



Title: Vegetation photosynthetic phenology metrics in northern terrestrial ecosystems: a dataset derived from a gross primary productivity product based on solar-induced chlorophyll fluorescence

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1 **Abstract**

2 Vegetation phenology can profoundly modulate the climate-biosphere interactions and
3 thus plays a key role in regulating the terrestrial carbon cycle and the climate. However,
4 most previous phenology studies are based on the traditional vegetation indices, which
5 are inadequate to characterize the seasonal activity of photosynthesis. Here, we
6 generated an annual vegetation photosynthetic phenology dataset with a spatial
7 resolution of 0.05 degree from 2001 to 2020, using the latest gross primary productivity
8 product based on solar-induced chlorophyll fluorescence (GOSIF-GPP). We combined
9 smoothing splines with multiple change-point detection to retrieve the phenology
10 metrics: start of the growing season (SOS), end of the growing season (EOS), and
11 length of growing season (LOS) for terrestrial ecosystems in the Northern Hemisphere.
12 We found that the derived phenology metrics agreed better with in situ observations



13 from the flux tower sites than vegetation indices and MODIS-GPP. Our phenology
14 metrics captured the spatial-temporal patterns of the single and double growing season
15 in the Northern Hemisphere. The double season was mainly from the cropland rotation
16 and ecosystems having two different phenological cycles. In addition, we observed a
17 trend toward advanced SOS in about 62.98% of the land area, with a mean rate of
18 0.14 ± 0.01 days year⁻¹, a trend toward delayed EOS in about 61.87% of the area, with a
19 mean rate of 0.19 ± 0.16 days year⁻¹, and a trend toward extended LOS in about 70.52%
20 of the area, with a mean rate of 0.33 ± 0.17 days year⁻¹. Our phenology product can be
21 used for validating and developing phenology models or carbon cycle models, for
22 evaluating satellite remote sensing phenology, and for monitoring climate change
23 impacts on terrestrial ecosystems. The data are available
24 at <https://doi.org/10.6084/m9.figshare.17195009.v2> (Fang et al. 2021).

25

26 **1. Introduction**

27 Vegetation phenology, the cycle sequence of plant vital activities, is a highly sensitive
28 indicator of the climate impacts on terrestrial ecosystems (Richardson et al. 2013, Piao



29 et al. 2019, Wang et al. 2019, Keenan et al. 2020). Most phenology studies focus on the
30 structural changes of plants, such as using the growth process of leaf represented by the
31 greenness indicators (Seyednasrollah et al. 2021, Yang and Noormets 2021). However,
32 recent studies found that the methods based on vegetation greenness have limited ability
33 to capture the photosynthesis changes in some vegetation types (e.g. evergreen forests)
34 since the greenness and photosynthesis are sometimes decoupled (Walther et al. 2016,
35 Smith et al. 2018). The inaccurate estimation of phenology can lead to substantial
36 uncertainties in the estimation of plant productivity and carbon sequestration
37 (Richardson et al. 2012, Wu et al. 2017, Fang et al. 2020).

38 The plant photosynthetic cycle on the seasonal time scale is termed as ‘vegetation
39 photosynthetic phenology’, which represents the functional aspects of plant activities
40 (Gu et al. 2009). This phenology definition is based on the photosynthesis transition
41 dates extracted from the gross primary productivity (GPP) time series. Thus, the
42 accuracy of extracted phenology metrics is largely dependent on the data source of GPP.
43 Currently, the GPP can either be derived from Eddy Covariance (EC) flux towers at the
44 ecosystem scale or from satellite remote sensing or modeling at the regional or global



45 scale (Xiao et al. 2019). The EC technique, which is considered as the most accurate
46 observation method (Baldocchi et al. 2001), has provided long-term GPP estimates for
47 more than 20 years. However, these observations are limited by their spatial distribution
48 and some key areas are still underrepresented (Xiao et al. 2019). For example, only a
49 few EC sites provide public datasets in the tropical and high latitude regions. GPP
50 derived from satellite remote sensing is able to investigate large-scale phenology across
51 the globe (Sjöström et al. 2013). Greenness-related vegetation indices such as the
52 normalized difference vegetation index (NDVI) and the enhanced vegetation index
53 (EVI) have been widely used to estimate GPP (Wu et al. 2017, Huang et al. 2019, Dai
54 et al. 2021). However, these indices work better for capturing the variations in
55 chlorophyll content or vegetation coverage and are not sufficient to track the
56 instantaneous physiological changes in vegetation photosynthesis, especially for
57 evergreen vegetation (Joiner et al. 2014, Li and Xiao 2020). Recently, the emergence
58 of satellite-based solar-induced chlorophyll fluorescence (SIF) has offered
59 unprecedented opportunities for developing more accurate photosynthetic phenology
60 data products on large scales (Joiner et al. 2011, Frankenberg et al. 2014, Li et al. 2018,



61 Köhler et al. 2018). SIF, a signal emitted by plant chlorophyll molecules after absorbing
62 photosynthetically active radiation (APAR), is considered to be an effective tool for
63 diagnosing terrestrial photosynthesis and estimating GPP more accurately (Meroni et
64 al. 2009, Verma et al. 2017, Wood et al. 2017, Li and Xiao 2020). Based on the SIF
65 product, recent studies used the relationship between the GPP and SIF to estimate the
66 regional or global GPP (SIF-GPP) (Li and Xiao 2019, Zhang et al. 2020). Previous
67 studies reported that SIF-GPP can better capture the GPP dynamics in evergreen
68 vegetation and dryland ecosystems than traditional vegetation indices (Bertani et al.
69 2017, Smith et al. 2018).

