coefficient) vectors of data variables from all the 10 km grid cells within the basin boundary. The Spearman’s correlation is suitable here because it quantifies the covariation of spatial patterns which, for WTD, are well constrained by topography and bedrock information and, for GPP, by remote sensing of land surface characteristics. We test the statistical significance of the correlation at 5% level of significance.

These GPP-WTD Spearman’s correlations were then controlled for five confounding climatic and land surface variables: precipitation, net radiation, topography/elevation, porosity, and leaf area index to eliminate the “background” collinearity. Precipitation and net radiation represent the moisture and energy available, respectively. Topography drives the spatial variation of WTD, and shallower WTDs usually occur at lower elevation. In addition, elevation differences also account for variation in temperature, which in turn relates
to vapor pressure deficit, a driver of transpiration and GPP. Soil porosity, the potential soil moisture content, affects groundwater recharge and resistance against evapotranspiration. Finally, leaf area index (LAI) accounts for differences in vegetation types within a basin. The data sets of these variables were obtained from several observation-based or model-based sources. A summary of these data is provided in Table 1, and they are explained in detail in section S1. As these data are either based on high-resolution satellite or topographic data, we assume that they represent the spatial variability well. The assumption is a necessity, as there are no global modeling or observation frameworks that could produce all the data needed in a consistent manner.

Since the GPP are positive and WTD are negative numbers while calculating (partial) correlation coefficient, the GPP-WTD correlation is positive for a basin when the grids with larger GPP have shallower WTD (smaller magnitude of a negative number). In contrast, a negative GPP-WTD correlation indicates that grids with larger GPP have deeper WTD. For each basin, the GPP-WTD correlation was calculated seasonally for four phenological stages of the vegetation derived from GPP (see sections S3 and S4 and Forkel et al. [2014]). The correlation coefficient over four seasons is then used to identify the dominant sign of GPP-WTD spatial correlation. The dominant GPP-WTD correlation for a basin is defined as the sign, positive or negative, of the correlation that dominates over four phenological stages. To identify the dominant correlation, we first record the sign of a significant correlation ($p < 0.05$) in each phenological stage, and then calculate the difference in number of seasons with positive correlation minus that with negative correlation. This generalization using the dominant GPP-WTD correlation is reasonable because the sign of GPP-WTD correlation does not change in different phenological stages for 89% of the basins (see section S5 for a detailed explanation). The main text here focuses on the dominant GPP-WTD correlation, while a complete analysis for each phenological stage is presented in the supporting information.

### 3. Results and Discussions

We first present the results for global distribution of the dominant GPP-WTD correlation. Then, we investigate the association of the dominant correlation with vegetation, surface, and climate characteristics. Finally, we evaluate the relative importance of vegetation, surface, and climate characteristics in determining the sign of the dominant GPP-WTD correlation.

#### 3.1. Global Distribution of GPP-WTD Correlation

The GPP-WTD correlations are statistically significant in at least one season for roughly two thirds of the vegetated area (Figure 1). Temporally, the correlation is statistically significant in only one season for 21% and in two seasons for further 15% of the basins. This shows that the groundwater-vegetation interaction,
Figure 1. Global distribution of the dominant GPP-WTD relationship. The red (blue) shades are used for basins where the number of seasons with positive correlation is larger (smaller) than that with negative correlation. The basins in which GPP-WTD correlation is statistically insignificant in all seasons are shaded grey. In the bar chart, the height of a bar shows the area coverage (relative to total global vegetated area), while the number above each bar is the number of basins.

either positive or negative, is significant under certain “ecohydrologically relevant” condition only and has strong temporal dynamics. Further, the area extent of significant positive correlation is the largest when GPP is minimum, showing that positive correlation predominates when vegetation activity is limited. In contrast, the negative correlation predominates when vegetation activity is maximum (see section S5 for details).

