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Many ecological systems follow a seasonal cycle affecting primary production, carbon flux, and vegetative gas emissions. The seasonal variation of ecological systems are both affected by and have effects upon climatic factors. A quantitative estimate of the seasonal variation of vegetation is required to characterize ecological systems and their interaction with climate. Monitoring the spatiotemporal variation of foliar biomass density (FBD) over one year will provide a quantitative estimate of the annual cycle and regional variation of photosynthetic activity. FBD is a quantitative measure of leafy material per unit of area produced by photosynthetically active vegetation. This seasonal variation in FBD is an important parameter for global and other large scale investigations of ecological, hydrological, and biogeochemical systems which require data and expertise from a variety of sources and disciplines. Therefore, FBD is potentially of great utility for ecologists, hydrologists, climatologists, and atmospheric scientists.

Recent regional scale investigations of ecological systems concluded that the repetitive coverage and synoptic view of remotely sensed measurements provide data to monitor the seasonal variation of biomass. A method to estimate the seasonal variation of FBD at global scales has not been developed. The objective of this research is to develop a methodology that could be used to estimate the
seasonal variation of FBD for the entire terrestrial biosphere. By coupling global satellite data, measured field data, and a vegetation classification, a model was developed to estimate the global spatiotemporal variation of FBD.

Comparisons between literature estimates of FBD and estimated FBD from this model were made as a means of validation. A more specific comparison was conducted between grasslands based on work conducted in the Senegalese Sahel region in Africa. Finally, a sensitivity analysis was performed to characterize the potential propagation of error associated with the literature FBD estimates used to drive this model.

# A Global Scale Analysis of the Spatiotemporal 

 Distribution of Foliar Biomass for 1988by

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I dedicate this thesis to my mother, Jody Pross, who by example, inspired in me an interest in the wonders of nature, and who has given me the perserverance and compassion to follow my present and future dreams to fruition. I also wish to dedicate this thesis to memory of my father, Donald Pross; a man whose life was taken before I had the chance to know him. My grandparents, Bertram and Alice Halberg, provided me with unlimited love and an interest in the "out-of-doors", and I thank them. Without my family, I would be a man of little substance. This thesis is really the accomplishment of my parents and grandparents in raising me as much as it is my personal scientific accomplishment.

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Corvallis, 1991

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# A GLOBAL SCALE ANALYSIS OF THE SPATIOTEMPORAL DISTRIBUTION OF FOLIAR BIOMASS FOR 1988 

## CHAPTER 1

## Introduction

There are many factors, both of natural and human origin, that determine the climate of the Earth. The driving energy for weather and climate comes from the Sun. Of the solar radiation that the Earth intercepts, about one third is reflected and the rest is absorbed by the components of the climate system (e.g. atmosphere, ocean, ice, land, and biota). As energy is absorbed from solar radiation it is balanced by outgoing radiation from the Earth and atmosphere. The temperature of the Earth-atmosphere system results from this balance (Gates, 1980).

There are several natural factors which can change the balance between the energy absorbed and emitted by the Earth. One of the most important important factors is the greenhouse effect. Simply, shortwave radiation can pass through a clear atmosphere relatively unimpeded, but the long wave terrestrial radiation emitted by the warm surface of the Earth is partially absorbed and then re-emitted by a number of trace gases in the cooler atmosphere above. However, if this balance is disturbed then warming or cooling will occur. The main concern over global climate change today is the quantity of water vapor, clouds, and trace gases in the atmosphere and the affect these parameters have upon the mean temperature of the Earth (Houghton et al., 1991).

The key greenhouse gases produced by natural and human activities are $\mathrm{CO}_{2}$, $\mathrm{CH}_{4}, \mathrm{~N}_{2} \mathrm{O}$, water vapor and ozone. All of these trace gases are determined to some degree by photosynthetically active foliar biomass, owing their quantities and locations of sources and sinks in part to the seasonal variation of terrestrial biota. Water vapor has the largest greenhouse effect but, on a global scale, is not affected
by human sources and sinks. The seasonal variation of terrestrial biota does, however, play a significant role in the regulation of the global water balance. Ozone affects incoming solar radiation, but quantifying the climatic effect of ozone change is not yet accurately defined. It is known, however, that biogenic gas emissions, such as isoprene, can affect the concentrations of tropospheric ozone. The sources of $\mathrm{CH}_{4}$ and $\mathrm{N}_{2} \mathrm{O}$ are the least well known but can be related to the burning and decay of biomass. While the anthropogenic sources and magnitudes of $\mathrm{CO}_{2}$ are known, the sources and sinks from the ocean and terrestrial biota are not well known, but it is recognized that the processes of photosynthesis and respiration fix and release carbon. In summary, the seasonal variation of photosynthetically active biota is an important parameter of the global climate system (Rosenzweig and Dickinson, 1986, and Berger et al., 1989).

The interaction between vegetation and climate are affected in both the short and long term. Biophysical processes such as photosynthesis and respiration are dependent on climatic factors and $\mathrm{CO}_{2}$ concentration in the short term. Photosynthesis captures atmospheric $\mathrm{CO}_{2}$, water, and solar energy and stores them in organic compounds which are then used for subsequent plant growth, animal growth, and growth of microbes in the soil. All of these organisms release $\mathrm{CO}_{2}$ via respiration into the atmosphere. Most land plants have a system of photosynthesis which will respond positively to increased atmospheric $\mathrm{CO}_{2}$ but the response varies with species. In the longer term, due to the species response, climate and $\mathrm{CO}_{2}$ are among the factors which control ecosystem structure (i.e. species composition, either directly by increasing mortality in poorly adapted species, or indirectly by mediating the competition between species) (Gates, 1980).

Because species respond differently to climatic change, some will increase in abundance and/or range while others will decrease. Ecosystems will therefore
change in structure and composition. For example, some species may be displaced to higher latitudes and altitudes, and may be more prone to local or global extinction whereas other species may thrive. In other words, ecosystem structure and species distribution are particularly sensitive to the rate of change of climate. As a result, the rate of temperature change can be deduced from the paleoclimatalogical records. These paleoclimatalogical records account for the photosynthetically active seasonal variation of global ecosystems (Rosenzweig and Dickinson, 1986).

A method is needed to characterize present distributions and seasonal variation of photosynthetically active vegetation because of the importance of accounting for this seasonal variation in order to achieve the best possible estimates for the vegetative component of the global climate system. Field sampling of vegetation is one way to estimate distribution and seasonal variation but problems of spatial and temporal sampling are impossible to overcome in the context of global ecosystems. However, by taking advantage of the same solar radiation characteristics that determine the temperature of the Earth-atmosphere system, it is possible to monitor the terrestrial biosphere with satellite instruments that record data in specific spectral windows to retrieve information relevant to the seasonal variation of photosynthetic processes of ecosystems (Rosenzweig and Dickinson, 1986).

Studies which use satellite data to quantitatively characterize the spatial and temporal variation for the entire terrestrial biosphere have not yet been conducted. Nevertheless, there has been a great deal of research performed on the quantification of foliar biomass from watershed to regional scales. These studies, in general, conclude that: 1) satellite imagery can be qualitatively and quantitatively related to foliar biomass, 2) large scale analysis of foliar biomass can be successfully conducted over large regions through time, 3) great care must be taken to normalize and reduce error within the satellite imagery in order that the data
remain reliable through time.
The objective of this study is to develop a methodology that could be used to model the quantitative seasonal variation of foliar biomass for the entire terrestrial biosphere. Coupling the satellite imagery with measured field data and a vegetation ecosystem classification provides the tools for developing a model to estimate the seasonal distribution of foliar biomass. It is hypothesized that: 1) foliar biomass density (FBD - a quantitative measure of the amount of foliar biomass per unit area) can be mathematically estimated with remotely sensed data, 2) these estimates will provide the spatiotemporal distribution of FBD at a monthly time step for one year, 3) the results will provide a realistic characterization of the seasonal varations of photosynthetically active foliar biomass for present ecosytems. These results are compared to other literature based estimates and another model of grasslands in order to test the validity of the model. Finally, a sensitivity analysis is performed to characterize the propagation of potential error associated with the foliar biomass estimates used to drive this model.

## CHAPTER 2

## Background

The research described in this document presents a model that characterizes the seasonal variation of FBD for the entire terrestrial biosphere with a one year time series of global satellite data. The methodology developed for this work was an extension of previous research that has been conducted to understand the physical and physiological basis of spectral response as a function of foliar biomass. The following discussion traces the establishment of both qualitative and quantitative use of these spectral windows for remote sensing of foliar biomass.

> Spectral Windows for Vegetative Remote Sensing

Two portions of the electromagnetic spectrum, the red (0.6-0.7 $\mu \mathrm{m}$ ) and the near-infrared (NIR, $0.7-1.5 \mu \mathrm{~m}$ ), are known to be highly sensitive to foliar biomass. The design of remote sensing instruments used to collect data relevant to vegetation using portions of these spectral windows relies upon the many studies conducted to analyze spectral response as a function of photosynthetically active foliar biomass.