70 In addition, the retrieval of phenology in previous studies mainly used a logistic
71 regression model to fit the time series of smoothed vegetation indices or GPP, and the
72 predetermined thresholds or inflection points are identified as the transition dates of
73 phenology in the fitted curve (Garrity et al. 2011, Wang et al. 2017, Yang and Noormets
74 2021). However, this method needs to reconstruct the original data sequence and thus
75 results in uncertainty from the model parameterization (Klosterman et al. 2014).
76 Furthermore, this method is usually used to capture a single growing season instead of



77 the multiple growing seasons in a given year (Yang and Noormets 2021).
78 Correspondingly, Richardson et al. (2018) proposed a method of smoothing spline and
79 multiple change-point detection to retrieve the transition dates of phenology from the
80 camera data. The strength of this method is not limited by the uncertainty of additional
81 model parameters and can also be applied in ecosystems having multiple growing
82 seasons. The method has been successfully used at multiple sites in North America
83 (Richardson et al. 2018) and needs to be extended to large scales.

84 Here, we aim to generate a photosynthetic phenology metrics dataset based on the
85 GPP product derived from satellite SIF data. Our data can detect multiple growing
86 seasons, which can be used to evaluate the photosynthesis activity of vegetation from
87 large scales. The metrics include the start state-transition dates of photosynthesis (SOS),
88 the end state-transition dates of photosynthesis (EOS), and the duration length of
89 photosynthesis (LOS). With this goal, we constructed a method combining smoothing
90 filter and change-point detection to retrieve photosynthetic phenology from a recently
91 developed SIF-based GPP product (GOSIF-GPP: 2001-2020) with a fine spatial
92 resolution (0.05°). This method enables us to acquire multiple photosynthesis activity



93 periods of vegetation within one year. The remainder of this paper describes the data of
94 SIF-GPP and land cover data, the adopted method for retrieving photosynthetic
95 phenology metrics, the results and discussion of the metrics and their uncertainties, and
96 the conclusions.

97

98

99 **2. Data**

100 We used the GOSIF-GPP dataset from 2001-2020 (Li and Xiao 2019) to derive the
101 phenology metrics on large scales in this study (<http://data.globalecology.unh.edu/>).
102 GOSIF-GPP was estimated from the GOSIF dataset based on eight linear SIF-GPP
103 relationships with 0.05° spatial and 8-day temporal resolutions (i.e., 46 GPP estimates
104 per year for each 0.05° grid cell). The GOSIF dataset was developed by using discrete
105 SIF soundings from the Orbiting Carbon Observatory-2 (OCO-2), remote sensing data
106 from MODIS, and reanalysis data from MERRA-2 based on machine learning method
107 (Li and Xiao 2019b). The GOSIF-GPP showed reasonable seasonal and spatial patterns
108 and was highly correlated with GPP from FLUXNET (Li and Xiao 2019). Here, we



109 identified the vegetation type of each grid cell according to the MODIS Land Cover
 110 Type Product Version 6 (MCD12C1: <https://lpdaac.usgs.gov/products/mcd12q1v006/>)
 111 (**Fig. S1**, 0.05° spatial resolution). The current study used six broad vegetation types
 112 (i.e. **forests**: evergreen needleleaf forests, evergreen broadleaf forests, deciduous
 113 needleleaf forests, deciduous broadleaf forests, and mixed forests; **shrublands**: closed
 114 canopy shrublands and open shrublands; **savannas**: savannas and woody savannas;
 115 **grasslands**; **wetlands**; **croplands**) in the Northern Hemisphere. For the sake of
 116 reducing noise generated by non-vegetation signals, we excluded the area covered with
 117 bare soil and sparse vegetation (i.e., maximum GPP lower than 2.0 g C m⁻² day⁻¹) (Liu
 118 et al. 2016). Since the seasonal variation of vegetation photosynthesis in the tropical
 119 region is relatively small (Piao et al. 2019), we focused on the area above 30° N latitude.
 120 The final dataset is provided at each 0.05° grid for 20 years in the six terrestrial
 121 ecosystems of the Northern Hemisphere.

122 To evaluate phenology estimates based on GOSIF-GPP, we used the daily GPP
 123 data from EC flux towers across the Northern Hemisphere based on the
 124 FLUXNET2015 Dataset (<https://fluxnet.org/data/fluxnet2015-dataset/>) (Pastorello et al.



2020). We retained the EC flux sites that were relatively homogeneous because the footprint of 0.05° GOSIF product and EC tower may not exactly match (Li and Xiao 2019). We selected the flux sites having available GPP data for more than one year. The selected flux tower GPP dataset includes 49 sites with 389 site-year data (the detailed information of these flux sites can be found in **Table S1**). As a comparison, we also compared the performance of GOSIF-GPP based phenology metrics with those based on the vegetation indices and GPP products from the MODIS datasets. For each site, we extracted and calculated three vegetation indices from the Nadir Bidirectional Reflectance Distribution Function (BRDF)-Adjusted Reflectance dataset MCD43A4 (produced daily and 500 m resolution) including the NDVI, the EVI, the near-infrared reflectance of vegetation (NIR_v) (Badgley et al. 2017), and the 8-day, 500-m MODIS GPP data (MOD17A2) (Zhao et al. 2005) from 2001 to 2014.

3. Method

3.1 Photosynthetic phenology metrics



141 The phenology metrics in this study include SOS, EOS, and LOS. Unlike the traditional
142 phenological events from the structural changes of leaf or flower, the photosynthetic
143 phenology is defined as the start (i.e. SOS) and end (i.e. EOS) state-transition dates of
144 the photosynthesis cycles. These transition dates are used as the phenology metrics.
145 One full cycle generally has five distinctive stages, including (1) photosynthesis
146 dormancy period, a season before the growing season; (2) photosynthesis development
147 period, a GPP rising stage; (3) photosynthesis peak period, a peak stage of GPP; (4)
148 photosynthesis recession period, a GPP falling stage; and (5) photosynthesis dormancy
149 period, the photosynthetically inactive stage after the growing season. Most previous
150 studies used the sigmoid-based methods (e.g., double-logistic model) to extract the
151 phenology, but these methods are limited to the single cycle (Yang and Noormets 2021).
152 Because some regions or ecosystems had multiple cycles in one year, we used the
153 smoothing splines and change points to identify the transition dates of photosynthesis.
154 In this study, all transition dates were extracted from the daily GPP sequence of each
155 grid cell. Thus, we interpolated the 8-day GOSIF-GPP data to the daily scale using
156 cubic spline interpolation before the extraction.