Even though the basins dominated by positive and negative GPP-WTD correlations have similar area extents globally, they have a distinct spatial footprint. The positive relationship prevails in ~36% of the vegetated area, located primarily in the Indian subcontinent, sub-Saharan Sahel regions, semiarid regions of eastern and southern Africa, noncoastal regions of southern and eastern Australia (in and around Lake Eyre basin), Eurasian midlatitudes, and northern parts of North America. In most of these regions with typically limited moisture supply, groundwater is likely supporting primary productivity through enhanced soil moisture availability. This is also consistent with earlier model-based findings of sizable groundwater-supplied evapotranspiration in the semiarid regions globally [Koirala et al., 2014] as well as those in seasonally dry northern high-latitude regions [Cohen et al., 2006] and regions with higher probability of using nonprecipitation moisture sources in Australia [Doody et al., 2017].

A negative GPP-WTD relationship predominates in ~36% of the global vegetated area, and it is mostly evident in humid regions concentrated in tropical regions of South America, Africa, Southeast Asia, and parts of northern and northeastern Eurasia. Such an extensive negative GPP-WTD relationship was unexpected, as the previous studies signify, almost exclusively, the positive effect of groundwater on productivity. However, a negative GPP-WTD relationship might emerge due to several mechanisms such as suppression of GPP via anaerobic stress under very shallow WTD [Naumburg et al., 2005] or via a loss of leaves [Parolin and Wittmann, 2010] and structural disturbances [Moreno-Casasola and Vázquez, 1999] in floodplains. Furthermore, and perhaps more essentially and widespread, the WTD may become deeper with increasing GPP through simultaneously increasing evapotranspiration [Jung et al., 2011] and decreasing groundwater recharge.

Nearly 30% of the total vegetated area shows no significant GPP-WTD correlation in any season. There is no spatial coherency of insignificant relationship apart from an extensive area within the contiguous U.S., which might be related to large-scale anthropogenic irrigation that also affects GPP-WTD relationship [Kath et al., 2014]. Uncertainties of the WTD and GPP products likely weaken correlation coefficients as well. We carried out an additional analysis using an independent WTD data from Fan et al. [2013] (section S7). Despite a significant difference in the magnitude of WTD in two data products, the analysis reveals consistent spatial patterns of positive and negative GPP-WTD relationships globally. This is due to a similar regional spatial variability of two data, as WTD is mainly driven by topography in both cases.
3.2. Association With Vegetation, Surface, and Climate Characteristics

To characterize the determinants of the dominant GPP-WTD correlation, we segregated the occurrences of dominant positive and negative relationships for different climate and vegetation classes. We find that negative relationships prevail in humid climate and positive relationships prevail in dry climate irrespective of the vegetation types (Figure 2). In transient climates that are neither strictly humid nor dry, there is a substantial heterogeneity with respect to different vegetation types. In these temperate and seasonally cold regions, there is a clear distinction between prevailing negative relationships for forests and prevailing positive relationships for shrub or grasslands. For Savannas, mixed tree-grass systems, and crops (which are highly managed), there is no clear pattern on whether positive or negative GPP-WTD relationships dominate.

To further identify possible driving mechanisms behind variability of GPP-WTD relationship, we dissected the occurrences of positive and negative relationships against gradients of vegetation characteristics (leaf area index, fractions of tree [Defries et al., 2000] and grass covers, relative fraction of vegetation with C4 photosynthesis [Jung et al., 2006]), and the most important climatic variables (precipitation and net radiation). Because the mean WTD and GPP are independent of the spatial GPP-WTD correlation, they were also included in the analysis.

In general, the dominant sign of GPP-WTD relationships relates more strongly to differences in vegetation characteristics than to differences in mean climate characteristics (Figure 3). The Kolmogorov-Smirnov two-sample test (see section S9 for details) also reveals that the differences in the distribution of the occurrences of positive and negative GPP-WTD relationships are statistically significant ($p < 0.05$) against the proxies of vegetation structure and productivity (except relative fraction of C4 vegetation). While the number of basins with positive GPP-WTD relationships decreases with increasing GPP, LAI, and tree cover (Figures 3a–3c), it increases with increasing grass cover and relative fraction of vegetation with C4 photosynthesis (Figures 3d and 3e).

