Detecting Ecosystem Reliance on Groundwater Based on Satellite-Derived Greenness Anomalies and Temporal Dynamics

S. Contreras
Centre of Pedology and Applied Biology of Segura, Spain

D. Alcaraz-Segura
University of Granada, Spain; University of Almería, Spain

B. Scanlon
The University of Texas at Austin, Texas

E. G. Jobbágy
San Luis Institute of Applied Mathematics (IMASL), Argentina

CONTENTS
13.1 Introduction ............................................................................................................. 284
13.2 Methods .................................................................................................................... 286
  13.2.1 Study Site ........................................................................................................... 286
  13.2.2 Climate and Satellite Dataset for Greenness Anomaly Estimation ....................... 288
  13.2.3 Greenness Timing and Metrics ........................................................................... 290
  13.2.4 Impact of Groundwater on Vegetation Dynamics: A Conceptual Model ............ 291
13.3 Results and Discussion ............................................................................................. 292
  13.3.1 MAP-EVI Regional Function .............................................................................. 292
  13.3.2 EVI Dynamics along a Groundwater Dependence Gradient at the Telteca Site .......... 292
13.1 Introduction

Groundwater-dependent ecosystems (GDEs) play a key role in human development, and are especially relevant in regions with low rates of rainfall, by providing a broad range of ecosystem services such as physical support for wildlife habitats and biodiversity hotspots, control of floods and erosion, regulation of nutrient cycling, or provision of landscape refuges for cognitive development (de Groot et al. 2002; Chen et al. 2004; Eamus et al. 2005; Bergkamp and Katharine 2006; Ridolfi et al. 2007). During the past decade, research on ecology and functioning of GDEs has received a growing interest from the scientific community and from landscape managers. However, in spite of their high intrinsic values, many of these ecosystems have been strongly impacted as a consequence of disruption of hydrological linkages with groundwater resources. This disruption has been generally promoted by excessive rates of groundwater extraction and depletion, for example, Las Tablas de Daimiel and Doñana National Reserves in Spain (Llamas 1988; Muñoz-Reinoso and García-Novo 2005); Swan Coastal Plain in southwest Australia (Groom et al. 2000); desert springs in the Mojave and Great Basin deserts in the United States (Patten et al. 2008); San Pedro River in the United States (Stromberg et al. 1996). It has also been caused by modification of morphology of stream channels or wetlands through dredging or artificial diversions (Ellery and McCarthy 1998) or as a consequence of changes in their water balance due to climatic factors (Murray-Hudson et al. 2006). A better understanding of the functioning and water consumption of GDEs is then critically required to evaluate the ecological services provided by them (Murray et al. 2006; Brauman et al. 2007) and, for developing adaptive management frameworks that reconcile compatible human activities, ecosystem conservation, and their underlying hydrological trade-offs under future scenarios of land use and climate change (MacKay 2006; Barron et al. 2012b).

GDEs are ecosystems that require groundwater inflows to maintain their current structure and functioning and the subsequent delivery of ecosystem services (Hatton and Evans 1997; Murray et al. 2003; Eamus et al. 2006). GDEs may display an obligate reliance requiring a constant groundwater presence, or a facultative one where they adapt their functioning to fluctuating groundwater availability (Murray et al. 2003; Bertrand
According to the aquifer–ecosystem interface relationship, GDEs include (Eamus et al. 2006; Eamus 2009): (a) caves and subterranean-aquatic ecosystems, including karst aquifers and rock-fractured systems; (b) ecosystems dependent on permanent or temporary surface expressions of groundwater, including baseflow riverine, spring, wetland or peatland, and estuarine/marine-shoreline ecosystems; and (c) ecosystems dependent on the subsurface presence of groundwater, also termed “terrestrial GDEs” or phreatophytic ecosystems (Richardson et al. 2011). Other pedological, morphological, hydrological, and biogeochemical criteria have been proposed for classifying GDEs from a functional point of view (Bertrand et al. 2012).