We constructed an automatic method to retrieve transition dates (i.e. SOS and EOS) of photosynthetic phenology using GPP data. The algorithm of this method is outlined in the flowchart in **Fig. 1**. The important basis for acquiring phenological events was the data reconstruction using smoothing methods to minimize the impact of abnormal values (Li et al. 2019). We applied the iterative procedure to conduct the smoothing process (**Fig. 1**): (1) Smoothing the GPP time series by the Savitzky-Golay filter, which can reflect the change characteristics of the original data sequence; (2) Calculating the ratio of the daily GPP value to the smooth value; (3) Identifying outliers in these ratios by using the Grubbs test; (4) Using the smooth value instead of the daily GPP value when the ratios were larger than one standard deviation below the mean ratio; (5) Applying the iterative procedure up to 20 times or until no outliers were detected from one iteration to the next. This procedure can largely keep the raw seasonal pattern of photosynthesis and avoid the uncertainty of parameter estimation by reconstructing the data time series by estimating parameters in the double logistic model.

The potential change points in the final smoothing splines were identified with the Pruned Exact Linear Time (PELT) method. This method can accurately detect the



173 significant change points in the data time series and does not need to preset the number
174 of change points. The PELT was first applied by Killick et al. (2012), and they described
175 in detail on how to find the change points in time series. For each photosynthesis cycle,
176 we followed Richardson et al. (2018) to set the penalty factor and the minimum segment
177 length of PELT as 0.5 and 14-days, respectively. We calculated the mean GPP value of
178 the adjacent change points as the potential peak and bottom baseline in one full cycle.
179 According to the time series of mean GPP value, we used the difference method to
180 detect the bottoms and peaks (i.e., minimum and maximum value in each cycle). The
181 adjacent bottoms and one peak were formed as a full cycle, and the value of these points
182 was considered as the baselines. Some GOSIF-GPP data affected by the weak
183 vegetation SIF signals could have unreliable cycles, and these cycles that had peaks less
184 than 0.25 of the maximum peak were excluded in the current study.

185 Here, the SOS and EOS dates of each cycle were determined by amplitude
186 thresholds. The amplitude was equal to the peak minus the bottom. Although the “true”
187 onset of photosynthesis may correspond most closely to the 10% amplitude threshold
188 (Wu et al. 2017), the most tightly-constrained transition dates tended to occur in the



189 later dates of the GPP rising stage and the earlier dates of the GPP falling stage
 190 (Richardson et al. 2018). Thus, we followed Richardson et al. (2018) to provide the
 191 SOS and EOS dates by using three amplitude thresholds: 10%, 25%, and 50%. The
 192 SOS and EOS were determined when the GPP smoothing splines reached the value of
 193 amplitude thresholds, and the LOS was defined as EOS minus SOS:

$$SOS_i = t, \text{ if } GPP_s(t) = (Peak - Bottom_1) \times threshold_i \quad (1)$$

$$EOS_i = t, \text{ if } GPP_s(t) = (Peak - Bottom_2) \times threshold_i \quad (2)$$

$$LOS_i = EOS_i - SOS_i \quad (3)$$

194 where i is the threshold (10%, 25%, and 50%); t is the day of the year (DOY); GPP_s is
 195 the daily value of the smoothing splines; $Bottom_1$ is the baseline for dormancy season
 196 before the growing season; $Bottom_2$ is the baseline for dormancy season after growing
 197 season. Note that we retrieved the phenology of vegetation indices (i.e. daily data) and
 198 MODIS-GPP (i.e. interpolating the 8-day data to the daily scale) by using the same
 199 method.

200

201 3.2 Uncertainty estimation



The uncertainties in the estimates of phenology metrics mainly arise from the gridded SIF-based GPP estimates, such as using the limited explanatory variables to acquire the gridded SIF estimates (i.e., GOSIF) and the relationship between the SIF and GPP. In this study, we did not assess the quality of the underlying SIF and GPP data, which was previously evaluated (Li and Xiao 2019); instead we used the Monte Carlo Bootstrapping method (Efron 1992) to estimate the related uncertainties. Bootstrapping provides valuable information about uncertainties without making assumptions about the underlying data distributions (Elmore et al. 2012). For each year of the individual grid cell, we used bootstrapping to replace the transition dates with 100 times random uniform sampling (Yang and Noormets 2021). The 5th and 95th percentiles of the 100 bootstrapped data were considered as the confidence interval of the mean estimated from the original transition dates.

4. Results and discussion

4.1 Comparison with phenology derived from vegetation indices, MODIS-GPP,



218 **and EC tower data**

219 We used the photosynthetic phenology metrics extracted from the daily GPP of the flux
 220 towers to examine the corresponding metrics extracted from the GOSIF-GPP product.
 221 We also use the same method to retrieve phenology from the NDVI, EVI, NIR_v, and
 222 MODIS GPP for the EC tower sites. According to the different thresholds, the metrics
 223 were divided in to nine groups (SOS_{10%}, SOS_{25%}, SOS_{50%}: SOS with 10%, 25%, and
 224 50% amplitude threshold; EOS_{10%}, EOS_{25%}, EOS_{50%}: EOS with 10%, 25%, and 50%
 225 amplitude threshold; EOS_{10%}, EOS_{25%}, EOS_{50%}: EOS with 10%, 25%, and 50%
 226 amplitude threshold) (**Fig. 2** and **Table 1**). Overall, the phenology metrics of GOSIF-
 227 GPP showed the highest correlations with the phenology metrics of EC tower GPP,
 228 while the phenology of NDVI showed the lowest correlations. For each metric, (1) SOS,
 229 the correlation coefficient (R) between the 10%, 25%, and 50% SOS of EC tower GPP
 230 and other data were (i.e. from high to low): GOSIF-GPP (0.78-0.80), MODIS-GPP
 231 (0.65-0.67), NIR_v (0.47-0.60), EVI (0.40-0.57), and NDVI (0.14-0.39); the root mean
 232 square error ($RMSE$) were (i.e. from low to high): GOSIF-GPP (14.99-18.03 days),
 233 MODIS-GPP (18.83-22.98 days), NIR_v (21.74-29.86 days), EVI (24.97-36.97 days),