Studies concerning vegetation in the 1960's and 1970's established the fact that different response characteristics of the red and NIR spectral windows are related to photosynthetically active foliar biomass. In one of the earlier studies, Gates et al. (1965) found that different plants display different spectral properties; a phenomenon upon which remote sensing of vegetation is based. Knipling (1970) noticed the spectral differences of plants in the visible ( $0.4-0.7 \mu \mathrm{~m}$ ) and the NIR (0.7-1.5 $\mu \mathrm{m}$ ) and discussed the physical and physiological reasons for the different spectral responses of plants in these two portions of the electromagnetic spectrum. The spectral reflectance and transmittance properties of leaves was researched by Woolley (1971), leading to a better understanding of light interaction
and instrument response. Tucker (1976) analyzed the $0.50-1.00 \mu \mathrm{~m}$ portion of the spectrum to report on the asymptotic nature of grass reflectance as a function of biomass, helping to define appropriate spectral windows for remote sensing purposes. Narrowing the spectral window to $0.750-0.800 \mu \mathrm{~m}$, Tucker (1977) was able to distinguish three quantitative classes of grass biomass. The spectral contribution of post-senescent grass to photosynthetically active grass was conducted by Tucker (1978), adding further to the foundation of spectral response as a function of photosynthetically active vegetation. Linear combinations of red and NIR spectral bands were shown to be highly sensitive to photosynthetically active biomass by Tucker (1979), who concluded that these combinations could be used to monitor biomass. Tucker (1979) reviewed remote sensing and other non-destructive techniques and concluded that satellite spectral methods work well for monitoring foliar biomass and allow for synoptic coverage of large areas.

Spectral Band Ratioing

The spectral windows (bands) recorded for remotely sensed imagery are analyzed by using digital imagery processing techniques. For an overview of digital image processing see Jensen 1986. One of the techniques commonly used in digital image analysis is band ratioing where, based on covariance among the spectral bands, the analyst may wish to ratio these bands together in some mathematical function. Rouse et al. (1973) was one of the first to ratio the red and the NIR to digitally distinguish vegetation types. This band ratio became known as the normalized difference vegetation index (NDVI)(equation 1).

$$
\begin{equation*}
N D V I=\frac{N I R-R E D}{N I R+R E D} \tag{1}
\end{equation*}
$$

where:

NIR = bandwidth corresponding to near-infrared
RED $=$ bandwidth corresponding to red.
Other band ratios have been used but NDVI is one of the computationally simplest and least instrument dependent, providing digital values that are highly correlated and directly related to photosynthetically active FBD. NDVI also minimizes the spatial heterogeneity of an image due to solar variation and topographic effects, increasing its utility for time series analysis. These factors make NDVI a desirable vegetation index. The following research is presented to explain the factors affecting NDVI.

One of the desirable features of the NDVI ratio is the minimization of solar variation across images through time. NDVI was used to effectively compensate for the variation in irradiational conditions through time to monitor photosynthetically active biomass dynamics (Tucker et al. 1979). Kimes (1980) reported on the spatial variability of vegetation canopy reflectance as a function of solar zenith angle. This was a key study for multitemporal image analysis, which is described later in this chapter. The study concluded that diurnal reflectances cannot be clearly understood until the bi-directional measurements of vegetation are more commonly known. However, if satellite data are acquired at the same time each day this problem is minimized. Holben and Justice (1980) used band ratioing as a means to reduce topographic effects on remotely sensed data. Tucker et al. (1981) used simple band ratios, including NDVI, to compensate for variation in solar intensities through time while estimating crop biomass accumulation. Kimes (1984) discovered that the NDVI is significantly less sensitive to solar variations than individual bands for all Sun angles at off-nadir viewing angles of less than $45^{\circ}$.

## Quantitative Applications of Remote Sensing Data

The other desirable feature of NDVI is that it can be used to estimate photosynthetically active foliar biomass. Two measures of foliar biomass are foliar biomass density (FBD) and leaf area index (LAI). FBD is a measure of the amount of leafy vegetation per unit area, and LAI is a measure of total leaf area per unit area. Both measurements quantify the amount of foliar biomass per unit area. The red and NIR spectral windows used in satellite remote sensing are sensitive to amount of photosynthetically active foliar biomass per pixel (when digitally analyzed), where the pixel is a measure of unit area. Clearly, the response characteristics of the red and NIR spectral windows are not changed when a different measurement is used to quantify foliar biomass. Therefore, these spectral response characteristics and the NDVI ratio should be similar when related to FBD and LAI.

An early attempt to assess foliar biomass was performed by Jordan (1969), where in situ measurements of LAI based upon spectral properties of trees were made. Pearson (1976) was one the first to model the relationship between airborne spectral remote sensing data and the amount of biomass using a nine band ratio. The study concluded that the ratio predicted 1.15 times the actual biomass present, with a coefficient of 0.98 for 26 biomass ground-truthed samples. Wiegand et al. (1979) reported one of the first succesful uses of NDVI as a quantitative measure correlating spectral reflectance of biomass to satellite sensors. Holben et al. (1980) discovered that the most significant correlations existed between NDVI and photosynthetically active foliar biomass.

Photosynthetically active LAI was estimated from remote sensing data by Curran (1983) where it was explained that reflectances and radiances in the red are inversely related to the in situ chlorophyll density and the NIR is directly related and proportional to photosynthetically active foliar biomass. Wardley and Curran
(1984) also used remote sensing techniques to estimate photosynthetically active LAI with an accuracy of $50-86 \%$ at the $95 \%$ confidence level. Conducting a timeseries analysis of spectral measurements, Hatfield (1985) discovered that a simple NIR/red ratio remained stable through time and at different locations for wheat.

Remote Sensing Analysis of Large Areas Over Time

Remote sensing can be used to characterize seasonal variation of vegetation provided that the temporal resolution is relatively high. Temporal spectral measurements of crop biomass development were conducted by Tucker et al. (1979) Kimes et al. (1981), and Markham et al. (1981) where significant relationships were found between NDVI and crop biomass development and crop chlorosis, displaying the usefulness of time series NDVI data. Biomass yield variation as a function of remote sensor response over growing seasons was also modeled by Tucker et al. (1980). A similar small scale analysis was conducted by Gallo et al. (1985) where estimates of photosynthetically active radiation in corn canopies were calculated, accentuating the utility of multi-temporal analyses.

Most of the studies up to this point were conducted on small areas and for only one homogeneous vegetation type, such as crops. Temporal analyses had also been conducted on these small areas and a new research direction to study large areas was initiated. The National Oceanic and Atmospherice Association (NOAA) satellites carrying the advanced very high resolution radiometer (AVHRR) provides remote sensing of large regions at a high temporal resolution compared to other remote sensing platforms. The choice is narrowed to one satellite platform when considering a suitable platform for monitoring the seasonal variation within global vegetation ecosystems where a high temporal frequency is needed along with synoptic coverage of large regions of the Earth (Table 1).

Table 1. The satellite and sensor are shown with the corresponding spatial and temporal resolutions and the number of spectral bands associated with the sensor. The NOAA AVHRR satellite platform has 14.5 orbits per day providing daily global coverage with some resampling discussed in the data section. The information in this table came from Jensen (1986).

| System Resolutions |  |  |  |
| :--- | ---: | :---: | :---: |
| Sensor | Spatial | Number <br> of bands | Temporal |
| SPOT Panchrom. | 10 m | 1 | 26 days |
| SPOT MSS | 20 m | 3 | 26 days |
| Landsat TM | 30 m | 7 | 16 days |
| Landsat MSS | 79 m | 5 | 18 days |
| NOAA AVHRR | 1100 m | 5 | 14.5/day |

The AVHRR NDVI data were studied to determine if various vegetation types could be differentiated with the coarse spatial resolution of these data. Norwine and Greegor (1983) stratified various vegetation types using the AVHRR imagery, showing the utility of low resolution satellite data to spectrally distinguish vegetation. Goward (1985) mapped different vegetation types for North America with AVHRR NDVI imagery supporting previous work. These studies show that while the spatial resolution is coarse the high temporal resolution more than makes up for any deficiences the spatial resolution introduces.

A long term regional scale spatiotemporal analysis of portions of Africa using AVHRR NDVI data began in 1980. It was hypothesized that remote sensing can provide invaluable ecological data. Tucker et al. 1983 used the AVHRR NDVI data to statistically model relationships of seasonal variation between the satellite data and grasslands. Tucker et al. 1985 used an NDVI time series to analyze both the spatial and temporal variability of grassland biomass. Tucker et al. (1985) integrated weekly satellite data with respect to time for a twelve month period and produced a remotely sensed estimate of primary production based upon the density and duration of foliar biomass. The dynamics of AVHRR data for Tunisia, Africa, were explained by a combination of vegetation and soil scattering components, and
the NDVI data were also shown to decrease signal variation (Kimes et al. 1985).
Further remote sensing research in Africa stresses the need for high frequency temporal satellite data. Tucker and Sellers (1986) estimated primary production under the limitations of off-nadir viewing and atmospheric conditions. These limitations coupled with the need to measure changing surface conditions emphasized the need for multitemporal measurements. Townshend and Justice (1986) showed NDVI response to different vegetative cover types has unique temporal profiles per vegetation type. An estimate of length of a growing season was made using time series NDVI in East Africa by Justice et al. (1986) by distinguishing levels of foliar biomass.

Holben and Fraser (1984) noted that cloud contamination, directional reflectance, off-nadir viewing, sun-angle, and shadow effects decrease values of NDVI, leading Holben (1986) to develop the maximum value composite procedure. The computations in this procedure retain the maximum value on a pixel by pixel basis for a number of images acquired for the same area of interest on different dates thereby providing the clearest and least shadowed view of the surface. Justice and Hiernaux (1986) showed the utility of high temporal resolution for monitoring at regional scales and emphasized the importance of the maximum value composite procedure. More recently Gutman (1987) reported that the maximum value composite procedure for AVHRR NDVI data is useful for minimizing cloud contamination, atmospheric scattering and absorption, and solar angle effects in the Great Plains region of the Western U.S.

