

1 **Title:** The Plant Phenology Monitoring Design for the National Ecological Observatory
2 Network

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1 **Abstract**

2 Phenology is an integrative science that comprises the study of recurring
3 biological activities or events. In an era of rapidly changing climate, the relationship
4 between the timing of those events and environmental cues such as temperature,
5 snowmelt, water availability or day length are of particular interest. This article provides
6 an overview of the plant phenology sampling which will be conducted by the National
7 Ecological Observatory Network NEON, the resulting data, and the rationale behind the
8 design. Trained technicians will conduct regular *in situ* observations of plant phenology
9 at all terrestrial NEON sites for the 30-year life of the observatory. Standardized and
10 coordinated data across the network of sites can be used to quantify the direction and
11 magnitude of the relationships between phenology and environmental forcings, as well as
12 the degree to which these relationships vary among sites, among species, among
13 phenophases, and through time. Vegetation at NEON sites will also be monitored with
14 tower-based cameras, satellite remote sensing and annual high-resolution airborne remote
15 sensing. Ground-based measurements can be used to calibrate and improve satellite-
16 derived phenometrics. NEON's phenology monitoring design is complementary to
17 existing phenology research efforts and citizen science initiatives throughout the country
18 and will produce interoperable data. By collocating plant phenology observations with a
19 suite of additional meteorological, biophysical and ecological measurements (e.g.,
20 climate, carbon flux, plant productivity, population dynamics of consumers) at 60
21 terrestrial sites, the NEON design will enable continental-scale inference about the status,
22 trends, causes and ecological consequences of phenological change.

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1 **Key words:** long-term monitoring; NEON; plant phenology; open-source data; sample
2 design

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1 **Introduction**

2 The overarching mission of NEON is to enable understanding and forecasting of
3 the impacts of climate change, land use change, and the introduction of invasive species
4 on ecosystem structure and function (see Thorpe et al., this issue). Tracking the timing of
5 seasonally recurring life cycle events (phenology) is thus a natural focal area of study for
6 the Observatory. Plant phenological transitions may be triggered by a variety of cues,
7 including chilling, spring temperature, growing degree days, and daylight cues (Chuine
8 2000); many of these factors are likely to shift significantly over the next 30 years (IPCC
9 2013). Changes in phenology have been observed for many taxa across the earth
10 (Parmesan and Yohe 2003). The onset of spring phenological events advanced at an
11 estimated mean rate of 1.2 days per decade from 1955-2002, across the Northern
12 Hemisphere, likely caused by recent climate warming (Schwartz et al. 2006).
13 Observational and experimental studies indicate that plants flower on average ~5 days
14 earlier per 1°C increase in spring temperature (Wolkovich *et al.* 2012) and current
15 projections indicate that spring phenology could advance by between 1 and 10 days over
16 the planned 30-year lifespan of the NEON observatory (IPCC 2013). Many species,
17 however, delay flowering in response to increases in winter or spring temperatures
18 (Mazer et al. 2013), and there is still much to learn about the causes of variation among
19 species and higher taxa in the direction and magnitude of their phenological responses to
20 both temperature and rainfall (Mazer et al., 2013, 2015).

21 Beyond providing an indicator of climate change, the timing of phenological
22 transitions is also a potentially important driver of demographic trajectories and
23 biogeographic distributions of individual taxa, and of ecological processes including
24 species interactions and rates of biogeochemical cycling (Morissette et al. 2008).

1 Phenological traits may physiologically constrain broad-scale distribution patterns of
2 species; phenology is consistently an important predictor in process-based species
3 distributions models (Chuine 2010 and references therein). Phenological plasticity may
4 be a beneficial trait; for example, species whose activity patterns closely track interannual
5 climate variability tend to have improved growth, productivity, or reproductive success
6 than those that do not (Cleland et al. 2012). In other cases, however, early greenup or
7 floral bud development in response to anomalously early arrival of spring can be
8 detrimental. Phenological advancement in response to warm spring temperatures
9 followed by a late frost can have catastrophic effects on fruit and seed production and
10 canopy development (Inouye 2008, Hufkens et al. 2012).