234 and NDVI (26.20-26.91 days). (2) EOS, the highest R between 10%, 25%, and 50%
235 EOS of EC tower GPP and other data was GOSIF-GPP (0.63-0.73) and the lowest R
236 was NDVI (0.42-0.56). (3) LOS, the highest R between 10%, 25%, and 50% EOS of
237 EC tower GPP and other data was GOSIF-GPP (0.65-0.76) and the lowest R was NDVI
238 (0.28-0.40). The comparisons indicated that GOSIF-GPP showed consistently better
239 performance than the vegetation indices (i.e., NDVI, EVI, and NIR_v) for different
240 phenology metrics and different thresholds. MODIS-GPP had larger deviations
241 compared to GOSIF-GPP, which highlights the need for the improvement on light use
242 efficient models. NIR_v, the product of near-infrared reflectance and NDVI (Badgley et
243 al. 2017), was slightly better to capture the phenology metrics of tower GPP than EVI
244 and NDVI. The results agreed with previous studies which showed a stronger ability of
245 SIF in responding to the environmental conditions such as water and heat stresses, and
246 thus in better capturing the seasonal and interannual photosynthetic activity (Walther
247 et al. 2016; Smith et al. 2018; Li et al. 2018).

248 The derived phenology of GOSIF-GPP and EC tower GPP showed a close
249 correspondence across the 389 site-years. The best performance of the different



thresholds in SOS, EOS, and LOS was 25% ($R=0.80$, 0.73 , and 0.76 ; $RMSE=15.83$,
21.89, and 29.14 days, respectively), and the threshold of 10% had relatively low
performance in SOS and EOS ($R=0.79$ and 0.63 ; $RMSE=18.03$ and 23.55 days,
respectively) and 50% had relatively low performance in LOS ($R=0.65$; $RMSE=27.89$
days). Our results showed that our method better captured the SOS than the EOS, which
was consistent with previous studies that uncertainty occurred in satellite-based EOS
estimations, especially for the evergreen vegetation such as tropical and boreal
evergreen forests (Liu et al. 2016, Piao et al. 2019). In addition, more tower sites need
to be considered in further studies so that the photosynthesis phenology metrics from
the SIF product can be better evaluated.

4.2 Number of growing seasons

We used the method to retrieve the multiple growing seasons in the Northern
Hemisphere. **Fig. 3** showed the spatial distribution of the number of growing seasons.
Most regions in the Northern Hemisphere had a single growing season, while part of
the cropland had a double growing season in a given year. The North China Plain (the



red part in the top right of **Fig. 3**) had the most areas with the double growing season because the wheat-maize rotation was the most important cropping system in this plain (Zhao et al. 2006). This artificial crop rotation pattern brought two photosynthesis cycles: wheat grows in winter and spring, and maize grows in summer and autumn. In addition to croplands, a small proportion of the grid cells also had double growing seasons, such as some areas in California. Turner et al. (2020) reported that the double growing season in California was due to two overlapping ecosystems in one grid, whereas they were out of phase with each other: grasslands showed a peak of the growing season in April and forests peak in June. The phenology retrieval of such mixed ecosystems is still challenging and requires further exploration (Piao et al. 2019).

4.3 Spatial patterns of photosynthetic phenology metrics

Here, we showed the spatial distribution of the first growing season in **Fig. 4**. Overall, phenology metrics (SOS, EOS, and LOS) in terrestrial ecosystems of the Northern Hemisphere exhibited a spatially explicit pattern from the high latitudes to the low latitudes. Limited by low temperature, the areas around the Arctic Circle had the latest



282 SOS (DOY>140), the earliest EOS (DOY<220), and the shortest LOS (days<120). LOS
 283 gradually increased as the climate conditions became more suitable for photosynthesis
 284 and then reached the longest in the subtropics. For different thresholds, the mean
 285 difference of SOS, EOS, and LOS between 10% and 50% was 30 days, 40 days, and
 286 70 days, respectively. For different ecosystems (**Table 2**), grasslands showed the
 287 earliest SOS and EOS among all biomes; forests and savannas had the latest SOS;
 288 croplands and forests exhibited the latest EOS and the longest LOS, while shrublands
 289 and grasslands had the shortest LOS.

290 **Fig. 5** showed the spatial distribution of the second growing season. We found that
 291 the second 10% SOS in North China Plain was in the end of May (DOY=150) and the
 292 second 10% EOS was in the middle of September (DOY=280). This was consistent
 293 with the emergence and dormancy of maize (i.e. the second growing season). The wheat
 294 would seed after the maize was harvested and the greenness of wheat was in the early
 295 March of the next year, which was the start time of the first growing season (Tang et al.
 296 2020). In California, the second 10% SOS of some areas was in the early June
 297 (DOY=160) and 10% EOS was in the middle of September (DOY=280); the second



298 10% SOS of other areas was in the late September (DOY=250) and 10% EOS was in
299 the late November (DOY=330). These results were from the two different mixed grids,
300 one included the evergreen forests and grasslands, another included the croplands and
301 grasslands. Turner et al. (2020) found that the growth of grasses provides the first
302 growing season for these grids. As the grids included evergreen forests entered summer,
303 the increase of the available water in the soil resulted in the growth of evergreen woody
304 plants, prompting these grids to enter the second growing season. Other ecosystems
305 were gradually entered the dormant stage in fall, but the crops still maintained
306 photosynthesis, making the grids containing croplands show the second growing season.
307

308 **4.4 Uncertainties of photosynthetic phenology metrics**

309 The uncertainty used in this study was defined as the 5th and 95th percentiles of the
310 100 Monte Carlo bootstrapping samples ranging from a few days to several weeks
311 (**Table 2**). The uncertainty was the lowest for SOS and the highest for LOS; EOS had
312 intermediate uncertainty. The highest uncertainty in LOS maybe because of the
313 compounding effect of SOS and EOS (Yang and Noormets 2021). Generally, metrics



314 of grasslands had the lowest uncertainty: SOS uncertainty ranged from 3.8 to 5.1 days,
315 EOS uncertainty ranged from 8.6 to 10.4 days, LOS uncertainty ranged from 13.7 to
316 14.2 days; forests have the largest uncertainty, with SOS uncertainty ranging from 7.3
317 to 9.4 days, EOS uncertainty from 16.0 to 18.4 days, and LOS uncertainty from 25.4 to
318 25.7 days. The high uncertainty in the forests was possibly because this ecosystem
319 included multiple mixed vegetation types and the phenology of these plants was more
320 difficult to retrieve (Piao et al. 2019).