Prince and Astle (1986) concluded that equations could be constructed to predict biomass, but only if the satellite data/vegetation relationship is applied to the same vegetation type by stratifying the region with a vegetation map. Prince and Tucker (1986) built regression models between the Senegalese grassland
biomass and NDVI which explained $93 \%-99 \%$ of the variation in NDVI. A qualitative approach suggested that by monitoring rangeland conditions with satellite data one could identify areas of deficiencies in primary production and provide synoptic information in support of regional drought monitoring (Tucker et al. 1986). Growing periods were defined by periods of minimum biomass and related to minimum NDVI values by Henricksen and Durkin (1986), emphasizing the ability of NDVI to be used to quantify measures of foliar biomass.

## Remote Sensing and Ecological Models

More recent studies have revealed the potential of the seasonal variation of the AVHRR NDVI as an input for ecological models needing spatiotemporal measures of foliar biomass. Running et al. (1986) reported on the first attempt to measure LAI of coniferous forests using satellite data. The study concluded that satellite derived measures of vegetation cover type and LAI may be used to provide more direct estimates of the carbon content and exchange rates of global vegetation than are possible with current data. Running (1988) related AVHRR NDVI to photosynthesis and transpiration of forests in different climates to assess carbon fixation in relation to the global carbon budget. Simulated photosynthesis and evapotranspiration was modeled by coupling the AVHRR NDVI data (used to determine LAI) with an ecosytem model (Running et al. 1989). Spanner et al. (1990) was able to monitor the seasonal variation of LAI in coniferous forests with AVHRR NDVI data. Goward (1989) states that foliar presence determines local rates of photosynthesis, affects surface albedo, and influences local rates of evapotranspiration as well as other elements of surface energy/mass balance. The report concludes that for the first time, through satellite acquired imagery, a consistent, global means to directly study interactions between climate and vegetation exists. The accuracy of the AVHRR NDVI data at a resolution of 1 km was assessed by

Box et al. (1989), who stated that NDVI data were relatively reliable for primary productivity except in areas of complex terrain, for seasonal values at high latitudes, and in extreme deserts. The study also concluded that total biomass (inclusive of woody material) was poorly correlated to the NDVI data. Primary productivity is associated with the amount of foliage present, lending credence to the reliability of using NDVI to estimate FBD.

## Global Vegetation Index (GVI) Data

The global vegetation index consists of resampled AVHRR NDVI imagery providing weekly global coverage. Malingreau (1986) determined that the GVI product provides a large set of useful information on ecosystem dynamics and cropping practices can be consistently derived from these time series data. Singh (1988 a,b) found that the GVI data remain reliable for high, medium, and low foliar biomass with solar zenith angles of less than $80^{\circ}$. The study also concluded that areas above solar zenith angles of $80^{\circ}$ make the GVI data unreliable because the optical depth increases such that the reflectance in the red and NIR is actually a measure of the atmosphere and not the land surface. Holben (1986) refered to this phenomena as the terminator effect. From this fact it was shown that for one GVI image a multitemporal composite image must be made from about four months of imagery. However, the areas of high solar angles lie in a hemisphere's winter during which there is little photosynthetic activity. Lloyd (1990) reported that the GVI data make possible, for the first time, a phenological approach in which classes are defined in terms of the timing, the duration, and the intensity of photosynthetic activity which minimizes this problem of reliability if one is interested only the photosynthetically active foliar biomass.

Goward (1990) cautions against using the GVI data for quantitative purposes due to reasons that Singh (1988 a, b) addresses, but provides methods and ideas to
make the data more reliable. The study stated that it is possible to reduce instrument calibration and off-nadir viewing angle error by approximately $10 \%$ with maximum value composite images of at least a one month time resolution. The conclusion suggested that if these errors were corrected a global measure of vegetation green foliage dynamics can be made within a precision of $10 \%( \pm 0.1 \mathrm{NDVI})$ at a monthly time resolution.

It has been shown that the NDVI ratio can be used to derive information about photosynthetic activity. The NDVI ratio has been used for both small and large scale analyses and it has been used successfully for multitemporal remote sensing studies. The maximum value composite procedure of the NDVI data minimizes effects of atmosphere and solar variation. There are, however, limitations which must be considered for large and global scale remote sensing analysis. Nevertheless, it is possible to make quantitative estimates of photosynthetic acivity at a global scale for the entire terrestrial biosphere.

## Foliar Biomass Estimates and Seasonal Variation

These quantitative estimates derived from the GVI data will be compared to other estimates of global foliar biomass found in the literature. One estimate of the total amount foliar biomass was found in the literature, and two other global estimates of leaf biomass were calculated by using other global estimates of biophysyical variables in the literature. Estimates of leaf area and chlorophyll exist (Whittaker 1975), as well as relationships between leaf biomass and leaf or leaf area or chlorophyll for the various biomes considered in this research (Whittaker 1962, Whittaker 1966, Whittaker and Woodwell 1969, Whittaker et al. 1974, Whittaker and Niering 1975, Lieth and Whittaker 1975, Blasco and Tassy 1975, Edwards and Grubb 1977, Edwards 1977, and Grubb 1977).

One final parameter must be taken into account before building the model. Most biomes occur in both hemispheres and in the Tropics (with a capital Treferring to the Tropics of Cancer and Capricorn). Biomes which occur in the northern and southern hemisphere must be analyzed separately by hemisphere because the seasonal variation is offset (i.e. during January the Northern Hemisphere experiences winter while the Southern Hemisphere is in summer). Further, it is clear that the tropics display a seasonal variation that is different than extratropical areas. The geographic classification of tropics is closely aligned with the Tropics due to characteristics of solar angles. The total amount of solar radiation received at any place depends on two factors: the duration and the intensity of insolation. Insolation, preciptation, and wind play key roles in the determination of climate, and hence, the determination of seasonal variation of vegetation within the tropics (Nieuwolt 1977). Therefore, the biomes will be stratified into a global category (e.g. Tropics, Northern or Southern Hemisphere) in order to analyze the seasonal variation of each biome.

The following chapters discuss the methodology to estimate photosynthetically active FBD by using a yearly time series of GVI, a vegetation map, literature values of FBD, and a stratification for seasonal variation to estimate photosynthetically active FBD, as well as validation procedures to assess the results.

## CHAPTER 3

## Methodology

Data
The modeling and analysis was performed in a raster based geographic information system (GIS). A raster based GIS provides the appropriate environment for modeling spatial data (Burrough, 1986). The GIS modeling and statistical analysis of these global data sets required approximately 200 megabytes of on-line memory and the CPU power to adequately run the algorithms.

The GVI satellite imagery used in this research was acquired from ACECERL in Geographic Resource Analysis Support System (GRASS) (ACE, 1988) raster format for the year 1988 in a weekly time series. The data originated at the National Oceanic and Atmospheric Association (NOAA) whose satellites carry the Advanced Very High Resolution Radiometer (AVHRR) from which daily global coverage is generated across a $56^{\circ}$ viewing track. Spatial resolutions are 1.1 km local area coverage (LAC), and $\sim 4 \mathrm{~km}$ global area coverage (GAC). Global daily coverage is available only through the GAC data that are generated on board the satellite by resampling the LAC data (Kidwell, 1984), where pixels with greater than $30^{\circ}$ off nadir view are not used (Goward et al., 1990). Further sampling, by taking the last pixel mapped into the new grid of 15 km or more in size (Townshend and Justice, 1986), and conversion to the NDVI results in the GVI product as described by Kidwell (1990). GVI data are spatially restricted to land masses between $75^{\circ}$ north and $55^{\circ}$ south with a final nadir resolution of approximately 8.5 minutes (approximately $256 \mathrm{~km}^{2}$ ). The GVI ratio uses the red ( $0.58 \mu \mathrm{~m}-0.68 \mu \mathrm{~m}$ ) and NIR $(0.725 \mu \mathrm{~m}-1.10 \mu \mathrm{~m})$ spectral bandwidths from the AVHRR instruments.

A maximum value compositing procedure (MVC) is used to minimize effects of topography, solar zenith and azimuth angles, and atmosphere on the NDVI ratio. The effects of topography, solar zenith and azimuth, and atmosphere have a tendency to reduce the NDVI value (Holben and Fraser, 1984). Therefore, by retaining the maximum value for each pixel these effects are minimized.

The MVC procedure used to compile the GVI data means that the pixel least affected by cloud or other atmospheric interference is automatically selected for a given seven day time period (Holben, 1986). Further, the GAC pixel with the highest value is used to represent the entire GVI pixel. The MVC procedure was used in this research to obtain monthly GVI images (Figures 1-2), to avoid atmospheric interference, and to retain spatial heterogeneity (Gutman, 1987, Goward et al., 1990). See Appendix A for all twelve monthly images.

Version WE3.0 of the Olson Database of World Ecosystems was used to distiguish different vegetation types in this study, as discussed by Prince and Asite (1986). This global vegetation database was chosen over others because it offered the highest resolution and because of its classification format. The spatial resolution of 30 minutes is much more coarse than the GVI data, but it offers the needed description of vegetation types at the highest possible resolution. The Olson classification is based on landscape and ecosystem while others like Matthews (1983) are based on potential vegetation excluding anthropogenic influences. The database consists of fifty-four categories of ecosystem/vegetation types at a resolution of 30 minutes which covers the entire earth (Olson et al., 1983, Olson 1990).