To preserve their ecological integrity and service provision, GDEs require water allocation plans and adaptive management strategies rooted in knowledge about their (a) typology and spatial distribution, (b) quantitative water requirements, and (c) resistance and resilience to natural and human perturbations on their groundwater regimes. A wide range of methodological approaches and techniques are commonly employed to accomplish these three aspects, including remote sensing, water balance analysis, hydrogeological modeling, tracer and isotopic studies, ecophysiological measurements, rooting system characterization, and aquatic fauna sampling (see Richardson et al. 2011 for a synthesis).

Tracking photosynthetic/greenness activity of vegetation using satellite-based indices such as Normalized Difference Vegetation Index (NDVI) or Enhanced Vegetation Index (EVI) offers a relatively inexpensive and effective way to characterize the functioning of riparian/wetland and terrestrial GDEs (Bradley and Mustard 2008; Barron et al. 2012a). These spectral indices are well correlated with aboveground net primary productivity (ANPP) and evapotranspiration (ET) in semiarid regions (Running and Nemani 1988; Paruelo et al. 1997; Jobbágy et al. 2002; Nagler et al. 2005; Guerschman et al. 2009) (see Chapter 18). When they are not influenced by the presence of groundwater and lateral-inflow resources, annual rates of ANPP and ET in those regions are primarily controlled by precipitation and, second, by radiation forcing and its seasonal coupling with rainfall inputs (Specht 1972; Specht and Specht 1989; Ellis and Hatton 2008; Palmer et al. 2010). Because groundwater supplies a more temporally reliable water source for terrestrial ecosystems than rainfall, higher and more stable ET and ANPP rates should be expected in GDEs compared to their nongroundwater ecosystem counterparts (Contreras et al. 2011; O’Grady et al. 2011). Field observations show that leaf area, ANPP, and water availability are closely correlated, supporting the use of satellite-based vegetation indices for the identification and characterization of GDEs, including the quantification of their water requirements at different temporal scales (Nagler et al. 2005; Contreras et al. 2011; Devitt et al. 2011; O’Grady et al. 2011). Field evidence also suggests that even when access to unlimited groundwater resources exists, the productivity of GDEs could be strongly constrained by other limiting resources or processes such...
as incoming energy, nutrient availability, morphological constraints, or disturbances, among others (Eamus et al. 2000; Do et al. 2008). Consequently, it seems that annual primary productivity estimates retrieved from annual summaries of spectral vegetation indexes could not be sufficient to identify and characterize the water requirements of terrestrial GDEs. To solve this potential constraint, complementary assessment of seasonal greenness timing can provide additional and valuable information on the functional response of ecosystems to their environment (Morisette et al. 2009). In these studies, in addition to primary productivity estimates, seasonality and phenology traits are commonly retrieved from annual greenness dynamics of ecosystems to classify and characterize ecosystem functional types, that is, patches of land surface with similar exchanges of matter and energy between the biota and the physical environment (Paruelo et al. 2001; Alcaraz-Segura et al. 2006; Fernández et al. 2010) (see Chapters 9 and 16).

This study aims to evaluate a satellite-based approach for identifying inflow-dependent ecosystems and to detect the type and degree of groundwater reliance of wetland and phreatophytic ecosystems. The approach consists of the complementary analysis of the annual greenness anomalies computed according to Contreras et al. (2011), and land surface phenological metrics retrieved from intra-annual and interannual greenness trajectories. The performance of this approach is tested in the lowlands of the central Monte Desert (Argentina), where a potential gradient of native inflow-dependent ecosystems has been previously identified. Finally, productivity, seasonality, and phenological metrics computed for a representative sample of those types of ecosystems are compared with those extracted from a sample of sites located at an upstream irrigated oasis that exploits surface and groundwater resources for its maintenance.