11 Climate-induced changes in phenology can create feedbacks that alter
12 biogeochemical cycling and species interactions (Melillo et al., 2014). Changes in the
13 timing of leaf budburst and senescence affect surface radiation, near surface temperature,
14 hydrology and carbon cycling (Churkina *et al.* 2005, Bonan 2008, Richardson et al. 2010,
15 Jeong et al. 2012, 2013). An analysis of more than a dozen models included in the North
16 American Carbon Program (NACP) Interim Synthesis indicated across all models, sites,
17 and years of data, for each forest type; errors of up to 25 days in predictions of “spring
18 onset” were common, and errors of up to 50 days were observed (Richardson *et al.* 2012).
19 From the general positive relationship between carbon uptake and season length derived
20 from a synthesis of a range of eddy covariance sites, the largest phenological errors in
21 current models would translate into between ~150 and ~450 g m⁻² of carbon annually
22 (Churkina et al. 2005). Differential responses to phenological cues between plants,
23 consumers, and/or pollinators can disrupt the overlap in activity periods among

1 interacting organisms, potentially resulting in changes in species fecundity and cascading
2 effects on the food chain (Strode 2003, McKinney et al. 2012) or local extinction of
3 consumer populations (Singer and Parmesan 2010).

4 Plant phenology has been studied at a range of geographic and temporal scales
5 and by employing a variety of tools, including: recording *in situ* observations,
6 experimental manipulation of abiotic factors, modeling, remote sensing, and digital
7 photography (Cleland et al. 2007). Understanding and reconciling the information
8 contributed at each scale is challenging (Morissette et al. 2008) and observations at
9 multiple scales are rare (but see Liang et al. 2011). This article provides an overview of
10 the plant phenology sampling that will occur within NEON sites, including observation
11 protocols, the spatial and temporal frequency of monitoring, and the taxa targeted for
12 observations, and the rationale for the sampling regime that was selected (Box 1). The
13 science design, developed by a technical working group of comprised of phenology
14 experts from academic institutions, government and non-profit agencies, reflects current
15 best practices in monitoring terrestrial plant phenology. By providing integrated and
16 multi-scale suites of measurements on the seasonal progression of a diversity of taxa and
17 ecosystem processes at intensively measured sites, data collected by NEON will enable
18 the scientific community to develop mechanistic linkages between the environmental
19 drivers that affect plant phenology, as well as the functional consequences of changing
20 phenology for a range of ecosystem types and processes. The resulting scientific
21 knowledge can inform decision-making processes related to natural resource
22 conservation and management, control of invasive species and infectious disease, and
23 efforts related to societal climate change adaptation (Enquist et al. 2014).

1 ***Box 1:NEON's contribution***

2 NEON is poised to advance the field of phenology by:

3 1) Accumulating high quality, long-term, standardized measurements
4 recorded by trained technicians across 20 major ecosystem types found within the
5 US;

6 2) Observing replicate individuals of select species to quantify
7 intraspecific variation in the timing of phenological events within and across
8 years, facilitating precise population-level estimates of phenology;

9 3) Observing multiple species to characterize the range of phenological
10 response patterns across species and functional groups and life history strategies;

11 4) Collocating plant phenological measurements with other terrestrial and
12 atmospheric measurements data, which may be used to understand relationships
13 between climate, phenology, ecosystem processes and biodiversity; and

14 5) Providing open-access, standardized datasets that easily integrate with
15 other large scale monitoring networks.

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18 ***Measurements***

19 Plant phenology is typically quantified by observing the date of onset and the
20 duration of particular phenophases, which may include both vegetative and reproductive
21 events. Specific phenophase definitions have not been universally adopted across
22 monitoring networks. Without common units, data interoperability becomes a limiting
23 factor in data integration. Consistent with NEON's commitment to use existing

1 nationally-accepted, vetted and standardized protocols wherever possible, NEON will
2 employ USA-NPN phenophase definitions and protocols (Denny *et al.* 2014).

3 Advantages of USA-NPN protocols and the reasons for selecting this standard for
4 NEON *in situ* phenology observations include: (1) status-based monitoring, or the
5 practice of reporting the phenological condition of an individual at any time that it is
6 monitored; (2) repeated tracking of marked and georeferenced replicate individual
7 perennials and patches of annual/clonal herbs and, (3) incorporation of both status and
8 ‘intensity’ definitions for phenophases (Kao *et al.* 2012, Denny *et al.* 2014). Using
9 status-based rather than first-event monitoring is a departure from many historical
10 phenological monitoring protocols, but has the advantage that events (such as leaf
11 emergence in Mediterranean climates, or flowering in many desert species) that may
12 occur multiple times during a single year can be captured. Status-monitoring also allows
13 the explicit quantification of uncertainties in phenophase transition dates (which occur in
14 continuous time) that are introduced by monitoring in discrete temporal bouts, as well
15 quantifying the duration of phenophases rather than just their date of onset. Monitoring
16 marked individuals/small patches ensures that the recorded dates of phenological events,
17 or their duration, are decoupled from population size (Miller-Rushing *et al.* 2008). The
18 protocols employed include intensity metrics (e.g. percentage of the canopy that is full
19 with leaves) along with phenophase status (e.g. one or more live, unfolded leaves
20 visible). These data can be used to estimate mean population onset and end dates for each
21 phenophase, as well as track the seasonal progression of development throughout the
22 active period. Together, these data should provide better linkages to ecosystem function
23 and remotely sensed phenological data than existing ‘first event’ phenological datasets,

1 which typically quantify the phenological status of only the most extreme individuals
2 within a population of unknown size (Miller-Rushing *et al.* 2008). While other
3 phenophase definitions exist (e.g. the BBCH scale, commonly used in agricultural
4 systems, as well as across Europe (Meier 2001; Koch *et al.* 2007)), the USA-NPN scales
5 were selected for interoperability with large-scale distributed monitoring datasets in the
6 continental US. Mapping from USA-NPN definitions to BBCH definitions is feasible for
7 many phenophases.