As discussed, the seasonal variation of foliar biomass is an important parameter to consider when characterizing photosynthetic activity. Because it is impossible to collect enough field data to characterize the seasonal variation of foliar biomass for the entire terrestrial biosphere, previous research from others must be


Figure 1. January GVI Image


Figure 2. July GVI Image
used. Measurements of FBD for large scale biomes were found in Box (1981) Cannell (1982), and Webb et al. 1983. The data found in these sources supplied the maxima and minima of FBD for large scale biomes. This range represents spatial heterogeneity expected of FBD in the biome at peak growth periods.

## Approach

The approach taken to model the spatiotemporal patterns of FBD for one year was to scale the range of FBD values to the range of GVI per biome defined with the Olson map. A general equation of empirical relationships which describes NDVI in terms of FBD exist based on regression analyses. There was a similarity amongst the individual empirical relationships for different vegetation types which confirms the reliability of NDVI for different vegetation types over time. The following discussion details the methodology of this approach.

Empirical relationships between AVHRR NDVI and FBD or LAI exist (for examples see: Tucker et al. 1983, Asrar et al. 1984, Wardley and Curran 1984, Tucker et al. 1985, Hatfield et al. 1985, Running et al. 1986, Peterson et al. 1987, Running et al. 1989, Spanner et al. 1990). Because FBD and LAI appear spectrally similar in the red and NIR bandwidths, they will also appear similar in the NDVI ratio, and therefore, a regression analysis between FBD, LAI and NDVI will also be quite similar. Regression analysis of both FBD and LAI in grasslands and conifer forests display similar response curves (figures 3-4).

They also have identical equations with different empirical constants (Tucker et al. 1983, Running et al. 1989). Representing the empirical constants with the variables $a$ and $b$, the similarities become more apparent (equations 2 a and 2 b ).

Tucker's model equation (see figure 3):

$$
\begin{equation*}
Y=a \ln X-b \tag{2a}
\end{equation*}
$$



Figure 3. Statistical relationship between AVHRR NDVI and the foliar biomass of grasslands. Modified from Tucker et al. (1983).

Running's model equation (see figure 4):

$$
\begin{equation*}
Y=a \ln \frac{X}{b} \tag{2b}
\end{equation*}
$$

where:

$$
\begin{aligned}
& \mathrm{Y}=\mathrm{NDVI} \\
& \mathrm{X}=\mathrm{FBD} \\
& \mathrm{a}, \mathrm{~b}=\text { empirical constants. }
\end{aligned}
$$

Comparing the two models, $a$ remains the same while $b$ is negative in Tucker's model and a negative in in Running's model. This implies that different vegetation types have a similar response curve relating AVHRR NDVI to the similar biophysical varibles; FBD and LAI. Further, the equation defining the response curve


Figure 4. Statistical relationship between AVHRR NDVI and LAI of Coniferous forests. Modified from Running and Nemani (1988).
remains the same while the values for $a$ and $b$ change. This thenomena suggests that the mathematical equation describing the response curves remains the same for FBD of different vegetation types. The empirical constansts, however, will change and may be thought of as variables in this case. Since the general form of the model equation remains constant, the equation may be manipulated to estimte FBD based on the AVHRR NDVI ratio, or GVI. Therefore, derivation of $a$ and $b$ based upon GVI will provide unique relationships between FBD and GVI for different vegetation types.

The model presented here is constructed by using the general form of these of equations which, for simplicity, is taken to be equation (2b). Inverting equation (2b), the FBD variable becomes dependent, or in other words, it may be predicted
(equation 3 ).

$$
\begin{equation*}
X=a \exp \frac{Y}{b} \tag{3}
\end{equation*}
$$

where:

$$
\begin{aligned}
& \mathrm{Y}=\mathrm{NDVI} \\
& \mathrm{X}=\mathrm{FBD}
\end{aligned}
$$

$$
\mathrm{a}, \mathrm{~b}=\text { biome specific constants. }
$$

As previously discussed, the range of FBD per vegetation type supplies only the maxima and minima to which the GVI data can be scaled. Derivation of the biome specific constants must be calculated to define the parameters of the model on a per vegetation type (biome) basis. Two new equations derived from equation 2 describe the biome specific constants in terms of both the biomass and GVI minima and maxima (equations $4 a$ and $4 b$ ), respectively.

$$
\begin{align*}
& F \min =a \exp \frac{N \min }{b}  \tag{4a}\\
& F \max =a \exp \frac{N \max }{b} \tag{4b}
\end{align*}
$$

where:
$\mathrm{Fmin}=\mathrm{FBD}$ minimum value per biome
Fmax $=F B D$ maximum value per biome
Nmin $=$ GVI minimum value per biome
Nmax $=$ GVI maximum value per biome
$\mathrm{a}, \mathrm{b}=$ biome specific constants.
In order to solve for the biome specific constants $a$ and $b$ equations 4 a and 4 b can be simultaneously solved to create equations 5 a and 5 b.

$$
\begin{align*}
& b=\frac{N \max -N \min }{\ln (F \max / F \min )}  \tag{5a}\\
& a=F \max \exp \frac{-N \max }{b} \tag{5b}
\end{align*}
$$

See appendix B for full the solution and appendix C for the biome specific parameters and constants. The model is now calibrated and can be driven by the monthly GVI images to create monthly FBD surfaces.

## Methods

The components of the model are now complete. The seasonal variation and the vegetation types will be stratified, the literature values of FBD will be used to parameterize the model, and the range of GVI values will be used to estimate the spatial heterogeneity of FBD by applying equation 3 on a per biome basis. The first step, however, is to equate the biomes described in the literature to the Olson classification. The Olson classification defines fifty-four categories which were aggregated into eighteen categories representing the literature biomes (Table 2).

The range of FBD in the literature describes the spatial heterogeneity one may find under normal peak growing conditions (see Chapter 2). Because GVI is positively related to FBD, the monthly image with the highest GVI values for each biome can be used to represent the time of peak growth. The month with the maximum mean value for each biome was used to represent the highest GVI values. These months were established by tracking the monthly distributions of GVI for the year (figure 5). These maximum mean data sets for each biome provided the range of GVI values to calibrate the model with the FBD values.

Table 2. Th. irst aggregation scheme based soley on literature descriptions of vegetation type. The second column displays the number of Olson categories which were aggregated into the global areal extents shown in the last column.

| First Aggregation of Olson Categories and their Areal Extent |  |  |  |
| :---: | :---: | :---: | :---: |
| Aggregated Category $\qquad$ | Olson <br> Cats. used | Areal Extent in $\mathrm{km}^{2}$ | Percent <br> Area * |
| Water; no data | 1 | 32,366,200 | 0.00 |
| Ice | 2 | 1,247,000 | 1.96 |
| Desert | 6 | 18,406,600 | 11.97 |
| Tundra | 4 | 10,057,700 | 13.96 |
| South Temperate BLF | 1 | 714,800 | 0.51 |
| Grassland | 3 | 21,356,200 | 14.16 |
| Farms/Towns | 2 | 12,260,700 | 8.69 |
| Nonpaddy Irrigated Dryland | 3 | 1,579,000 | 1.07 |
| Forest/Fields/Woods | 4 | 9,196,300 | 6.60 |
| North Temperate BLF | , | 786,900 | 0.64 |
| Cool Conifer Hardwood | 2 | 3,550,900 | 2.70 |
| Tropical Montane | 1 | 1,175,400 | 0.66 |
| Wetlands/Hinterlands/Shore | 7 | 3,578,700 | 2.60 |
| Woodlands | 6 | 19,902,500 | 11.62 |
| Warm Conifer | 1 | 399,200 | 0.28 |
| Paddyland | 1 | 1,994,100 | 1.19 |
| Taiga | 5 | 11,489,500 | 12.94 |
| Trop. Seasonal Humid BLF | 1 | 6,173,800 | 3.47 |
| Trop/Subt Humid BLF | 2 | 4,237,200 | 2.34 |
| Cool Conifer | 1 | 3,102,100 | 2.63 |
| Total |  | 454,871,000 | 100.00 |

[^0]

Figure 5. The temporal profile of GVI for the Northern Temperate Broadleaf Forest. (see appendix D for the temporal profiles of all the biomes.)

The possibility exists, however, that two or more of Olsons original vegetation types may have been classified as one due to descriptive ambiguity of the definitions for the vegetation types. These categories were digitally analyzed to determine if the classication agreed with a spectral classification using GVI. Histograms of GVI were produced for the fifty-four original Olson categories and the aggregate biomes (Table 2). These distributions of GVI were used to examine the original classifications and the aggregates. If the distribution of GVI is clearly bimodal in the aggregations then the possibility exists that there are two spectrally distinct vegetation types as defined by the original Olson classification.

Bimodality was noticed for the woodlands (figure 6), grasslands, and the desert biomes suggesting the aggregation scheme incorporated dissimilar biomes.


Figure 6. The histogram for the Woodlands biome displaying the bimodality associated two distinct spectral signatures.

Checking these aggregates where bimodality occurred against histograms of the original Olson biomes, two distinct groups could be separated representing each mode (figures 7a-b and figures $8 \mathrm{a}-\mathrm{d}$ ).


Figure 7a. Olson biome number 32, skewed left with a mean of 26.0.


Figure 7b. Olson biome number 43, skewed left with a mean of 26.2.


Figure 8a. Olson biome number 46, skewed right with a mean of 17.4.


Figure $\mathbf{8 b}$. Olson biome number 47 , skewed right with a mean of 16.4.


Figure 8c. Olson biome number 48 , skewed right with a mean of 7.5 .