The stronger association of vegetation characteristics with the difference in the sign of GPP-WTD relationship might result from different connections between WTD, root depth [Fan, 2015; Tron et al., 2014], and distribution [Grimald et al., 2015], as well as from plant physiological factors affecting water use [Rodriguez-Iturbe et al., 2007]. For example, deeper tree roots (compared to those of shrub or grass) provide access to a
deeper soil column [Neill et al., 2013] under both normal and dry conditions [Baldocchi et al., 2004; Weltzin and McPherson, 1997], which increases the potential water uptake. This large water uptake decreases soil moisture and, consequently, groundwater recharge and results in a deeper WTD leading to a negative GPP-WTD relationship. On the other hand, the enhanced water use efficiency of vegetation with C4 photosynthesis [Still et al., 2003] facilitates a higher productivity per additional unit of soil moisture supply from groundwater, which could explain the prevalence of positive GPP-WTD relationship at the highest relative fraction of C4 vegetation (>90%).

The number of both positive and negative GPP-WTD relationships peaks at relatively low precipitation (300–400 mm/yr: Figure 3g) and shallow WTD (<10 m; Figure 3f), while no clear pattern is evident for net radiation (Figure 3h). We observe a statistically significant pattern (p < 0.05) of more negative GPP-WTD relationships with increasing precipitation (Figure 3g), suggesting that WTD deepens due to a large productivity and transpiration loss when the moisture supply from rainfall is abundant. Positive relationships prevail at shallow WTD (6–8 m), where groundwater capillary flux can be expected to enhance root zone moisture and surface water fluxes [Koirala et al., 2014]. Such shallow WTD conditions are particularly necessary for grass and shrub vegetation with shallower roots and limited direct access to groundwater.

Figure 3. Distribution of vegetation, surface, and climate characteristics in basins dominated by positive and negative GPP-WTD relationship. In the left axis, the number of basins (red line for positive and blue line for negative) is plotted against (a) GPP, (b) leaf area index, (c) tree cover fraction, (d) grass cover fraction, (e) relative fraction of C4 vegetation, (f) 25th percentile WTD, (g) mean annual precipitation, and (h) net radiation. In the right axis, the difference in number of basins (positive minus negative) is plotted (grey bars). The P values of the Kolmogorov-Smirnov test of significance between the distributions of positive and negative relationships are provided on the top-right section of each plot. Note the irregular vertical axis limits, as well as the logarithmic horizontal axis in Figure 3f.
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3.3. Relative Importance of Vegetation, Surface, and Climate Characteristics

Finally, we quantify the relative importance of climate (three variables), vegetation (seven variables), and surface (three variables) characteristics for the dominant sign of GPP-WTD correlation. To do so, we first trained a random forest classification algorithm [Breiman, 2001] to predict whether the GPP-WTD correlation is dominated by a positive or negative relationship. The relative importance of the groups of variables was then calculated using two methods; permutation and leave out approaches (see section S10). Irrespective of the metric chosen, vegetation-related predictors have a much larger relative importance than climate and land surface-related predictors in determining the dominant GPP-WTD correlation (Figure 4), which is also evident at the seasonal scale (Figures S13 and S14 in the supporting information). An additional calculation using only two principal components of each group of variables also produces a similar result (Figure S15), showing that the collinearity and difference in the number of variables within each group have little effect on relative importance.

4. Conclusions

This analysis has uncovered patterns of complex but spatially coherent and statistically significant distribution of positive or negative groundwater-vegetation relationships during at least one season in ~72% of the global vegetated land surface. This clearly demonstrates a global relevance of, often excluded, groundwater-vegetation interactions.

Whether groundwater may be supporting vegetation (yielding a positive relationship) or vegetation may be affecting groundwater (yielding a negative relationship) relates with climate, and more importantly, with characteristics of the vegetation itself. Because causality cannot be inferred from correlation coefficients alone, appropriate ecosystem model developments are urgently needed to quantify the groundwater-vegetation interactions at the global scale. It is especially critical because both positive and negative relationships are equally extensive at the current state, but their patterns and extents can be altered due to global climate and land cover changes. It implies that the recent trends and future projections of declining groundwater resources [Richey et al., 2015] and forest cover [Hansen et al., 2013], along with an increasing vegetation water use efficiency due to CO2 fertilization [Keenan et al., 2013], would potentially shift the groundwater-vegetation interactions. The understanding of how groundwater-vegetation interactions respond to changes in climate and anthropogenic influence is crucial to quantify ecosystem services of groundwater and vegetation resources.

References