8 The phenology protocol includes repeated assessment of phenophase status and
9 intensity on each individual (see section Temporal distribution of sampling, below, for
10 more details), as well as an annual assessment of individual-level covariates that can
11 affect phenology. Due to resource constraints, only a subset of the USA-NPN-defined
12 phenophases (as described by Denny *et al.* 2014) will be targeted in NEON phenology
13 sampling protocols, with the greatest focus on leaf phenology. The focus on canopy
14 development was selected based on recommendations in the NSF Research Coordination
15 Network Report (2012), to facilitate linkages with NEON's measurements of ecosystem
16 processes such as landscape phenology and carbon cycling. To connect phenological
17 measurements to plant health, productivity and canopy position, NEON will measure the
18 size (stem diameter, % cover, height and canopy dimensions), disease status, health
19 condition and structure of each individual plant or patch once per year. These annual
20 measurements will be consistent with those taken on other plants at NEON sites as part of
21 the vegetation structure and productivity protocol (see Meier and Jones 2015 for details).

22 ***Phased sampling design***

1 Two priorities were identified for NEON's plant phenology observations:
2 *Phenology of dominants*, which includes estimating the mean and intraspecific variance
3 of phenological timing in dominant species within each site (see Phase I, below), and
4 *Community phenology*, focused on capturing a range of species-specific phenologies that
5 represent the plant community at each NEON site (Phase II). Dominants are targeted
6 specifically to facilitate linkages to ecosystem function based on the assumption that
7 species contribute to ecosystem properties roughly in proportion to their relative
8 abundances (Grime 1998). Sampling of dominant species' phenology will enable linking
9 phenological events and patterns observed above-ground to processes captured at other
10 scales by other NEON measurement systems (including root phenology, ecosystem
11 productivity and respiration, and carbon, water and nutrient cycling) and to the ground-
12 based land-surface phenology signal observed via remote sensing methods. It will also
13 provide critical information on intraspecific variation in phenology patterns, which are
14 poorly captured when monitoring efforts are limited to a census of one to several
15 individuals per site. Sampling of community-level phenology will inform questions
16 regarding interspecific variation in the timing and duration of phenological phases and
17 their sensitivity to climate. The resulting dataset will enable assessment of the degree to
18 which phenological timing and climate sensitivity vary based on functional groups or
19 growth forms (e.g. natives/exotics, overstory/understory, perennial/annual,
20 deciduous/evergreen, herbaceous/woody, early and late-season). These patterns can
21 enable generalizations regarding the likely phenological responses and sensitivities of
22 species beyond those targeted for regular observation.

1 NEON will implement phenological monitoring in two phases in order to
2 accomplish both inter- and intra-specific sampling goals. During Phase I (Phenology of
3 dominants), implemented during the first three full (i.e., all sites operational) years of
4 sampling, phenological observations will concentrate on intensive monitoring of three
5 dominant species at each of the 60 terrestrial sites.. The NSF Research Coordination
6 Network (RCN) report (2012) recommends a minimum of 5-10 replicate individuals
7 sampled for vegetative phenology per site per species, with an ideal sampling intensity of
8 20-30 individuals. In the absence of existing data sufficient to statistically determine
9 smaller minimum sample sizes for particular species and sites, NEON will target the
10 higher end of this range in order to quantify intraspecific variation in phenological timing
11 for the three most dominant species at each site (see section ‘Temporal distribution of
12 sampling, below, for details of monitoring frequency).

13 Phase II (community phenology), will follow Phase I and consist of more limited
14 sampling than Phase I in terms of frequency and the number of replicate individuals per
15 species (minimum of 5 individuals per species per site), but will have an increased
16 number of species. The focal shift will alter which individuals are monitored, but keep
17 the total number of plants monitored per site at ~90-100 due to budgetary limitations.
18 Phase II monitoring will commence in the 4th year of operational sampling and will
19 continue for the remainder of NEON operations at each site. Species to be monitored in
20 Phase II will include dominant species (the three species studied as part of Phase I at each
21 site) and up to 17 additional species per site that collectively represent a range of
22 functional groups and life history strategies. Phase II will inform both the range of

1 phenological patterns occurring at a site, as well as predictive models of the sensitivities
2 of particular species based on their traits (Buckley and Kingsolver 2012).