Figure 8d. Olson biome number 59, skewed right with a mean of 14.9 .
The same analysis separated the grassland and desert biomes into two new distinct classes as well. These new groups were then added to the aggregation scheme
resulting in the new aggregates (Table 3).
Table 3. The second aggregation scheme based on the analysis of GVI histograms. The second column displays the number of Olson categories which were aggregated.

| Second Aggregation of Olson Categories and their Areal Extent |  |  |  |
| :---: | :---: | :---: | :---: |
| Aggregated Category <br> Names | Olson Cats. used | Areal Extent in $\mathrm{km}^{2}$ | Percent Area * |
| Water; no data | 1 | 32,366,200 | 0.00 |
| Ice | 2 | 1,247,000 | 2.00 |
| Nonpolar Desert | 4 | 2,252,700 | 1.93 |
| Polar Desert | 2 | 16,153,900 | 10.30 |
| Tundra | 4 | 10,057,700 | 12.14 |
| South Temperate BLF | 1 | 714,800 | 0.52 |
| Grassland | 2 | 4,084,200 | 3.41 |
| Shrubland | 1 | 17,272,000 | 11.06 |
| Farms/Towns | 2 | 12,260,700 | 8.87 |
| Nonpaddy Irrigated Dryland | 3 | 1,579,000 | 1.09 |
| Forest/Fields/Woods | 4 | 9,196,300 | 6.73 |
| North Temperate BLF | 1 | 786,900 | 0.66 |
| Cool Conifer Hardwood | 2 | 3,550,900 | 2.76 |
| Tropical Montane | 1 | 1,175,400 | 0.68 |
| Wetlands/Hinterlands/Shore | 7 | 3,578,700 | 2.66 |
| Dry Forest and Woodlands | 2 | 11,427,500 | 6.59 |
| Semi-arid Woodlands | 4 | 8,474,900 | 5.28 |
| Warm Conifer | 1 | 399,200 | 0.29 |
| Paddyland | 1 | 1,994,100 | 1.22 |
| Taiga | 5 | 11,489,500 | 13.22 |
| Trop. Seasonal Humid BLF | 1 | 6,173,800 | 3.54 |
| Trop/Subt Humid BLF | 2 | 4,237,200 | 2.39 |
| Cool Conifer | 1 | 3,102,100 | 2.69 |
| Total |  | 454,871,000 | 100.00 |

* The water class was excluded from the percentage calculations.

The aggregated version of Olson vegetation types was then used to group the GVI data into unique spatial categories corresponding to the biomes of the literature.

These biomes were then stratified into global categories (e.g. Northern Hemisphere, Southern Hemisphere, or the Tropics) that represent distinct seasonal variation within each geographic area. The areal extents, by percentage, of biomes in the northern hemisphere and in the Tropics were used to determine in which category the majority of the biome occurred (Table 4). The area where the majority
of the biome occurred was used to calibrate the model by extracting the range of GVI values from inside the geographic constraints of these global categories.

Table 4 . Percentage of biome located in global categories of unique seasonal variations.

| Percentage of Biome in the Tropics, Northern, and Southern Hemsiphere |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: |
| Aggregated Category |  | Tropics | Northern <br> Nemi. |  |
| Nonpolar Desert | 1.8 | Southern <br> Hemi. |  |  |
| Polar Desert | 35.7 | 66 | 32.2 |  |
| Tundra | 1.1 | 64.5 | 0.8 |  |
| Southern Temperate BLF | 0.0 | 50.1 | 38.8 |  |
| Grassland | 0.0 | 96.7 | 42.3 |  |
| Shrubland | 42.4 | 42.8 | 14.3 |  |
| Farms/Towns | 25.7 | 63.8 | 10.5 |  |
| Nonpaddy Irrigated Dryland | 19.8 | 68.6 | 11.6 |  |
| Forest/Fields/Woods | 25.3 | 64.9 | 9.8 |  |
| Northern Temperate BLF | 0.0 | 93.8 | 6.2 |  |
| Cool Conifer Hardwood | 10.9 | 82.0 | 7.1 |  |
| Tropical Montane | 91.2 | 8.8 | 0.0 |  |
| Wetlands/Hinterlands/Shore | 42.8 | 45.6 | 11.6 |  |
| Dry Forest and Woodlands | 97.1 | 1.4 | 1.5 |  |
| Semi-arid Woodlands | 47.1 | 18.7 | 34.2 |  |
| Warm Conifer | 5.2 | 92.3 | 2.5 |  |
| Paddyland | 51.4 | 48.2 | 0.4 |  |
| Taiga | 0.0 | 100 | 0.0 |  |
| Trop. Seasonal Humid BLF | 94.6 | 2.9 | 2.5 |  |
| Trop-Subt Humid BLF | 99.7 | 0.3 | 0.0 |  |
| Cool Conifer | 0.0 | 100 | 0.0 |  |

For example, $96.7 \%$ of the grassland biome is present in the northern hemisphere, therefore, all the GVI data extracted to characterize grasslands was taken from the northern hemisphere. The stratification allows for a more realistic characterization of seasonal variation rather than arbitrarily dividing the biome at the equator and analyzing two different data sets. The biomes are now geographically restricted to one of the global categories for the rest of the calibration procedure.

Again the mean of the GVI values per biome, along with the fifth and ninetyfifth percentile, were tracked by month for the year to determine the peak growth period. The fifth and ninety-fifth percentiles were chosen to represent the entire range of GVI to remove any noise associated with the two distinct boundaries caused by the resolution differences between the Olson map and the GVI imagery
(Table 5).
Table 5. The biomes with ranges of GVI values used to calibrate the model. Also, the global category in which the range occurred is shown as well as the maximum mean GVI value and the month when the maximum mean occurred.

| Biome | Global <br> Category | Month of <br> Max. Mean | Maximum <br> Mean | 5th <br> \%ile | 95 th <br> \%ile |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Nonpolar Desert | North | Jun | 19.5069 | 5 | 36 |
| Polar Desert | North | Mar | 8.0313 | 4 | 12 |
| Tundra | North | Jul | 21.5753 | 6 | 38 |
| Southern TemperateBlf | North | Jun | 31.1673 | 8 | 46 |
| Grassland | North | Jul | 26.1649 | 10 | 43 |
| Shrubland | North | Aug | 17.0670 | 4 | 35 |
| Farms/Towns | North | Jul | 27.6886 | 9 | 42 |
| Nonpaddy Irrigated Dryland | North | Aug | 19.4412 | 4 | 37 |
| Forest/Fields/Woods | North | Jun | 32.5523 | 11 | 46 |
| Northern Temperate Blf | North | Jun | 39.5085 | 20 | 50 |
| Cool Conifer Hardwood | North | Jun | 32.2999 | 11 | 46 |
| Tropical Montane | Tropic | Nov | 32.0610 | 15 | 45 |
| Wetlands | North | Jul | 27.7999 | 7 | 42 |
| DryForest and Woodland | Tropic | Nov | 25.7458 | 10 | 42 |
| Semi-arid Woodland | South | Jan | 19.5504 | 8 | 36 |
| Warm Conifer | North | Jun | 29.7677 | 14 | 41 |
| Paddyland | North | Jul | 26.4018 | 10 | 38 |
| Taiga | North | Jul | 29.3820 | 20 | 40 |
| Trop. Seasonal Humid BLF | Tropic | Nov | 34.6479 | 22 | 45 |
| Trop/subt Humid BLF | Tropic | Dec | 34.5576 | 20 | 44 |
| Cool Conifer | North | Jul | 33.0470 | 17 | 45 |

Ranges of FBD from the literature (Table 6) were then scaled to the GVI by equation 2. See appendix C for the derived constants.

For values of GVI that are greater than the value of the 95 th percentile, FBD is set to a constant value in order to eliminate anomalous occurrences of exceedingly high GVI values that would convert to exaggerated FBD values. This upper FBD limit was specific for each biome. Using the model equation response curves displaying FBD in terms of GVI were calculated (figure 9).

Table 6. The maximum and minimum values for FBD expressed in $\mathrm{kg} / \mathrm{m}^{2}$.

| Biome | Mimimum <br> FBD | Maximum <br> FBD |
| :--- | :---: | :---: |
| Nonpolar Desert | 0.01 | 0.05 |
| Polar Desert | 0.01 | 0.10 |
| Tundra | 0.01 | 0.05 |
| Southern Temperate BLF | 0.2 | 0.8 |
| Grassland | 0.05 | 0.5 |
| Shrubland | 0.1 | 0.5 |
| Farm/Town | 0.1 | 0.5 |
| Nonpdy Irrigated Dryland | 0.1 | 0.5 |
| Forest/field/woods | 0.1 | 0.8 |
| Northern Temperate BLF | 0.3 | 0.6 |
| Cool Conifer Hardwood | 0.3 | 1.0 |
| Tropical Montane | 0.3 | 0.7 |
| Wetlands | 0.1 | 0.5 |
| Dry Forest and Woodland | 0.1 | 0.5 |
| Semi-arid Woodlands | 0.1 | 0.3 |
| Warm Conifer | 0.1 | 1.0 |
| Paddyland | 0.1 | 0.5 |
| Taiga | 0.3 | 1.5 |
| Trop Seasonal Humid BLF | 0.3 | 0.8 |
| Trop/subt Humid BLF | 0.3 | 1.3 |
| Cool Conifer | 0.5 | 2.0 |



Figure 9. The calibration curve for the Cool Conifer boime derived from the literature. See appendix E for all the calibration curves for the biomes.