321

322 **4.5 Changes in photosynthetic phenology metrics**

323 We conducted the linear regression analysis by using the transition dates of phenology
324 and the time series in each grid cell, and the regression coefficient was considered as
325 the changing trend of the grid cell (**Fig. 6**). Here, we only showed the changes of the
326 dominant single growing season. For the spatial distribution of the phenology metrics
327 with the three thresholds, 61.71-64.25% of the study area experienced advanced trends
328 of SOS, with a large advanced trend in northwestern North America, northern Siberia,
329 and eastern Europe (changes > 0.6 days year⁻¹); 57.97%-65.89% of the study area



330 experienced delayed trends of EOS, with a large delayed trend in the northern North
331 America and northern Siberia (changes >0.4 days year⁻¹); 70.29-70.76% of the study
332 area experienced extended trends of LOS, with a large extended trend in northern China,
333 northern North America, and northern Siberia (changes >0.6 days year⁻¹). Note that the
334 inconsistent climate change trends in different seasons may lead to advanced or delayed
335 SOS and EOS in some regions simultaneously, such as eastern Europe (Cohen et al.
336 2012).

337 We spatially averaged the phenology metrics for the terrestrial ecosystems across
338 the Northern Hemisphere to assess the interannual variation of phenology metrics (**Fig.**
339 **7**). During the period 2001-2020, the mean SOS of all thresholds significantly advanced
340 by 0.13-0.16 days year⁻¹ ($p < 0.05$); the mean EOS of 10% and 25% significantly
341 advanced by 0.03-0.35 days year⁻¹ ($p < 0.05$); the mean LOS of all thresholds
342 significantly extended by 0.16-0.51 days year⁻¹ ($p < 0.01$). These findings are consistent
343 with previous studies (Zhu et al. 2012, Liu et al. 2016). For example, Liu et al. (2016)
344 indicated that the EOS delayed by 0.18 days year⁻¹ across the Northern Hemisphere of
345 1982-2011.



346

347

348 **5. Data availability**

349 This dataset is divided into single and double growing seasons. The entire dataset is
350 deposited at the open-access repository Figshare
351 (<https://doi.org/10.6084/m9.figshare.17195009.v2>; Fang et al. 2021).

352

353

354 **6. Conclusions**

355 This study used a long-term (2001-2020) SIF-based GPP product (GOSIF-GPP) to
356 generate annual photosynthetic phenology of vegetation with a high spatial resolution
357 (0.05°) in the Northern Hemisphere. Here, we applied a method combining filter
358 smoothing and change point detection to determine the annual dynamics of phenology
359 metrics (i.e., SOS, EOS, and LOS). This method avoided the re-modeling of the GPP
360 time series and allowed the extraction of metrics with different thresholds in multiple
361 growing seasons. We provided data users with three choices (10%, 25%, and 50%



threshold) of the metrics most appropriate for their specific application. Overall, the photosynthetic phenology metrics based on GOSIF-GPP agree with those extracted from in situ observations of EC towers. Compared to the metrics of vegetation indices and MODIS-GPP, the GOSIF-GPP metrics can provided more accurate phenology in most EC tower sites. The comparison with field data acquired at the EC towers suggests the 25% threshold of GOSIF-GPP can better capture the dynamics of photosynthetic phenology than other thresholds. In addition, the results showed a spatially explicit pattern from the north to the south in Northern Hemisphere. The SOS of all thresholds presented a significant advanced trend in the past 20 years; the EOS of 50% threshold showed an insignificant delayed trend; the LOS of all thresholds had a significant extended trend.

The phenology product based on GOSIF-GPP in our study is of great use in vegetation phenology studies because the SIF can directly reveal seasonal variations in vegetation vital activities (Mohammed et al. 2019). With these metrics, the response of vegetation phenology to climate change can be further investigated such as the importance of precipitation in spring phenology (Li et al. 2021). It will also be useful



for developing and validating dynamic vegetation models. Our phenology metrics could be further improved when more accurate SIF-based GPP estimates are available.

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Reference

- Badgley, G., Field, C. B., & Berry, J. A. 2017. Canopy near-infrared reflectance and terrestrial photosynthesis. *Science Advances*, **3**:e1602244.
- Baldocchi, D., Falge, E., Gu, L., Olson, R., Hollinger, D., Running, S., ... & Wofsy, S. 2001. FLUXNET: A new tool to study the temporal and spatial variability of ecosystem-scale carbon dioxide, water vapor, and energy flux densities. *Bulletin of the American Meteorological Society*, **82**: 2415-2434.
- Bertani, G., F. H. Wagner, L. O. Anderson, and L. E. O. C. Aragão. 2017. Chlorophyll Fluorescence Data