3 *Spatial distribution of sampling*

4 A common critique of much of the existing ground-phenology observation data is
5 that observations are limited in space and are reported as points, whereas remote sensing
6 data pixels from commonly used satellite products used to model phenology range from
7 30m to >1km (Schwartz and Hanes 2010). While some studies have found little spatial
8 autocorrelation in a single plant species' phenological response given uniform
9 temperature over small areas (Schwartz et al. 2013), dispersion of monitored individuals
10 throughout a larger area is important to encompass variation in plant phenology within
11 the sampling area caused by microenvironmental variation, genetic variation, or both. To
12 facilitate repeatable observation of multiple individuals over a relatively large area, while
13 keeping travel time to a minimum, marked individuals will be situated along a fixed, 800-
14 meter square 'loop' transect (200 meters on a side), with the 4 edges oriented in the four
15 cardinal directions. This size is comparable to the ~250m modis pixel size, which is
16 commonly used in satellite-based phenology assessments.

17 This loop will be situated within or near NEON's flux tower footprint whenever
18 possible. The distance of the transect from the tower will be site specific based on
19 identified exclusion areas around tower instrumentation, and will be placed to facilitate
20 inclusion of individuals located within sampling plots used for NEON's biomass and
21 productivity measurement (see Meier et al. this issue) (Figure 1). Collocation of the
22 phenology transect with the instrument tower will allow meteorological and biophysical
23 data collected by tower-mounted sensors to be used directly in analysis of phenological

1 data (e.g. how local climate affects phenology) and vice versa (e.g. how leaf status affects
2 daily carbon flux). NEON's tower locations are positioned such that the tower air-shed is
3 situated in a spatially and structurally homogenous area with the goal of a minimum of
4 80% contribution from the representative ecosystem, ensuring that plants selected for
5 phenological monitoring are also located within a regionally representative habitat type.
6 The assumption is that the intraspecific variation in phenological responses will, in
7 general, be from individuals subject to similar environmental conditions. Even so,
8 microtopographic features may still affect variation in observed phenological response.
9 Additional information such as slope, aspect, community composition, above-ground
10 biomass, and canopy chemistry as derived from NEONs airborne observation system may
11 provide additional insight into the realized environmental heterogeneity of the various
12 sites.

13 ***Temporal distribution of sampling***

14 A standard sampling frequency for phenology has not been prescribed by the
15 ecological community. Typically, sampling frequency varies by species, environment,
16 sampling objectives, and budgetary and logistical constraints. The ideal frequency of
17 sampling depends on analysis goals (e.g. fitting a thermal forcing model vs. long-term
18 trend detection vs. quantifying intraspecific variation in phenology), as well as the degree
19 of intraspecific and interannual variation in phenology. Mazer *et al.* (2015) found that
20 twice-weekly sampling over a three-year period was sufficient to detect statistically
21 significant associations between winter monthly rainfall and/or mean temperature (and
22 their interactions) and the onset dates of vegetative growth, flowering, and fruiting in four
23 species monitored in California across broad environmental conditions. An NSF

1 Research Coordination Network (RCN) report on phenology (2012) suggests a sampling
2 interval of 2-4 times per week. Miller-Rushing *et al.* (2008) recommend sampling every
3 2nd day to ensure a 97% chance of detecting a significant change in flowering date over
4 10 years of sampling, based on existing long-term flowering data collected in
5 Massachusetts and Colorado. These recommendations assumed realistic anticipated rates
6 of climate warming and interannual variability in temperature, in addition to a sensitivity
7 of flowering date to temperature of 1 day/°C. A more recent synthesis of long-term
8 phenology datasets worldwide (Wolkovich *et al.* 2012), however, suggests that flowering
9 phenology will, on average, shift at a rate of 5-6 days/°C. Therefore less frequent
10 sampling may be adequate for many species for simple trend detection.

11 Following the RCN recommendations, the first three years of sampling the
12 phenological status of dominant species (Phase I) will be observed 3 times a week during
13 key transition periods (i.e. leaf emergence and senescence, Table 1). Resulting data will
14 be used to inform the sampling intensity necessary to characterize the mean (+/- 3 days
15 S.E.) for leaf phenology transition dates for the 3 dominant species at the site in
16 subsequent years. This target is based on a recent analysis by Jeong *et al.* (2012), who
17 concluded that when observational error in estimating population mean transition days
18 for key phenological events (e.g. budburst) is greater than +/- 3 days, parameterizing
19 phenological forcing models is compromised. During Phase II, the frequency of
20 phenological observations will be reduced to 2 times a week during transitional phases in
21 order to accommodate sampling of a greater number of species.