These biome specific relationships were used to spatially distribute the estimates of FBD for each of the monthly GVI images in the GRASS raster GIS environment. GVI has now been used to estimate FBD values between $75^{\circ} \mathrm{N}$ and $55^{\circ} \mathrm{S}$ for the terrestrial biosphere in a monhly time series for twelve months. These FBD surfaces display values from $0-2.0 \mathrm{~kg} / \mathrm{m}^{2}$ simulating the seasonal variation of GVI values (Appendix F).

## CHAPTER 4

## Results

Results and Discussion
Spatiotemporal patterns of FBD that follow characteristics of seasonal variation can be observed for the terrestrial biosphere. The seasonal variation is clear when comparing the January FBD surface (Figure 10) to the July FBD surface (Figure 11). For example, the northern hemisphere in the January surface is void of any high values of FBD while the southern hemisphere and the Tropics contain the high FBD values. The July surface shows the opposite situation with the equatorial region showing high values again.

The MVC procedure used to make these monthly images highlights a problem caused by solar angles that present a limitation of this procedure. The GVI images during winter in the northern hemisphere, particularly January (Figure 10) have a band of high values in the extreme northern latitudes over the terrestrial biosphere. The solar angles are so low that the optical depth of the atmosphere is increased. The satellite sensor at this point is essentially recording atmospheric reflectance rather than surface reflectance. Applying the MVC procedure retains the highest noise values and creates bogus data in these high latitudes during times of low solar zenith angles. Holben (1986) referred to this phenomena as the terminator effect. Because the GVI data are the NDVI ratio, they cannot be broken apart into the original red and NIR values which could be corrected. This is a limitation of using just the NDVI ratio without having the red and NIR spectral bands to analyze.

The total foliar biomass calculated for the terrestrial biosphere during 1988, considering the discussed limitations, is 47.7 gigatons (Gt). Assuming maximum FBD values per biome a global total for foliar biomass was calculated to be 77.9 Gt. Accounting for the seasonal variation of FBD decreases the yearly global


Figure 10. January FBD Surface


Figure 11. July FBD Surface
esimate for foliar biomass, as expected. The next step is to compare these estimates to other global estimates of foliar biomass for validation. Comparison on a biome specific basis would be advantageous, but the problem of non-comparability of classification schemes would introduce additional uncertainties.

The one global total of foliar biomass found in the literature was 75 Gt (Box, 1981). Lieth and Whittaker (1975) and Whittaker (1975) provided terrestrial biome and global totals of leaf area and chlorophyll. Other sources contained relationships between foliar biomass and leaf area or chlorophyll (using their biome classification scheme). Foliar biomass/area values were determined from data on tropical rain forests (Grubb 1977, Edwards and Grubb 1977, Blasco and Tassy 1975, and Tanner 1977), and temperate deciduous forests, temperate evergreen forest, woodlands, and shrublands (Lieth and Whittaker 1975, Whittaker and Niering 1975, Whittaker and Woodwell 1969, Whittaker et al. 1974, Whittaker 1966, Whittaker 1962). The values ranged from 50 tons $/ \mathrm{km}^{2}$ leaf area to 178 tons $/ \mathrm{km}^{2}$ leaf area. A value of 100 tons $/ \mathrm{km}^{2}$ leaf area was taken as representative for forests, woodlands, and shrublands. These values were multiplied by the total leaf area estimates for each biome indicated in the literature to calculate total foliar biomass of 87.5 Gt for the globe (Table 7).

Another comparison involves converting chlorophyll mass to foliar biomass. Foliar biomass conversion factors were used only for forests, woodlands and shrublands. These values ranged from $127 \mathrm{~g} / \mathrm{g}$ (foliar biomass/chlorophyll mass) to 366 $\mathrm{g} / \mathrm{g}$ for temperate deciduous forests and woodlands. $250 \mathrm{~g} / \mathrm{g}$ was taken as representative for these biomes as well as the tropical rain and seasonal forests. A value of $453 \mathrm{~g} / \mathrm{g}$ (rounded to $450 \mathrm{~g} / \mathrm{g}$ ) was used for the temperate evergreen forest (sprucefir). These were used to calculate total foliar biomass for the forest biomes (Table 8).

Table 7. The total foliar biomass for the globe was calculated by converting LAI into foliar biomass and totalling the leaf biomass of each biome.

| Biome | Areal Extent <br> in $\mathrm{km}^{2} \times 10^{6}$ | LAI | LAI $\times$ Area <br> in $\mathrm{km}^{2} \times 10^{6}$ | Tons/km ${ }^{2}$ <br> Leaf Area | Foliar Biomass <br> Gigatons |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Tropical Rain Forest | 17.0 | 8 | 136 | 100 | 13.6 |
| Tropical Seasonal Forest | 7.5 | 5 | 38 | 100 | 3.8 |
| Temperate Evergreen Forest | 5.0 | 12 | 60 | 100 | 6.0 |
| Temperate Deciduous Forest | 7.0 | 5 | 35 | 100 | 3.5 |
| Boreal Forest | 12.0 | 12 | 144 | 100 | 14.4 |
| Woodland/Shrubland | 8.5 | 4 | 34 | 100 | 3.4 |
| Savanna | 15.0 | 4 | 60 | 100 | 6.0 |
| Temperate Grassland | 9.0 | 3.5 | 32 | 460 | 14.7 |
| Tundra and Alpine | 8.0 | 2 | 16 | 300 | 4.8 |
| Desert/Semi-desert | 18.0 | 1 | 18 | 100 | 1.8 |
| Extreme Desert | 24.0 | 0.05 | 1.2 | 100 | 0.1 |
| Cultivated Land |  |  |  |  |  |
| Swamp/Marsh | 14.0 | 4 | 56 | 250 | 14.0 |

Table 8. The total foliar biomass for the forest biomes was calculated by multiplying the total chlorophyll per biome by the foliar biomass/chlorophyll mass values.

| Biome | Chlorophyll <br> tons $\times 10^{6}$ | Foliar biomass/ <br> Chlor. mass | Foliar biomass <br> Gigatons |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Tropical Rain Forest | 51.0 | 250 | 12.8 |  |  |
| Tropical Seasonal Forest | 18.8 | 250 | 4.7 |  |  |
| Temperate Evergreen Forest | 17.5 | 450 | 7.9 |  |  |
| Temperate Deciduous Forest | 14.0 | 250 | 3.5 |  |  |
| Boreal Forest | 36.0 | 450 | 16.2 |  |  |
| Woodland/Shrubland | 13.6 | 250 | 3.4 |  |  |
| Total Foliar Biomass for the Forests |  |  |  |  | 48.5 Gt |

The total foliar biomass estimates for these six biomes using the chlorophyll conversion equalled 48.5 Gt , comparing favorably to the total of 34.2 Gt using the leaf area conversions for global total foliar biomass estimates considering the uncertainties associated with the calculations and the potential noncomparability between the biome classifications.

These two comparisons produced similar results. Each estimate using leaf area or chlorophyll conversion is higher than the estimates produced in this research. Nevertheless, considering the uncertainties associated with conversion estimates and that the estimates from both conversion calculations and the model estimates are of the same order of magnitude, the results are considered satisfactory.

In the final validation procedure, the grassland biome described in this study is compared to a regional analysis of the Senegalese Sahel grassland described by Tucker et al. (1983) and Tucker et al. (1986). The grassland biome presented in this study was isolated for this analysis. Tucker's model (equation 6) was run using the GVI data for the grassland biome and compared to the grassland model derived in this analysis (equation 7).

$$
\begin{align*}
& F B D=\exp \frac{G V I+1.0107}{0.1857}  \tag{6}\\
& F B D=0.2485 \exp \frac{G V I * 100}{14.3317} \tag{7}
\end{align*}
$$

GRASS raster (cell) format requires integers, therefore, the GVI data in equation 2 is multiplied by 100. (Consult Appendix C to see all the derived biome specific coefficients.)

It is not possible to compare both models within the geographic confines of Tucker's study area. Tucker describes his study area as grassland while the aggregated Olson classification used in this analysis describes this same geographic area as $15 \%$ Farms/Towns, $26 \%$ Nonpaddy Irrigated Dryland, and $59 \%$ Dry Forest and Woodlands.

However, it is possible to apply Tucker's model to the entire grassland biome described by Olson for comparison. The resulting estimated range of Tucker's
model as applied to the grassland biome is $.02-0.58 \mathrm{~kg} / \mathrm{m}^{2}$ while the range taken from the literature for this analysis was $.05-0.5 \mathrm{~kg} / \mathrm{m}^{2}$. Running Tucker's model at the same monthly time step produces a global grassland estimate of 0.528 Gt of FBD for 1988 , while this model estimates 0.947 Gt of FBD in 1988.

It appears, however, that this model overestimates grassland with the GVI data. One explanation is that the entire grassland biome which this model is based on, likely has different characteristics than the geographically restricted grasslands of the Senegalese Sahel. This would result in different empirical constants creating the different results. Another explanation is that the grassland that is described by Tucker is really what Olson would call Dry Woodland and Forest. Either of these explanations would account for differences in the results. Considering the limitations of these comparison, it is difficult to say whether the model developed in this research overesitimates or underestimates foliar biomass. Nevertheless, the results of this model were always of the same order of magnitude with the validation estimates, lending credence to its validity.

The results of the statistical analysis to isolate biomes of similar seasonal variations are presented. As previously discussed, the tropical biomes were determined by the majority of their areal extent with respect to their geographic locations. Nevertheless, the descriptive statistics of the monthly GVI data sets concerning the tropical biomes, as defined in this study, were also used to determine and justify which biomes should be considered tropical (Table 9).