- 398 Reveals Climate-Related Photosynthesis Seasonality in Amazonian Forests. *Remote Sensing*
 399 **9**:1275.
- 400 Chen, A., J. Mao, D. Ricciuto, D. Lu, J. Xiao, X. Li, P. E. Thornton, and A. K. Knapp. 2021. Seasonal changes
 401 in GPP/SIF ratios and their climatic determinants across the Northern Hemisphere. *Global*
 402 *Change Biology* **27**:5186-5197.
- 403 Cohen, J. L., J. C. Furtado, M. Barlow, V. A. Alexeev, and J. E. Cherry. 2012. Asymmetric seasonal
 404 temperature trends. *Geophysical Research Letters* **39**.
- 405 Dai, J., M. Zhu, W. Mao, R. Liu, H. Wang, J. M. Alatalo, Z. Tao, and Q. Ge. 2021. Divergent changes of the
 406 elevational synchronicity in vegetation spring phenology in North China from 2001 to 2017 in
 407 connection with variations in chilling. *International Journal of Climatology* **41**:6109-6121.
- 408 Efron, B. 1992. Bootstrap Methods: Another Look at the Jackknife. Pages 569-593 in S. Kotz and N. L.
 409 Johnson, editors. *Breakthroughs in Statistics: Methodology and Distribution*. Springer New
 410 York, New York, NY.
- 411 Elmore, A. J., S. M. Guinn, B. J. Minsley, and A. D. Richardson. 2012. Landscape controls on the timing of
 412 spring, autumn, and growing season length in mid-Atlantic forests. *Global Change Biology*
 413 **18**:656-674.
- 414 Fang, J., J. A. Lutz, L. Wang, H. H. Shugart, and X. Yan. 2020. Using climate-driven leaf phenology and
 415 growth to improve predictions of gross primary productivity in North American forests. *Global*
 416 *Change Biology* **26**:6974-6988.
- 417 Fang, J., Li, X., X. Xiao, J., X. Yan, B. Li, F. Liu. 2021. Vegetation photosynthetic phenology metrics in
 418 northern terrestrial ecosystems: a dataset derived from a gross primary productivity product
 419 based on solar-induced chlorophyll fluorescence. figshare. Dataset.
 420 <https://doi.org/10.6084/m9.figshare.17195009.v1>
- 421 Frankenberg, C., C. O'Dell, J. Berry, L. Guanter, J. Joiner, P. Köhler, R. Pollock, and T. E. Taylor. 2014.
 422 Prospects for chlorophyll fluorescence remote sensing from the Orbiting Carbon Observatory-
 423 2. *Remote Sensing of Environment* **147**:1-12.
- 424 Garrity, S. R., G. Bohrer, K. D. Maurer, K. L. Mueller, C. S. Vogel, and P. S. Curtis. 2011. A comparison of
 425 multiple phenology data sources for estimating seasonal transitions in deciduous forest carbon
 426 exchange. *Agricultural and Forest Meteorology* **151**:1741-1752.
- 427 Gu, L., Post, W. M., Baldocchi, D. D., Black, T. A., Suyker, A. E., Verma, S. B., ... & Wofsy, S. C. 2009.
 428 Characterizing the seasonal dynamics of plant community photosynthesis across a range of
 429 vegetation types. In *Phenology of ecosystem processes* (pp. 35-58). Springer, New York, NY.
- 430 Huang, X., J. Xiao, and M. Ma. 2019. Evaluating the Performance of Satellite-Derived Vegetation Indices



- 431 for Estimating Gross Primary Productivity Using FLUXNET Observations across the Globe.
 432 Remote Sensing **11**:1823.
- 433 Joiner, J., Y. Yoshida, A. P. Vasilkov, K. Schaefer, M. Jung, L. Guanter, Y. Zhang, S. Garrity, E. M. Middleton,
 434 K. F. Huemmrich, L. Gu, and L. Belelli Marchesini. 2014. The seasonal cycle of satellite
 435 chlorophyll fluorescence observations and its relationship to vegetation phenology and
 436 ecosystem atmosphere carbon exchange. Remote Sensing of Environment **152**:375-391.
- 437 Joiner, J., Y. Yoshida, A. P. Vasilkov, Y. Yoshida, L. A. Corp, and E. M. Middleton. 2011. First observations
 438 of global and seasonal terrestrial chlorophyll fluorescence from space. Biogeosciences **8**:637-
 439 651.
- 440 Keenan, T. F., A. D. Richardson, and K. Hufkens. 2020. On quantifying the apparent temperature
 441 sensitivity of plant phenology. New Phytologist **225**:1033-1040.
- 442 Killick, R., P. Fearnhead, and I. A. Eckley. 2012. Optimal detection of changepoints with a linear
 443 computational cost. Journal of the American Statistical Association **107**:1590-1598.
- 444 Klosterman, S. T., K. Hufkens, J. M. Gray, E. Melaas, O. Sonnentag, I. Lavine, L. Mitchell, R. Norman, M.
 445 A. Friedl, and A. D. Richardson. 2014. Evaluating remote sensing of deciduous forest phenology
 446 at multiple spatial scales using PhenoCam imagery. Biogeosciences **11**:4305-4320.
- 447 Köhler, P., Frankenberg, C., Magney, T. S., Guanter, L., Joiner, J., & Landgraf, J. 2018. Global retrievals of
 448 solar-induced chlorophyll fluorescence with TROPOMI: First results and intersensor
 449 comparison to OCO-2. Geophysical Research Letters, **45**: 10-456.
- 450 Li, X., Y. H. Fu, S. Chen, Xiao J., G. Yin, X. Li, X. Zhang, X. Geng, Z. Wu, X. Zhou, J. Tang, and F. Hao. 2021.
 451 Increasing importance of precipitation in spring phenology with decreasing latitudes in
 452 subtropical forest area in China. Agricultural and Forest Meteorology **304-305**:108427.
- 453 Li, X., Xiao, J., He, B., Altaf Arain, M., Beringer, J., Desai, A. R., ... & Varlagin, A. 2018. Solar-induced
 454 chlorophyll fluorescence is strongly correlated with terrestrial photosynthesis for a wide variety
 455 of biomes: First global analysis based on OCO-2 and flux tower observations. Global change
 456 biology, **24**: 3990-4008.
- 457 Li, X., and J. Xiao. 2019. A global, 0.05-degree product of solar-induced chlorophyll fluorescence derived
 458 from OCO-2, MODIS, and reanalysis data. Remote Sensing **11**: 517.
- 459 Li, X., and J. Xiao. 2020. Global climatic controls on interannual variability of ecosystem productivity:
 460 Similarities and differences inferred from solar-induced chlorophyll fluorescence and enhanced
 461 vegetation index. Agricultural and Forest Meteorology **288-289**:108018.
- 462 Li, X., Y. Zhou, L. Meng, G. R. Asrar, C. Lu, and Q. Wu. 2019. A dataset of 30 m annual vegetation
 463 phenology indicators (1985–2015) in urban areas of the conterminous United States. Earth