22 Phenologically active periods will vary among species both spatially across the
23 continent, and inter-annually at each site. In order to catch the full growing season for all

1 selected species, NEON will aim to commence weekly sampling three weeks prior to the
2 earliest anticipated onset of the first phenophase (based on the earliest date observed in
3 recent records for the species). This date will be determined using local information,
4 where available (such as at LTER sites where historical phenological data exist, or
5 indicator plants at a nearby, lower elevation site), or from historical MODIS data, in sites
6 where local information is not available to guide sampling. Start of season metrics based
7 on remote sensing data are typically biased towards early dates (White *et al.* 2009;
8 Ganguly *et al.* 2010), so this should provide an ‘earliest’ outer bound on start of season.

9 Once bud break or initial growth is observed, the observation frequency will
10 increase from once a week to either three times (Phase I) or two times (Phase II) a week.
11 The intensive sampling stage ends once full-sized leaves have emerged/full canopy has
12 formed, and sampling frequency is reduced to once a week or once every other week to
13 survey for open flowers. Three weeks before the anticipated first date of senescence,
14 based on local and/or MODIS data, sampling frequency will increase again to weekly (if
15 previously reduced to every other week). At the first sign of leaf senescence (i.e. fall
16 color), observation frequency will, once more, increase to 2 times a week sampling until
17 <5% of leaves remain or until three consecutive censuses of no change have been
18 observed.

19

20 ***Species selection***

21 Prior to commencing phenology observations at a given site, NEON will conduct
22 quantitative vegetation surveys within 20-30 randomly placed plots within the tower
23 footprint to assess species abundance. Three dominant species will be identified at each

1 site for Phase I phenology monitoring. The dominant species selected will include the
2 two most abundant canopy species plus the single most abundant understory species for
3 sites with greater than 50% canopy closure, and the two most abundant understory
4 species plus the single most dominant overstory species for sites with less than 50%
5 canopy closure. At sites with no defined woody overstory, e.g. grasslands, all three
6 species will be selected from the herbaceous community. Understory and canopy species
7 frequently occupy discrete temporal niches, with the understory species, or in some cases
8 understory individuals, showing advanced phenology relative to that of canopy-forming
9 individual (Richardson and O'Keefe 2009).

10 Additional species to be sampled for Phase II will be selected from the whole
11 community of species present within the tower footprint using a random selection
12 procedure, weighted by abundance. Abundance of woody vegetation with stem diameter
13 >1 cm at a height of 130cm along the stem will be determined by biomass, calculated
14 from stem diameters, according to Jenkins (2003) allometric equations per species.
15 Because biomass is more difficult to assess for shrubs and herbaceous species, abundance
16 in these growth forms is assessed based on total areal cover by species (surveyed as
17 percent cover / m² for herbaceous species and measurement of canopy area within
18 defined survey plots for shrubs). Species are then re-grouped into a single list, ordered by
19 their absolute abundance rank as estimated within the 20-30 plots surveyed. The
20 abundance values will then be used to identify species for targeted selection (Phase I) or
21 to weight species for random selection (Phase II). By stratifying in this way, common
22 species with very low biomass have a greater likelihood of selection than infrequent high
23 biomass individuals.

1 Exceptions to the randomized selection process will be made to intentionally
2 target species that either contribute to NEON’s ability to address grand challenge
3 questions (e.g. invasive species) or contribute to NEON’s ability to align data collection
4 with existing national citizen science data collection efforts. Invasive species, USA-NPN
5 campaign taxa and PBB ‘10 most wanted’ species will be preferentially selected from the
6 species list prior to weighted random selection. In order to avoid species that are not
7 present in sufficient quantities to maintain monitoring of replicate individuals, NEON
8 will limit potential community members for monitoring to those species found in more
9 than 10% of the surveyed plots. The weighted random selection procedure should ensure
10 that a diversity of plant growth forms, invasives and natives are selected at sites where
11 they are present, without requiring any *a priori* definition of ‘functional group’, a concept
12 which is not yet well understood for predicting phenology. It will also serve to
13 concentrate monitoring efforts on species that are relatively common, while also
14 including some rare species.