Comparing the months of maximum mean GVI and the coefficient of variation it is clear that these biomes were similar before being combined into the tropical category. For example, the Tropical Montane biome has months of maximum mean and coeffecient of variation for the northern and southern hemisphere of, November, 0.283 , and November, 0.303 , respectively. When combined into the

Table 9 . This table displays some descriptive statistics in order to quantitatively compare the global categories of seasonal variation of GVI data within the tropical biomes.

| Monthly GVI Data Sets |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Descriptive Statistics of Tropical Biomes by Hemispheres |  |  |  |  |  |  |
| Biome | Global <br> Category | Month of Max. Mean | Coefficient of Variation | $\begin{gathered} 5 t h \\ \% \text { ile } \end{gathered}$ | $\begin{aligned} & 95 t h \\ & \% \text { ile } \end{aligned}$ | Median GVI |
| Trop. Montane | North | November | 0.283 | 15 | 44 | 36 |
| Trop. Montane | South | November | 0.303 | 13 | 46 | 32 |
| Dry Forest/Wdld | North | August | 0.324 | 7 | 37 | 26 |
| Dry Forest/Wdld | South | December | 0.318 | 12 | 44 | 32 |
| Trop. Seas. Frst | North | November | 0.183 | 24 | 44 | 36 |
| Trop. Seas. Frst | South | December | 0.198 | 22 | 45 | 36 |
| Trop/Subt. Frst | North | December | 0.224 | 19 | 43 | 35 |
| Trop/Subt. Frst | South | December | 0.197 | 22 | 44 | 37 |
| Descriptive Statistics of Tropical Biomes by Tropics |  |  |  |  |  |  |
| Biome | Global Category | Month of Max. Mean | Coefficient of Variation | $5 t h$ \%ile | 95th \%ile | Median |
| Trop. Montane | Tropic | November | 0.293 | 15 | 45 | 33 |
| Dry Forest/Wdld | Tropic | November | 0.375 | 10 | 42 | 26 |
| Trop. Seas. Frst | Tropic | November | 0.205 | 22 | 45 | 36 |
| Trop/Subt. Frst | Tropic | November | 0.209 | 20 | 44 | 36 |

tropic category the month of maximum mean and the coeffecient of variation are November and 0.293 , respectively. This similarity of all the tropical biomes is an indication that the arbitrary north-south division is not necessary, and perhaps wrong. The Dry Forest and Woodland biome displays the greatest variance because it is located in higher latitudes of the tropics.

## Sensitivity Analysis

A sensitivity analysis was performed to characterize the sensitivity of the model to the maximum and minimum estimates of FBD found in the literature. To assess the sensitivity of global estimates of foliar biomass, the biome specific FBD values were arbitrarily changed by $\pm 10 \%$ around the minima and maxima (Table 10). For each one of the four deviations about the maxima and minima, new coefficients were derived to run the model. Global totals of foliar biomass were
calculated to compare the magnitude of change associated with each of these deviations (Table 11).

Clearly, movement about the maxima has the greatest effect on overall quantities of estimated FBD. A $20 \%$ change in the maxima causes about 7 Gt difference to be estimated while the minima causes approximately 2 Gt of change. The $5.54 \%$ change of foliar biomass associated with $+10 \%$ maxima shown in table 11 would be greater if the maximum values of estimated FBD were not held constant to the maximum limit defined by the 95 th percentile GVI value on a per biome basis (Figure 9).

Table 10. The estimates of FBD per biome are shown with a $\pm 10 \%$ deviation from the minima and maxima. These percentages in FBD are used to characterize the potential of error in the estimates of FBD presented in this research.

| Ten Percent Change About the Minima and Maxima Foliar Biomass Estimates |  |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Biome | Minima <br> $\mathrm{kg} / \mathrm{m}^{2}$ | $-10 \%$ <br> $\mathrm{~kg} / \mathrm{m}^{2}$ | $+10 \%$ <br> $\mathrm{~kg} / \mathrm{m}^{2}$ | Maxima <br> $\mathrm{kg} / \mathrm{m}^{2}$ | $-10 \%$ <br> $\mathrm{~kg} / \mathrm{m}^{2}$ | $+10 \%$ <br> $\mathrm{~kg} / \mathrm{m}^{2}$ |  |
| Nonpolar Desert | .01 | .009 | .011 | .05 | .045 | .055 |  |
| Polar Desert | .01 | .009 | .011 | .10 | .09 | .11 |  |
| Tundra | .01 | .009 | .011 | .05 | .045 | .055 |  |
| South Temperate BLF | .2 | .18 | .22 | .8 | .72 | .88 |  |
| Grassland | .05 | .045 | .055 | .5 | .45 | .55 |  |
| Shrubland | .1 | .09 | .11 | .5 | .45 | .55 |  |
| Farms/Towns | .1 | .09 | .11 | .5 | .45 | .55 |  |
| Nonpaddy Irrigated Dryland | .1 | .09 | .11 | .5 | .45 | .55 |  |
| Forest/Fields/Woods | .1 | .09 | .11 | .8 | .72 | .88 |  |
| North Temperate BLF | .3 | .27 | .33 | .6 | .54 | .66 |  |
| Cool Conifer Hardwood | .3 | .27 | .33 | 1.0 | .9 | 1.1 |  |
| Tropical Montane | .3 | .27 | .33 | .7 | .63 | .77 |  |
| Wetlands/Hinterlands/Shore | .1 | .09 | .11 | .5 | .45 | .55 |  |
| Dry Forest and Woodlands | .1 | .09 | .11 | .5 | .45 | .55 |  |
| Semi-Arid Woodlands | .1 | .09 | .11 | .3 | .27 | .33 |  |
| Warm Conifer | .1 | .09 | .11 | 1.0 | .9 | 1.1 |  |
| Paddyland | .1 | .09 | .11 | .5 | .45 | .55 |  |
| Taiga | .3 | .27 | .33 | 1.5 | 1.35 | 1.65 |  |
| Trop seas Humid BLF | .3 | .27 | .33 | .8 | .72 | .88 |  |
| Trop/subt Humid BLF | .3 | .27 | .33 | 1.3 | 1.17 | 1.43 |  |
| Cool Conifer | .5 | .45 | .55 | 2.0 | 1.8 | 2.2 |  |

Table 11. This table displays a $10 \%$ movement around the maxima and minima FBD estimates found in the literature. The estimated global totals are shown each $10 \%$ change in maxima and minima as well as the percent change.

| Change in Global Quantities of Estimated FBD |  |  |
| :--- | :---: | :---: |
| \% Change of <br> Foliar Biomass | Foliar Biomass <br> in Gigatons | \% Change in <br> Total Foliar Biomass |
| Estimated Foliar Biomass | 47.7 | 0 |
| Maxima $+10 \%$ | 50.5 | 5.54 |
| Maxima $-10 \%$ | 43.7 | 8.39 |
| Minima $+10 \%$ | 48.8 | 2.86 |
| Minima $-10 \%$ | 46.4 | 2.72 |

## CHAPTER 5

## Conclusions

This study represents a first attempt to model the seasonal variation of FBD across the terrestrial biosphere by quantitatively using global satellite data and other global vegetation data. The approach has proven useful to capture the gross spatiotemporal patterns of the distribution of photosynthetically active FBD. The seasonal variation of GVI was shown to adequately estimate the seasonal variation of FBD. The model estimates of FBD were sufficient when compared to other estimates of global totals of FBD considering uncertainties involved with both estimates.

The GVI data does, however, have these limitations as presented by Singh (1988a), Singh (1988b), and Holben (1986). It is suspected that the addition of snow in the winter months also has a strong effect on the NDVI ratio by decreasing the GVI values for conifer forests where such a drastic decrease is not expected (Appendix D).

The division of the Earth into three unique zones of seasonal variation was shown to be acceptable. The Northern and Southern Hemispheres were obviously different, but the Tropics were shown to present their own seasonal variation. The tropical biomes showed little seasonal variation. This can be seen by looking at the annual time course for the tropical biomes where GVI data and estimated FBD remained relatively stable. (see appendix G and H for monthly distributions of GVI and FBD, respectively.) The abitrary equatorial division through the tropical biomes was statistically shown to be unnecessary and perhaps wrong. Therefore, the equatorial region should not be divided at the equator and, further, the region between $23^{\circ} \mathrm{N}$ and $23^{\circ} \mathrm{S}$ should isolated for analysis of seasonal variation. Also, the Northern and Southern Hemispheres should start at $23^{\circ} \mathrm{N}$ and $23^{\circ} \mathrm{S}$,
respectively, and extend to the poles for analysis of seasonal variation. This provides three global bands described by latitude that represent three unique zones of seasonal variation.

The biggest limitation in this approach is the model itself because the model depends upon scarce data. not the quantitative use of the imagery, but instead the data on spatial heterogeneity for FBD and on the global vegetation maps. After scanning the literature for spatial FBD data only the sources documented in this manuscript were consulted. Other sources were inappropriate or not available, indicating an extremely low availibility of this kind of vegetation data. Because there is such a scarcity of global vegetation data, this model was calibrated with these data, and as a result a high amount of faith was placed in their validity. With the upcoming EOS platforms researchers may better relationships between biophysical variables and the remotely sensed data. But unless there is future research undertaken to establish ranges of FBD, LAI, and other variables and uncertainty associated with these ranges, quantitative estimates of biophysical variables may not drastically improve.