- 464 System Science Data **11**:881-894.
- 465 Liu, Q., Y. H. Fu, Z. Zhu, Y. Liu, Z. Liu, M. Huang, I. A. Janssens, and S. Piao. 2016. Delayed autumn
 466 phenology in the Northern Hemisphere is related to change in both climate and spring
 467 phenology. *Global Change Biology* **22**:3702-3711.
- 468 Meroni, M., M. Rossini, L. Guanter, L. Alonso, U. Rascher, R. Colombo, and J. Moreno. 2009. Remote
 469 sensing of solar-induced chlorophyll fluorescence: Review of methods and applications.
 470 *Remote Sensing of Environment* **113**:2037-2051.
- 471 Mohammed, G. H., R. Colombo, E. M. Middleton, U. Rascher, C. van der Tol, L. Nedbal, Y. Goulas, O.
 472 Pérez-Priego, A. Damm, M. Meroni, J. Joiner, S. Cogliati, W. Verhoef, Z. Malenovsky, J.-P.
 473 Gastellu-Etchegorry, J. R. Miller, L. Guanter, J. Moreno, I. Moya, J. A. Berry, C. Frankenberg, and
 474 P. J. Zarco-Tejada. 2019. Remote sensing of solar-induced chlorophyll fluorescence (SIF) in
 475 vegetation: 50 years of progress. *Remote Sensing of Environment* **231**:111177.
- 476 Pastorello, G., Trotta, C., Canfora, E., Chu, H., Christianson, D., Cheah, Y. W., ... & Law, B. 2020. The
 477 FLUXNET2015 dataset and the ONEFlux processing pipeline for eddy covariance data. *Scientific*
 478 *data* **7**:225.
- 479 Piao, S., Q. Liu, A. Chen, I. A. Janssens, Y. Fu, J. Dai, L. Liu, X. Lian, M. Shen, and X. Zhu. 2019. Plant
 480 phenology and global climate change: Current progresses and challenges. *Global Change*
 481 *Biology* **25**:1922-1940.
- 482 Richardson, A. D., R. S. Anderson, M. A. Arain, A. G. Barr, G. Bohrer, G. Chen, J. M. Chen, P. Ciais, K. J.
 483 Davis, A. R. Desai, M. C. Dietze, D. Dragoni, S. R. Garrity, C. M. Gough, R. Grant, D. Y. Hollinger,
 484 H. A. Margolis, H. McCaughey, M. Migliavacca, R. K. Monson, J. W. Munger, B. Poulter, B. M.
 485 Raczka, D. M. Ricciuto, A. K. Sahoo, K. Schaefer, H. Tian, R. Vargas, H. Verbeeck, J. Xiao, and Y.
 486 Xue. 2012. Terrestrial biosphere models need better representation of vegetation phenology:
 487 results from the North American Carbon Program Site Synthesis. *Global Change Biology*
 488 **18**:566-584.
- 489 Richardson, A. D., K. Hufkens, T. Milliman, D. M. Aubrecht, M. Chen, J. M. Gray, M. R. Johnston, T. F.
 490 Keenan, S. T. Klosterman, and M. Kosmala. 2018. Tracking vegetation phenology across diverse
 491 North American biomes using PhenoCam imagery. *Scientific data* **5**:1-24.
- 492 Richardson, A. D., T. F. Keenan, M. Migliavacca, Y. Ryu, O. Sonnentag, and M. Toomey. 2013. Climate
 493 change, phenology, and phenological control of vegetation feedbacks to the climate system.
 494 *Agricultural and Forest Meteorology* **169**:156-173.
- 495 Seyedinasrollah, B., D. R. Bowling, R. Cheng, B. A. Logan, T. S. Magney, C. Frankenberg, J. C. Yang, A. M.
 496 Young, K. Hufkens, M. A. Arain, T. A. Black, P. D. Blanken, R. Bracho, R. Jassal, D. Y. Hollinger, B.