15

16 ***Site-specific modifications***

17 Modifications will be made for sites with growing seasons or species with life
18 histories that differ from the typical temperate deciduous model. For example, sampling
19 may begin earlier than described above to capture flowering phenophases for plants that
20 flower prior to leaf production. Additionally, sampling frequency will need to be
21 modified at sites without a clear seasonal greening pattern (e.g. tropical ecosystems, or
22 Mediterranean climates where species may leaf out or flower multiple times per year in
23 response to episodic rainfall); in these cases, year-round sampling with longer intercensus

1 intervals will be necessary to capture phenological trends. Modifications will also need to
2 be made for cropped (agricultural) sites. At these sites, NEON will monitor the cultivated
3 species; in most cases, the selected species will vary by year to track crop rotations and
4 will likely not have the diversity to support Phase II sampling. Details of monitoring,
5 including frequency and replication, may be adjusted based on the initial data collected at
6 each site and budgetary constraints. All site specific details including site-specific
7 modifications, species selection and targeted sampling windows will be captured,
8 tracked, and made available to end users as part of the NEON phenology sampling
9 protocol (available through the NEON web portal; www.neoninc.org).

10

11 **Applications of phenology data**

12 NEON plant phenology data will provide foundational information about the
13 variability in plant phenology across populations, communities, and landscapes, which
14 can be used to validate remotely-sensed land surface phenology measures and better
15 inform terrestrial biosphere models. To date, realistic parameterization of phenological
16 models for wild species is limited to the very few species for which relevant data are
17 available (Jeong et al. 2012). NEON will expand the taxonomic representation of
18 phenological data, measuring as many as 20 plant species at each of 60 sites across the
19 continent. Quantifying the range of phenological responses across a diversity of species
20 and sites also will aid in the development of more general phenological forcing models
21 based on species and site characteristics, as well as understanding of the degree to which
22 these models can be used to estimate phenology where direct measurements are not
23 available. Bayesian hierarchical models are a promising avenue forward in community

1 phenology forecasting (see Ibáñez et al. 2010, Diez et al. 2012 for examples applied to
2 individual sites with multiple taxa, or single taxa measured across multiple sites). Multi-
3 site, multi-species datasets provided by NEON can form the basis of an expanded
4 phenological modeling framework across sites and species. Accurate representation of
5 intra- and inter-annual variability in vegetation phenology is critical for correctly
6 predicting net CO₂ uptake (Desai 2010). An evaluation of vegetation phenology in 14
7 terrestrial biosphere models found that for deciduous forests an early start of season bias
8 of two weeks or more was typical across all models which resulted in a 13% over
9 estimate of gross ecosystem productivity (Richardson et al. 2012). Such
10 misrepresentation of phenology has consequences beyond ecosystem productivity
11 estimates. When terrestrial and atmospheric models are not properly coupled, reductions
12 in temperature associated with the onset of leaf emergence and associated increases in
13 transpiration are often misrepresented (Levis and Bonan 2004). This insufficient coupling
14 during critical phenological stages can lead to errors in modeled microclimate and
15 weather patterns and thus present cascading effects on other model components. High
16 quality, long-term, standardized phenological measurements across major ecosystem
17 types will be critical components for improving model development and accuracy.

18 The dominant species in all plant communities generally represent key resources
19 for animals that depend on them for food or shelter. Consequently, phenological shifts in
20 the onset, duration, and abundance of vegetative and reproductive resources detected by
21 NEON's phenological monitoring program can alert resource managers of changes that
22 may affect the community composition and population dynamics persistence of insects,
23 pollinators, birds, and mammals at site or regional scales. This goal requires monitoring

1 of the animals that interact with the focal plant species at NEON sites. In addition to
2 plant phenology observations (the focus of this manuscript), terrestrial protocols that
3 contribute to phenological monitoring at NEON sites include trapping of (1) mosquitoes
4 and (2) small mammals throughout the active growing season; these data may be used to
5 track phenology of mosquito emergence and annual population dynamics and small
6 mammal reproductive periods, respectively (Hoekman *et al.*, this issue, Thibault *et al.*,
7 this issue). Integration of NEON phenology data with surveillance data on other taxa,
8 conducted either by NEON or by PIs working at NEON sites, can help track phenological
9 asynchrony between interacting species and potential consequences to shifts in
10 overlapping activity periods throughout the duration of the observatory.

11 The development of integrated, interoperable datasets will enhance the utility of
12 data collected by NEON and other programs. A number of other programs (e.g. USA
13 National Phenology Network (<https://www.usanpn.org/>), Long Term Ecological
14 Research (LTER) Network sites (<http://www.lternet.edu/>), National Parks
15 (<http://science.nature.nps.gov/im/monitor/>), the Pan European Phenology Project
16 (PEP725; <http://www.pep725.eu/>)), as well as multiple longterm PI-directed research
17 projects also take phenology measurements. NEON data will augment and compliment
18 these efforts, providing replication and longevity of measurements that are difficult to
19 achieve without a centralized source of funding. Because of NEON's planned
20 infrastructure, its potential to link ground-based measurements, landscape green-up and
21 brown-down metrics, and ecosystem processes is unique (Keller et al. 2008). NEON will
22 also collect biweekly leaf area index (LAI) digital hemispherical photos, landscape
23 images collected multiple times per day using stationary cameras (phenocams), and

1 carbon flux estimates processed at half-hour intervals. These data streams, augmented
2 with annual sub-meter hyperspectral and LiDAR remote sensing data will be valuable in
3 determining statistical and mechanistic associations between aboveground, belowground
4 and landscape scale seasonal dynamics.