A sensitivity analysis was performed on the FBD parameters taken from the literature in light of the high amount of faith placed in the accuracy of these FBD ranges. The potential error was characterized in percentages and can be used as a guide to assess the effect that error may have upon the global totals of FBD that were estimated by this model.

In the interest of global climate change and tropospheric chemistry, the FBD surfaces were used as variables in a biogenic gas emissions model. The results of this research used as an input for an isoprene model yielded favorable results (Turner et al., submitted). The results are also being input into a water balance model and a carbon flux model (Marks, submitted, and Gucinski, submitted).

There is a need to refine the relationships between the satellite data and FBD, as well as other biophysical variables. The new EOS platforms will provide much improved data over present systems that will not only provide better relationships for biophysical variables but will demand that the scientific community clearly understand concepts associated with all aspects of remote sensing.

To go beyond empirical relationships based on satellite vegetation indices to physcial relationships based on radiative properties of vegetation is a highly desirable research priority that would enhance the quantitative application of satellite data. Ongoing remote sensing expirements such as FIFE, HAPEX-MOBLY, and BOREAS may help to establish a more physiologically based model. The more physiologically based a model becomes the less dependent the model becomes on vegetation maps and other vegetation data. This is highly desirable for global and regional remote sensing studies of vegetation where, as previously discussed, there is a scarcity of physiological data to couple with the remote sensing data, because one may not have to couple physiological data with remote sensing data.

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APPENDICES

## APPENDIX A

Monthly Images of GVI, January through December













## APPENDIX B

Derivation of FBD-GVI relationship
Original equation expressed generically:

$$
\begin{equation*}
Y=a \ln \frac{X}{b} \tag{1}
\end{equation*}
$$

where:

$$
\begin{aligned}
& \mathrm{Y}=\mathrm{NDVI} \\
& \mathrm{X}=\mathrm{FBD}
\end{aligned}
$$

$$
\mathrm{a}, \mathrm{~b}=\text { biome specific constants }
$$

Equation 1 must now be manipulated to bring $X(F B D)$ to the outside. The following equations show the steps of this process.

Divide both sides by $a$.

$$
\begin{equation*}
\frac{Y}{a}=\ln \frac{X}{b} \tag{2}
\end{equation*}
$$

Exponentiate the equation.

$$
\begin{equation*}
\exp \frac{Y}{a}=\frac{X}{b} \tag{3}
\end{equation*}
$$

Multiply both sides by $b$.

$$
\begin{equation*}
X=a \exp \frac{Y}{b} \tag{4}
\end{equation*}
$$

Variable Substitution and further algebraic manipulation.
Set up the equations in order to simultaneously solve for $a$ and $b$.
(the equations are manipulated with the X and Y variables for ease)

$$
\begin{equation*}
X=a \exp \frac{Y}{b} \tag{5}
\end{equation*}
$$

remove the exponent

$$
\begin{equation*}
\ln X=\ln \left\{a \exp \frac{Y}{b}\right\} \tag{6}
\end{equation*}
$$

final equation

$$
\begin{equation*}
\ln X=\ln a+\frac{Y}{b} \tag{7}
\end{equation*}
$$

Substitute in the known model variables.

$$
\begin{align*}
& \ln F \max =\ln a+\frac{N \max }{b}  \tag{8}\\
& \ln F \min =\ln a+\frac{N \min }{b} \tag{9}
\end{align*}
$$

where:
Fmin $=$ FBD minimum value per biome
Fmax $=$ FBD maximum value per biome
$\mathrm{Nmin}=$ GVI minimum value per biome
Nmax $=$ GVI maximum value per biome
$\mathrm{a}, \mathrm{b}=$ biome specific constants
Solve for for $b$.

$$
\begin{align*}
& \ln F \max =\ln a+\frac{N \max }{b}  \tag{10}\\
& \ln F \min =\ln a+\frac{N \min }{b} \tag{11}
\end{align*}
$$

Subtract equation 11 from equation 10

$$
\begin{gather*}
\ln \left\{\frac{F \max }{F \min }\right\}=\frac{N \max -N \min }{b}  \tag{12}\\
b=\frac{N \max -N \min }{\ln F \max / F \min }
\end{gather*}
$$

Solve for $a$ in terms of $b$. (leave $b$ as the variable " $b$ ").

$$
\begin{gather*}
\ln F m a x=\ln a \frac{N m a x}{b}  \tag{14}\\
\ln a=\ln F m a x-\ln b \tag{15}
\end{gather*}
$$

expontiate

$$
\begin{equation*}
a=F \max \frac{-N \max }{b} \tag{16}
\end{equation*}
$$

The components of the model are now ready to be solved with the data presented in the methods section of this manuscript.

## APPENDIX C

1) Table of biome specific variables and constants

| Biome | Fmax | Fmin | Nmax | Nmin | B | A |
| :--- | :---: | :--- | :---: | :---: | :--- | :---: |
| NonpolarDesert | 0.05 | 0.01 | 33 | 5 | 0.0077137 | 19.2614 |
| PolarDesert | 0.10 | 0.01 | 17 | 4 | 0.0031623 | 3.47438 |
| Tundra | 0.05 | 0.01 | 37 | 6 | 0.0073951 | 19.8827 |
| SouthernTemperateBLF | 0.80 | 0.20 | 45 | 8 | 0.14938 | 27.4112 |
| Grassland | 0.5 | 0.05 | 42 | 9 | 0.024885 | 14.3317 |
| Shrubland | 0.5 | 0.10 | 39 | 4 | 0.081248 | 19.2614 |
| Farms/Towns | 0.5 | 0.1 | 42 | 8 | 0.064472 | 20.5041 |
| NonpaddyIrrigatedDryland | 0.5 | 0.1 | 37 | 4 | 0.082277 | 20.5041 |
| Forest/Fields/Woods | 0.8 | 0.1 | 45 | 10 | 0.05202 | 16.8314 |
| NorthernTemperateBLF | 0.6 | 0.3 | 50 | 8 | 0.18899 | 43.2809 |
| CoolConiferHardwood | 1.0 | 0.3 | 46 | 7 | 0.20549 | 29.0704 |
| TropicalMontane | 0.7 | 0.3 | 45 | 15 | 0.1964 | 35.4067 |
| Wetlands/Hinterlands/Shore | 0.5 | 0.1 | 40 | 7 | 0.072478 | 21.7467 |
| DryForestandWoodland | 0.5 | 0.1 | 42 | 10 | 0.060474 | 19.8827 |
| Semi-aridWoodland | 0.3 | 0.1 | 33 | 6 | 0.073060 | 25.4867 |
| WarmConiferForest | 1.0 | 0.1 | 41 | 12 | 0.030303 | 11.726 |
| Paddyland | 0.5 | 0.1 | 38 | 10 | 0.056282 | 17.3974 |
| Taiga | 1.5 | 0.3 | 40 | 20 | 0.060 | 12.4267 |
| TropicalSeas.HumidBLF | 0.8 | 0.3 | 45 | 22 | 0.1174 | 23.4495 |
| Trop/Subt.HumidFrst | 1.3 | 0.3 | 44 | 20 | 0.088397 | 16.3673 |
| CoolConiferForest | 2.0 | 0.5 | 45 | 17 | 0.21549 | 20.1977 |

## APPENDIX D

1) Table of Olson categories by number used in the aggregation.

| Biome | Olson Categories |
| :--- | :--- |
| NonpolarDesert | 49526971 |
| PolarDesert | 5051 |
| Tundra | 42535463 |
| SouthernTemperateBLF | 26 |
| Grassland | 4064 |
| Shrubland | 41 |
| Farms/Towns | 3031 |
| NonpaddyIrrigatedDryland | 373839 |
| Forest/Fields/Woods | 55565758 |
| NorthernTemperateBLF | 25 |
| CoolConiferHardwood | 2324 |
| TropicalMontane | 28 |
| Wetlands/Hinterlands/Shore | 44456566676872 |
| DryForestandWoodland | 3243 |
| Semi-aridWoodland | 46474859 |
| WarmConiferForest | 27 |
| Paddyland | 36 |
| Taiga | 2021606162 |
| TropicalSeas.HumidBLF | 29 |
| Trop/Subt.HumidFrst | 3373 |
| CoolConiferForest | 22 |

## APPENDIX E

Temporal Profiles for Each Biome












> GV and Olson aggregation data All GV values of 0 deleted
> Area - weighted mean GU Vs Month HEM $=\mathrm{S}$ BOME $=(1 \in)$ Semi - arid Woodinnd


## APPENDIX F

Calibration Curves for Each Biome








## FBD vs GVI

Using the 5th and 95th Percentile BIOME $=$ (18) Paddyland $\mathrm{CAT}=\mathrm{N}$






## APPENDIX G

Monthly FBD Surfaces, January through December













## APPENDIX H

Monthly Histograms of GVI per Biome




$$
\frac{L}{L} \frac{L}{L} L
$$






|  |  |  |  |
| :---: | :---: | :---: | :---: |
|  |  |  |  |
|  |  |  |  |
















## APPENDIX I

Monthly Histograms of FBD per Biome

|  |  |  |  |
| :---: | :---: | :---: | :---: |
|  |  |  |  |
|  |  |  |  |




















| , |  | mid | BIOME $=$ (21) Trop/subt. Humid Frist MONTH |
| :---: | :---: | :---: | :---: |
|  |  |  |  |
|  |  |  |  |
|  |  |  |  |




[^0]:    * The water class was excluded from the percentage calculations.