13.2 Methods

13.2.1 Study Site

The study region covers an area of 87,500 km$^2$ of lowlands ($\leq$ 1000 m a.s.l.) and expands over the central Monte Desert in Argentina between 31° S and 36° S (Figure 13.1). The region is bounded by the Andes Cordillera to the west and by the Sierras Pampeanas to the east. Precipitation in the region ranges from 150 to 400 mm y$^{-1}$, most of it concentrated in the austral summer (from October to March), and mean annual temperature ranges from 13°C to 19°C. Potential evapotranspiration reaches 1400 mm y$^{-1}$ in the driest parts of the study region. A detailed review of the main biophysical and socioeconomic characteristics of the Monte Desert is provided by Abraham et al. (2009), while Villagra et al. (2009) review some of the effects that land use and disturbance factors have had on the dynamics of the natural ecosystems of this desert.
The area is crossed by five major rivers (from north to south: San Juan, Mendoza, Tunuyán, Diamante, and Atuel) with their origins in the Andes Cordillera. After crossing the mountains, these rivers reach the alluvial fans and sedimentary plains of the central Monte Desert to finally discharge into the Desaguadero-Salado River system. Andean rivers are the main sources of water for four large artificial oases located in the foothills of the region (Figure 13.1), with vineyards, olives, and fruit trees being the main crops. These oases represent approximately 90% of the economic activity in the region, and more than 1.5 million people live there. Along their route in the alluvial fans, rivers recharge large unconsolidated aquifers that extend downstream of the artificial oases to reach the lowlands of the region that are covered by sandy plains of fluvial, lacustrine, and eolian origin.

Alluvial and lowland plains at the foothills of the Andes are mainly covered by three types of ecosystems: (a) shrub-steppes dominated by *Larrea* spp. (*jarillales*), (b) open phreatophytic woodlands of *Prospolis* spp. trees (locally known as *algarrobales*), and (c) marshes and wetlands that are along the main rivers. Two of the largest wetland systems are the Rosario system—at the last section of the Mendoza River just before its confluence with the
San Juan River, and the Guanacache system at the end of the San Juan River. Because of the regulation of the river upstream of the irrigated oases and the great water diversion for agriculture, the Mendoza River has an ephemeral hydrological regime downstream from Mendoza city. Riparian vegetation along the distal section of this river and the Rosario wetlands at its end are supplied with surface waters only after intense rainfall events. Nevertheless, the San Juan River has a permanent water regime acting as a constant source of surface water to the Guanacache wetlands. However, in the past decades, similar to the occurrence in the Mendoza River, the discharge values to the wetland system have been strongly affected by hydraulic regulation and irrigation agriculture in the San Juan oasis. Lacustrine vegetation, such as *Scirpus californicus* and *Typha dominguensis*, dominates wetlands but alien species of the genus *Tamarix* are invading those areas more and more because of changes in the water regime of the river and the streams that fed them. Open *Prosopis* woodlands are mostly located in the alluvial plains on soils that are > 90% sand. These woodlands have different structures depending on their reliance on groundwater resources and show higher growth rates and health status in the Telteca National Reserve and the distal section of the Tunuyán River (phreatic level at 6–15 m depth) than at the Nácuñán National Reserve (water table at 70–80 m depth) (Villagra et al. 2005). In the Telteca area, where an extensive dune system dominates, those open woodlands are well developed in the interdune valleys. The strong reliance of these woodlands on groundwater has been demonstrated by isotopic and hydrochemical profiling studies (Aranibar et al. 2011; Jobbágy et al. 2011). Both open woodlands and wetlands have historically provided the local settlements and economies with timber, peat, and charcoal, and food and water, and also with the physical support required for domestic livestock (Villagra et al. 2009).

In the framework of this study, satellite-based metrics of vegetation dynamics were extracted at the Rosario (R_wet) and Guanacache (G_wet) wetland systems and at the open *Prosopis* woodlands located at the Telteca National Reserve and surroundings (T_wood). Vegetation dynamics were equally characterized for a representative sample of irrigated crops located in the Mendoza irrigated oasis (MIO_agr) (Figure 13.1).