5 One limitation of the NEON design for phenology is that the financial and
6 logistical commitment required to measure phenology alongside a large suite of other
7 parameters (see Lunch et al. 2014 for the full list of NEON data products) constrains the
8 total number of NEON sites. As a result, NEON sites are spatially sparse compared to
9 continent-wide citizen-science observation efforts, such as the USA National Phenology
10 Network (www.usanpn.org; hereafter USA-NPN), Project BudBurst (www.budburst.org;
11 hereafter PBB) and affiliated national and regional monitoring networks. Because
12 NEON uses nationally standardized protocols, however, data from the intensively studied
13 NEON sites can be readily combined with existing and ongoing efforts to facilitate
14 continental-scale analysis and forecasting. By integrating ground-based observations with
15 other North American plant phenological monitoring programs (e.g., USA-NPN),
16 existing datasets (e.g. Wolkovich et al. 2012), the PhenoCam network
17 (<http://phenocam.sr.unh.edu/webcam/>), satellite imagery (e.g. MODIS land cover
18 dynamics <http://modis.gsfc.nasa.gov/data/dataproduct/>), and/or models (e.g. the Growing
19 Season Index; Jolly et al. 2005), *in situ* phenology observations made by NEON can
20 contribute critical information to an annual ‘green wave’ (Schwartz 1998; Ault et al *in*
21 *press*) projection over the continent.

22 Phenological data can also be used in a number of natural resource management
23 activities (Enquist et al. 2014). Accurate phenological forecasts can aid land managers in

1 timing controlled burns, mechanical harvesting, pesticide and/or herbicide applications
2 for maximum efficiency in controlling invasive species. Data on seasonal growth and
3 senescence patterns can inform wildfire predictions. Similarly, information on peak
4 flowering and leaf color change dates can help promote and plan for seasonal tourism
5 coincident with wildflower or fall foliage viewing. Last, recent studies theorize that a
6 species' ability to make appropriate phenological adjustments to a changing climate may
7 be predictive of its future success in a changing climate (Willis et al. 2010; Pau et al.
8 2011). This suggests that an improved understanding of species-specific phenological
9 sensitivities could be used to identify particularly vulnerable native taxa for protection, or
10 prioritize invasive species for removal.

11 Changes in plant phenology are widely regarded as 'fingerprints of climate
12 change' or 'climate change indicators'(e.g., U.S. Environmental Protection Agency
13 2014); indeed, plant phenology is an exemplary essential species trait in the ongoing
14 development of Essential Biodiversity Variables (EBV's) targeted for international
15 monitoring (Pereira et al. 2013). Ongoing efforts both nationally (e.g. USA-NPN, Project
16 Budburst) and internationally (e.g. PEP725), will continue to document patterns of plant
17 phenology over large spatial extents. Leveraging data from NEON will enable the
18 extrapolation not only of patterns of plant phenological shifts across the continent (e.g.
19 Jeong et al. 2013, Ault et al. *in press*), but potentially also of the functional consequences
20 of these shifts. Collocated measurements conducted by NEON will elucidate the degree
21 to which plant phenological status is broadly indicative of related ecosystem processes
22 for which continent-wide data are sparse (e.g. below-ground phenology, carbon flux,
23 seasonal biomass accumulation. In turn, the analysis, synthesis, and application of

1 phenological information will facilitate decision-making related to critical ecological
2 issues that affect societal well-being now and into the future.

3

4

5

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10

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1 Table 1. Proposed rule sets for specific growth forms for phenology sampling at sites with a well-defined growing season