### 13.2.2 Climate and Satellite Dataset for Greenness Anomaly Estimation

According to Contreras et al. (2011), we define “greenness anomaly” as the absolute difference between mean annual greenness observed at any pixel of the landscape and a site-specific reference greenness value estimated depending on the local precipitation. For this study, we used the EVI as an indicator of vegetation greenness. The precipitation-based reference greenness value is assumed to be linearly related to mean annual precipitation (MAP) as follows:

\[
\text{EVI}_{\text{ref}} = a \text{MAP} + b
\]  

(13.1)
where $\text{EVI}_{\text{ref}}$ is precipitation-based EVI and $a$ and $b$ are fitted-parameters computed empirically from a quantile regression analysis developed over the observed EVI-MAP scatterplot defined for a set of reference sites. We assumed a linear relationship between EVI and evapotranspiration based on the field data support available for semiarid regions (e.g., Nagler et al. 2005; Guerschman et al. 2009; O’Grady et al. 2011). With such assumption, we are able to estimate the expected EVI value for a vegetation cover that exclusively uses local precipitation and is in equilibrium with long-term precipitation (Boer and Puigdefábregas 2003; Contreras et al. 2008). This condition, in which annual ET approaches the MAP, has been proposed to be reached at our study region for 75th quantile threshold value (Contreras et al. 2011). At an annual timescale, we defined the concept of $\text{EVI}$ anomaly as follows:

$$\text{EVI}_a = \text{EVI}_{\text{ma}} - \text{EVI}_{\text{map}}$$

(13.2)

where $\text{EVI}_{\text{ma}}$ is the observed annual average of EVI computed from satellite images at each pixel, and $\text{EVI}_{\text{map}}$ is the $\text{EVI}_{\text{ref}}$ in Equation 13.1 estimated using the 75th quantile threshold value. From a functional point of view, both metrics, $\text{EVI}_a$ and $\text{EVI}_{\text{map}}$, are considered here as surrogates of primary productivity.

A map of MAP for the region, which is required to estimate $\text{EVI}_{\text{map}}$, was calculated from long-term average monthly values reported in the CRU CL 2.0 dataset (New et al. 2002), which were previously corrected with data from the CLIMWAT 2.0 database (FAO 2006) and local meteorological stations. Maps of precipitation were finally resampled to a 250-m spatial resolution, which is compatible with the satellite data (Contreras et al. 2011).

The EVI MOD13Q1 land product from MODIS Collection 5 (Solano et al. 2010) was extracted for the region (tile h12v12) covering nine hydrological years from September 2001 through August 2009 (23 scenes per hydrological year). Before processing, raw EVI data at 250-m spatial resolution were filtered using a local polynomial function based on an adaptive Savitzky–Golay filter using the TIMESAT software (Jönsson and Eklundh 2004). EVI, which combines data from the blue, red, and infrared spectral bands, was preferable to NDVI because atmospheric interferences and soil background signal are more effectively removed and because of its greater sensitivity to high biomass situations (Huete et al. 2002).

Equation 13.1 was parameterized across 125 reference sites that meet the criteria of having low disturbance rates and lacking artificial or runoff water supplies (Contreras et al. 2008, 2011). From the resulting EVI-MAP scatterplot, three quantile threshold values were used here to propose a preliminary gradient of classes of groundwater reliance. First, as stated earlier, we used the 75th quantile regression of mean annual EVI versus MAP function as a conservative value to generate $\text{EVI}_{\text{map}}$ values. Observed EVI values lower than $\text{EVI}_{\text{map}}$ (i.e., negative greenness anomaly values) were assumed
not to have any dependency on groundwater resources. For values higher than $EVI_{map}$, a gradient of three potential classes of low, moderate, and high degree of reliance were established using the 75th, 90th, and 99th quantile thresholds, respectively. In this study, the former quantile threshold values were arbitrarily selected in order to evaluate the potential agreement between the resulting reliance levels and the phenological metrics extracted from the greenness timing analysis.

### 13.2.3 Greenness Timing and Metrics

A representative sample of pixels for the four study systems (Telteca woodlands, Rosario and Guanacache wetlands, and irrigated crops at the Mendoza oasis) was selected for retrieving metrics, or phenometrics, related to vegetation traits of primary productivity, seasonality, and phenology (Figure 13.2). At the Telteca Reserve, 78 pixels were sampled to cover all the potential groundwater-reliance degrees identified by the anomaly greenness (no reliance, low, moderate, and high) although pixels...
with moderate- and high-reliance degrees were finally grouped together to obtain a more robust comparison among classes. In the wetland systems, vegetation metrics were only extracted from pixels (Guanacache, n = 40; Rosario, n = 10) with a potential high degree of reliance on groundwater. Metrics related to vegetation primary productivity, seasonality, and phenology were extracted at intra-annual and interannual timescales from the average seasonal trajectory resulting from September 2000 to August 2009 (intra-annual variability) and from the annual trajectories for the nine hydrological years covered by the study (interannual variability). In this study, we extracted the following EVI metrics related to traits of: (a) vegetation productivity: \( \text{EVI}_{ma} \) and \( \text{EVI}_{gs} \); (b) vegetation seasonality: \( \text{EVI}_{\text{max}}, \text{EVI}_{\text{min}}, \text{EVI}_{\text{max}} \) and \( \text{EVI}_{\text{nrange}} \); and (c) vegetation phenology: \( L_{gs} \) and \( T_{\text{max}} \) (see Figure 13.2 for more details).

### 13.2.4 Impact of Groundwater on Vegetation Dynamics: A Conceptual Model

From a functional point of view, a water table close to the land surface is expected to impact intra-annual (seasonal) and interannual (multiyear) variability of the EVI dynamics in several ways. The following hypothesis guided our analyses (Table 13.1).

At the intra-annual scale, we hypothesize that MAP (\( \text{EVI}_{ma} \)), the cumulated productivity during the growing season (\( \text{EVI}_{gs} \)), and maximum (\( \text{EVI}_{\text{max}} \)) and minimum (\( \text{EVI}_{\text{min}} \)) values of greenness are expected to increase.

<p>| TABLE 13.1 |</p>
<table>
<thead>
<tr>
<th>Trends in Vegetation Traits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vegetation Traits</td>
</tr>
<tr>
<td>---</td>
</tr>
<tr>
<td>Productivity</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Seasonality</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Phenology</td>
</tr>
<tr>
<td></td>
</tr>
</tbody>
</table>

**Note:** Trends (arrows) measured by satellite-based metrics expected in terrestrial GDEs as groundwater reliance increases. Metrics are related to *productivity* traits: \( \text{EVI}_{ma} \) = mean annual EVI; \( \text{EVI}_{gs} \) = mean EVI accumulated during the growing season; *seasonality* traits: \( \text{EVI}_{\text{max}} \) and \( \text{EVI}_{\text{min}} \) = maximum and minimum EVI values; \( \text{EVI}_{\text{nrange}} \) = annual amplitude of EVI (\( \text{EVI}_{\text{max}} - \text{EVI}_{\text{min}} \)); and *phenology* traits: \( T_{\text{max}} \) = time at which maximum value is reached.
with greater reliance of ecosystems on groundwater. Because shallow water tables represent a perennial source of water for ecosystems, we also hypothesize that vegetation with any reliance on groundwater would show less variable seasonal trajectories of productivity ($\text{EVI}_{\text{range}}$) as groundwater reliance increases. As a consequence of less variability in the greenness trajectory (less $\text{EVI}_{\text{range}}$), a longer period should be required to reach 50% of the total annual productivity, here defined as the growing season length ($L_{gs}$). At the interannual scale, we expect that variability of all vegetation metrics described for productivity, seasonality, and phenology traits should be lower in GDEs than in non-GDEs and should decrease as reliance on groundwater increases. The matrix of conceptual rules proposed here to evaluate ecosystem reliance on groundwater has been designed under the assumption that the access to groundwater by vegetation remained relatively constant without large changes in the water table depth. Then, changes in the water table depth or in the hydrological regime of those ecosystems are expected to be followed by modifications in their greenness dynamics and phenological patterns.

### 13.3 Results and Discussion

#### 13.3.1 MAP-EVI Regional Function

According to the MAP-EVI function described for the region (Figure 13.3), positive EVI anomalies cover 26,000 km² (~30% of the total area) with 36% distributed over the irrigated oases of the region (Table 13.2; Figure 13.1). High positive anomalies represent almost 24% of the total positive anomalies mapped on natural ecosystems/rangelands, but almost 95% of the total area at irrigated oases, which proves the important role that irrigation has on agricultural development in the region.