Growth form	Monitor indicator individual for:	Sample 3x/week until all tagged individuals show:	Sample 1x/week until all tagged individuals show:	Then ² :	Then:	Sample 2x/week until all individuals show:	Sample 1x/week until:	Then:
Cactus	Breaking flower buds	NA	End sampling season when no more fresh flowers are present	NA	NA	NA	NA	NA
Deciduous broadleaf	Breaking leaf or flower buds	>50% of canopy is full with leaves or three consecutive bouts of no change	95% or more of canopy is full with leaves	Commence every-other week monitoring for open flowers	Monitor indicator individuals for one or more colored leaves	One or more colored leaves	<5% of canopy full with green or colored leaves	End sampling season
Deciduous conifer	Breaking needle buds	>50% of canopy is full with needles or three consecutive bouts of no change	95% or more of canopy is full with needles	Commence every-other week monitoring for open pollen cones	Monitor indicator individuals for one or more colored needles	One or more colored needles	<5% of canopy full with green or colored needles	End sampling season
Drought deciduous broadleaf	Breaking leaf buds	Young leaves	No more young leaves	Commence every-other week monitoring for open flowers	Monitor indicator individuals for one or more colored leaves ³	One or more colored leaves	<5% of canopy full with green or colored leaves	End sampling season
Evergreen Broadleaf	Breaking leaf buds	Young leaves	No more young leaves	Commence every-other week monitoring for open flowers	End sampling season when no more fresh flowers are present	NA	NA	NA
Evergreen conifer	Breaking needle buds	Young needles	No more young needles	Commence every-other week monitoring for open pollen cones	End sampling season when no more fresh pollen cones are present	NA	NA	NA

Growth form	Monitor indicator individual for:	Sample 3x/week until all tagged individuals show:	Then ² :	Then:	Sample 2x/week until all individuals show:	Sample 1x/week until:	Then:
Evergreen forb	Breaking leaf buds	Young leaves	Commence every-other week monitoring for open flowers	End sampling season when no more fresh flowers are present	NA	NA	NA
Forb	Initial growth	One or more fully unfolded leaves	Commence every-other week monitoring for flowering phenology	Monitor indicator individuals evidence of senescence	NA	No more full sized leaves are present	End sampling season
Graminoid	Initial growth	>50% of plant is green or three consecutive bouts of no change	Commence every-other week monitoring for flowering phenology	Monitor indicator individuals for >5% Leaf senescence (i.e. percentage of plant that is green <95%)	<95% green leaves	<5% of plant is green	End sampling season
Pine	Emerging needles or pollen cone development	Young needles	Commence every-other week monitoring for open cone	End sampling season when no more fresh pollen cones visible	NA	NA	NA
Semi-evergreen broadleaf⁴	Breaking leaf or flower buds	Young leaves OR >50% of canopy is full with leaves OR three consecutive bouts of no change	Commence every-other week monitoring for open flowers	Monitor indicator individuals for one or more colored leaves ³	One or more colored leaves	<5% of canopy full with green or colored leaves	End sampling season

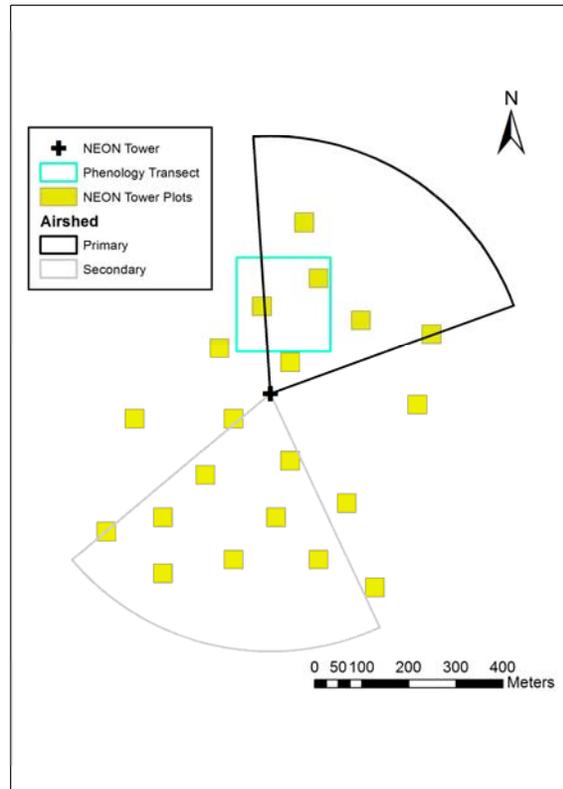
1 This is generally applicable to temperate or boreal systems; sites lacking a distinct growing season where growth occurs year-round or is episodic such that a growing season cannot be defined will be monitored on a weekly basis.

2 If flowering phenology precedes leaf/needle bud break skip the steps outlined in this column and decrease monitoring to watching indicator individuals for fall senescence or end monitoring for the season as specified in the following column.

3 Seasonal monitoring may end at this point if senescence does not occur.

4 Semi-evergreen broadleaf growthform may be used for species in which life history varies with latitude. Monitoring strategy should be driven by phenophase observations.

1 Figures



2

3 **Figure 1.** Layout of phenology transect (teal square) with respect to the NEON
4 Tower (cross shape), the airshed (wedge shapes) and the Tower Plant Productivity plots
5 (yellow squares) (figure credit: Rachel Krauss, 2015)

6

7