#### 13.3.2 EVI Dynamics along a Groundwater Dependence Gradient at the Telteca Site

Almost synchronous intra-annual (Figure 13.4) and interannual (Figure 13.5) trajectories of EVI were found in *Prosopis* woodlands located at the Telteca natural reserve, with annual and growing season values higher than those observed at the control sites (sites with no positive greenness anomalies) as EVI anomalies increased (Figure 13.6a). Trends in productivity metrics were also confirmed for seasonality values with higher $\text{EVI}_{\text{max}}$ and $\text{EVI}_{\text{min}}$ values, but lower seasonality variation ($\text{EVI}_{\text{range}}$), as EVI anomalies increased (Figure 13.6b). No significant trends were found, however, for phenological metrics, that is, growing season length ($L_{gs}$) and time at which maximum
FIGURE 13.3 (See color insert.)
Mean Annual Precipitation–Enhanced Vegetation Index (MAP-EVI) quantile regression functions for the study region. Sample of pixels selected at each control area (brown symbols) are embraced by dashed lines. Functions were computed from MAP-EVI values measured at 125 reference sites (black-white circles). EVI thresholds corresponding to 75th, 90th, and 99th quantile regressions are used to classify systems into their low, moderate, and high reliance on water inputs besides local precipitation, respectively.

TABLE 13.2
Negative and Positive Enhanced Vegetation Index (EVI) Anomalies

<table>
<thead>
<tr>
<th>Type of Land Cover</th>
<th>Negative Anomaly</th>
<th>Positive Anomaly</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Low</td>
</tr>
<tr>
<td>Natural ecosystems</td>
<td>61.526</td>
<td>6.416</td>
</tr>
<tr>
<td>or rangelands</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Irrigated oases</td>
<td>265</td>
<td>139</td>
</tr>
<tr>
<td>Total</td>
<td>61.791</td>
<td>6.555</td>
</tr>
</tbody>
</table>

Note: Total area (in km²) with negative and positive EVI anomalies in the study region. Positive and negative anomalies were computed from EVI_map values estimated from 75th quantile regression function. Positive anomalies, which represent portions of landscape with any reliance degree on water inputs besides precipitation, have been divided into low, moderate, and high levels of reliance if EVI is higher than 75th, 90th, and 99th quantile threshold values, respectively.
EVI is reached ($T_{max}$), although average values for both were higher than control sites as EVI anomalies increased (Figure 13.6c). Because patterns and trends predicted by our conceptual model were matched at the Telteca woodlands, it seems that greenness anomaly may be a good surrogate for the reliance that woodland ecosystems have on groundwater resources. In addition to the metrics and trends recorded, a higher increase rate in the greenness trajectory was also observed during the late spring period, from November to December, in sites with moderately high EVI anomalies than in the control sites (Figure 13.4). This “early upraise” makes EVI differences among sites the highest during this seasonal period when energy constraints (low temperatures) start to disappear for phreatophytic woodlands, but rainfall inputs are still low for promoting vegetation growth in their nonphreatophytic counterparts.

Significant differences in EVI annual values were found throughout the entire study period between control sites and sites with moderately high EVI anomalies (Figure 13.5). Although weak absolute differences in EVI were observed, those differences were not significant between sites with
low positive EVI anomalies and control sites. As EVI anomalies increased, interannual variability (measured as the coefficient of variation among annual values of each hydrological year) decreased for growing season EVI (EVI_{gs}), minimum EVI (EVI_{min}) and the intra-annual variability (EVI_{intra}) (Table 13.3). The remaining EVI metrics did not show any clear trends, although values for sites with moderately high EVI anomalies were slightly lower than for control sites (Table 13.3).

### 13.3.3 Intercomparison among Phreatophytic Woodlands, Wetlands, and Irrigated Crops

Wetlands and irrigated sites at the Mendoza oasis showed EVI trajectories clearly different from those of open woodlands of Prosopis (Figure 13.4). Annual productivity (EVI_{ma}) at the wetlands of Guanacache and Rosario and at irrigated sites was 8.7, 9.4, and 12.4 times higher than those measured at the Telteca woodlands, respectively (Figure 13.6a). According to
FIGURE 13.6
Whisker plots showing average and confidence intervals (at 95% level) for (a) productivity metrics ($EVI_{ma}$ and $EVI_{gs}$), (b) seasonality metrics ($EVI_{min-a}$, $EVI_{mix-a}$, $EVI_{range}$), and (c) phenological metrics ($Lgs$, $T_{max}$). Metrics were computed at Prosopis woodlands at Telteca ($T_{wood}$), Guanacache and Rosario wetlands ($G_{wet}$, $R_{wet}$), and the Mendoza irrigated oasis ($MIO_{agr}$). Samples at each system were selected for different levels of reliance on water inputs besides local precipitation: (0: no reliance; +: low reliance; ++/+++: moderate and high reliance). For comparison purposes, all average metrics were computed in terms of their corresponding EVI anomalies.
Phenometrics in Groundwater Dependent Ecosystems

their greenness anomalies, mean annual evapotranspiration rates reported for open *Prosopis* woodlands reached approximately 185 mm y$^{-1}$, of which approximately 25 mm y$^{-1}$ are estimated to be supplied by shallow groundwater reserves (Contreras et al. 2011). These results agree with independent estimates computed from independent isotopic and hydrochemical evidences (Jobbágy et al. 2011). Supplementary water consumption of wetlands is even higher than in open woodlands, with rates that can reach up to 450–500 mm y$^{-1}$ in addition to rainfall inputs. No accurate data exist on the relative contribution of groundwater supplies to the average productivity of the Guanacache and Rosario wetlands, but the observation of different average seasonal EVI trajectories suggests two patterns of ecological functioning: vegetation at the Guanacache wetland is characterized by higher intra-annual variability ($EVI_{\text{range}}$), lower minimum EVI values ($EVI_{\text{min}}$, Figure 13.6b), and a shorter growing season ($L_{gs}$, Figure 13.6c) than at the Rosario wetlands. Although no significant differences in maximum greenness were found between both wetland systems (Figure 13.6b), the time at which they were reached was approximately 16 days earlier at the Guanacache system than at the Rosario system (Figure 13.6c). The lower interannual variability found for all EVI metrics at the Rosario wetlands compared to the Guanacache wetlands would suggest that ecological functioning of the Rosario system relies more on groundwater resources than the Guanacache system. This fact is confirmed by the interannual greenness dynamics at the Guanacache system (Figure 13.5), where abrupt rises and falls in the mean annual EVI values suggest a higher dependence on the water discharges

### TABLE 13.3
Coefficients of Variation for Greenness Metrics

<table>
<thead>
<tr>
<th>Sites</th>
<th>Productivity</th>
<th>Seasonality</th>
<th>Phenology</th>
</tr>
</thead>
</table>
|             | $EVI_{\text{ma}}$ | $EVI_{\text{gs}}$ | $EVI_{\text{max}}$ | $EVI_{\text{min}}$ | $EVI_{\text{range}}$ | $L_{gs}$ | $T_{\text{max}}$
| $T_{\text{wood}}$ | 6.02 | 0.89 | 11.88 | 6.36 | 25.17 | 5.96 | 18.42 |
| (0)         | (0.78) | (0.21) | (2.01) | (0.85) | (3.58) | (1.39) | (5.05) |
| $T_{\text{wood}}$ | 5.61 | 0.69 | 10.67 | 5.86 | 22.56 | 5.05 | 15.67 |
| (+)         | (0.79) | (0.31) | (1.85) | (1.29) | (3.62) | (1.08) | (4.30) |
| $T_{\text{wood}}$ | 6.00 | 0.50 | 10.72 | 5.55 | 22.12 | 5.34 | 18.97 |
| (++/+++/+++)| (1.23) | (0.25) | (3.52) | (1.40) | (5.52) | (1.08) | (5.50) |
| $G_{\text{wet}}$ | 26.52 | 1.36 | 23.31 | 30.49 | 20.43 | 9.46 | 30.65 |
| (+++)       | (15.53) | (1.25) | (12.90) | (19.64) | (12.59) | (4.40) | (12.15) |
| $R_{\text{wet}}$ | 9.62 | 0.51 | 11.75 | 8.20 | 14.49 | 5.80 | 22.65 |
| (+++)       | (3.55) | (0.41) | (5.68) | (2.12) | (5.53) | (1.44) | (7.57) |
| $MIO_{\text{agr}}$ | 7.54 | 1.08 | 9.85 | 12.61 | 16.13 | 7.56 | 26.45 |
| (+++)       | (5.31) | (0.72) | (4.79) | (6.15) | (6.51) | (2.84) | (10.47) |

*Note:* Values were computed from the annual metrics computed from September 2000 to August 2009 (nine hydrological years). Standard deviations of the coefficients of variation (spatial variability observed at each ecosystem type) are shown between parentheses.
supplied by the San Juan River and, consequently, by the water abstractions accounted for irrigation at the upstream San Juan oasis.

Greenness seasonal dynamics observed at the Mendoza irrigation oasis are characterized by its lack of coupling with the rest of the ecosystem types. Irrigated oases showed similar values for productivity metrics (Figure 13.6a) and seasonality (Figure 13.6b) compared to those observed in both wetland systems. However, average intra-annual trajectory of EVI (Figure 13.4) in the Mendoza oasis highlights an earlier maximum vegetation activity and a higher activity during the growing season. The existence of a phase difference between the seasonal EVI trajectories of the irrigated oasis and the natural wetland systems would suggest a competitive process for water resources. This fact was stressed earlier between the San Juan oasis and the Guanacache wetland but would be equally expected for the Mendoza oasis and the Rosario system. In the Guanacache-San Juan case, consequences of water abstraction on the wetland productivity are more clearly depicted because of limited reliance of wetlands on groundwater resources. In the Rosario-Mendoza case, where the Rosario wetlands seem to rely more on groundwater resources, it is expected that consequences of agricultural development on the wetland productivity are less evident during wet or average-rainfall hydrological years than during dry years. A more detailed study identifying those differences during the driest periods would help to identify GDEs and the effects that irrigation development could have on their ecological functioning and the services they provide.

13.4 Conclusions

GDEs offer an outstanding example of the dependence of human well-being on ecosystem services. In this study, we demonstrated the usefulness of the annual greenness anomaly concept (Contreras et al. 2011) to identify landscape systems where vegetation activity depends on abnormally high inputs of water apart from precipitation: that is, riparian ecosystems, wetlands (see Chapter 17), phreatophytic woodlands, and irrigated oases. Particularly in low-precipitation regions, provisioning services (such as biomass or water availability), regulating services (such as the maintenance of lifecycles, habitats, and gene pools, or the local climate regulation), and cultural services (intellectual, spiritual, or recreational interactions with distinctive landscapes) are locally concentrated in these ecosystems, which are tightly coupled to the groundwater dynamics. Annual greenness anomaly estimated from satellite data was demonstrated to be a simple yet robust measurement for mapping those inflow-dependent systems at vast regions with limited field data and for providing a first estimate of their water requirements. Additional information on the reliance of those ecosystems on groundwater
can be obtained from complementary analysis of their EVI intra-annual and interannual trajectories and from the extraction of metrics related to productivity, seasonality, and phenology of carbon gains in those ecosystems (see Chapter 9). In this study, we showed how the average greenness during the growing season, the annual minimum greenness, and the intra-annual variability (normalized range) were higher in phreatophytic woodlands and wetlands than in their nonphreatophytic counterparts. Hence, we suggest the use of these metrics to quantify and map the ecosystem's reliance on groundwater resources and the degree of dependence of ecosystem services on the groundwater dynamics.

Satellite-based approaches based on the spatial analysis of vegetation greenness anomalies and the tracking of their phenologies during time periods explicitly selected to cover dry rainfall conditions provide an initial characterization of natural ecosystems that show any reliance on groundwater resources. Both methods are extremely useful as a first step in building conceptual models on the functioning of GDEs, to quantify their water requirements, and to evaluate the ecosystem services trade-offs that can emerge between their conservation and agricultural development options.

Acknowledgments

S. Contreras acknowledges the support given by a Juan de la Cierva postdoctoral fellowship (JCI-2009-04927) funded by the Spanish Ministry of Science and Innovation. Pablo E. Villagra, Erica Cesca, and three anonymous reviewers are acknowledged for their help in supplying technical data and for their insightful comments.

References