- 497 E. Law, Z. Nesic, and A. D. Richardson. 2021. Seasonal variation in the canopy color of
 498 temperate evergreen conifer forests. *New Phytologist* **229**:2586-2600.
- 499 Sjöström, M., M. Zhao, S. Archibald, A. Arneth, B. Cappelaere, U. Falk, A. de Grandcourt, N. Hanan, L.
 500 Kergoat, W. Kutsch, L. Merbold, E. Mougin, A. Nickless, Y. Nouvellon, R. J. Scholes, E. M.
 501 Veenendaal, and J. Ardö. 2013. Evaluation of MODIS gross primary productivity for Africa using
 502 eddy covariance data. *Remote Sensing of Environment* **131**:275-286.
- 503 Smith, W. K., J. A. Biederman, R. L. Scott, D. J. P. Moore, M. He, J. S. Kimball, D. Yan, A. Hudson, M. L.
 504 Barnes, N. MacBean, A. M. Fox, and M. E. Litvak. 2018. Chlorophyll Fluorescence Better
 505 Captures Seasonal and Interannual Gross Primary Productivity Dynamics Across Dryland
 506 Ecosystems of Southwestern North America. *Geophysical Research Letters* **45**:748-757.
- 507 Tang, J., Zeng, J., Zhang, Q., Zhang, R., Leng, S., Zeng, Y., ... & Wang, Q. 2020. Self-adapting extraction of
 508 cropland phenological transitions of rotation agroecosystems using dynamically fused NDVI
 509 images. *International journal of biometeorology*, 64: 1273-1283.
- 510 Verma, M., D. Schimel, B. Evans, C. Frankenberg, J. Beringer, D. T. Drewry, T. Magney, I. Marang, L. Hutley,
 511 C. Moore, and A. Eldering. 2017. Effect of environmental conditions on the relationship
 512 between solar-induced fluorescence and gross primary productivity at an OzFlux grassland site.
 513 *Journal of Geophysical Research: Biogeosciences* **122**:716-733.
- 514 Walther, S., Voigt, M., Thum, T., Gonsamo, A., Zhang, Y., Köhler, P., ... & Guanter, L. 2016. Satellite
 515 chlorophyll fluorescence measurements reveal large-scale decoupling of photosynthesis and
 516 greenness dynamics in boreal evergreen forests. *Global change biology*, **22**: 2979-2996.
- 517 Wang, X., Xiao, J., Li, X., Cheng, G., Ma, M., Che, T., ... & Wu, J. 2017. No consistent evidence for
 518 advancing or delaying trends in spring phenology on the Tibetan Plateau. *Journal of*
 519 *Geophysical Research: Biogeosciences*, **122**: 3288-3305.
- 520 Wang, X., Xiao, J., Li, X., Cheng, G., Ma, M., Zhu, G., Arain, M.A., Black, T.A., Jassal, R.S. 2019. No trends
 521 in spring and autumn phenology during the global warming hiatus. *Nature communications* **10**:
 522 1-10.
- 523 Wood, J. D., T. J. Griffis, J. M. Baker, C. Frankenberg, M. Verma, and K. Yuen. 2017. Multiscale analyses
 524 of solar-induced florescence and gross primary production. *Geophysical Research Letters*
 525 **44**:533-541.
- 526 Wu, C., D. Peng, K. Soudani, L. Siebicke, C. M. Gough, M. A. Arain, G. Bohrer, P. M. Lafleur, M. Peichl, A.
 527 Gonsamo, S. Xu, B. Fang, and Q. Ge. 2017. Land surface phenology derived from normalized
 528 difference vegetation index (NDVI) at global FLUXNET sites. *Agricultural and Forest*
 529 *Meteorology* **233**:171-182.



530 Xiao, J., Chevallier, F., Gomez, C., Guanter, L., Hicke, J.A., Huete, A.R., Ichii, K., Ni, W., Pang, Y., Rahman,
 531 A.F., Sun, G., Yuan, W., Zhang, L., Zhang, X. 2019. Remote sensing of the terrestrial carbon cycle:
 532 A review of advances over 50 years. *Remote Sensing of Environment*, 233:111383.
 533 Yang, L., and A. Noormets. 2021. Standardized flux seasonality metrics: a companion dataset for
 534 FLUXNET annual product. *Earth System Science Data* **13**:1461-1475.
 535 Zhao, M., Heinsch, F. A., Nemani, R. R., & Running, S. W. 2005. Improvements of the MODIS terrestrial
 536 gross and net primary production global data set. *Remote Sensing of Environment*, **95**:164-176.
 537 Zhao, R.-F., X.-P. Chen, F.-S. Zhang, H. Zhang, J. Schroder, and V. Römheld. 2006. Fertilization and Nitrogen
 538 Balance in a Wheat–Maize Rotation System in North China. *Agronomy Journal* **98**:938-945.
 539 Zhu, W., Tian, H., Xu, X., Pan, Y., Chen, G., & Lin, W. 2012. Extension of the growing season due to delayed
 540 autumn over mid and high latitudes in North America during 1982–2006. *Global Ecology and*
 541 *Biogeography*, **21**: 260-271.
 542
 543

Table 1. Statistical comparison of the phenology metrics retrieved from EC tower GPP and GOSIF-GPP, NDVI, EVI, NIR_v, and MODIS-GPP. 10%, 25%, and 50% mean the three thresholds. The bold means the highest *R* and the lowest *RMSE*. *R*: correlation coefficient; *RMSE*: root mean square error.

Data source	SOS			EOS			LOS		
	10%	25%	50%	10%	25%	50%	10%	25%	50%
	<i>R</i>								
GOSIF-GPP	0.79	0.80	0.78	0.63	0.73	0.63	0.72	0.76	0.65
NDVI	0.14	0.25	0.39	0.45	0.42	0.56	0.28	0.32	0.40
EVI	0.40	0.46	0.57	0.57	0.60	0.66	0.37	0.37	0.38
NIR _v	0.47	0.51	0.60	0.63	0.66	0.67	0.51	0.48	0.41
MODIS-GPP	0.66	0.67	0.65	0.29	0.55	0.61	0.47	0.55	0.49
	<i>RMSE (days)</i>								
GOSIF-GPP	18.03	15.83	14.99	23.55	21.89	24.38	33.93	29.14	27.89



NDVI	36.91	32.18	26.20	34.13	39.68	41.86	53.87	53.92	52.39
EVI	36.97	31.34	24.97	31.75	29.41	26.09	58.53	49.04	39.28
NIRv	29.86	26.00	21.74	27.26	25.67	24.95	46.11	40.03	35.12
MODIS-GPP	22.98	20.56	18.83	30.22	24.76	23.88	43.79	36.33	32.25

Table 2. The mean value and uncertainty of photosynthetic phenology metrics in the different terrestrial ecosystems.

Terrestrial ecosystems	Threshold	Mean SOS (uncertainty)	Mean EOS (uncertainty)	Mean LOS (uncertainty)
Forests	10%	108.26 (7.34)	271.29 (18.39)	163.03 (25.73)
	25%	122.12 (8.28)	255.40 (17.31)	133.28 (25.59)
	50%	138.44 (9.38)	236.24 (16.01)	97.80 (25.40)
Shrublands	10%	75.10 (5.09)	144.33 (9.78)	69.22 (14.88)
	25%	80.97 (5.49)	137.14 (9.30)	56.17 (14.79)
	50%	88.14 (5.97)	129.02 (8.75)	40.89 (14.72)
Savannas	10%	106.37 (7.21)	244.25 (16.56)	137.88 (23.77)
	25%	117.25 (7.95)	230.10 (15.60)	112.85 (23.55)
	50%	130.85 (8.87)	213.10 (14.45)	82.24 (23.32)
Grasslands	10%	56.72 (3.84)	153.48 (10.40)	96.76 (14.25)
	25%	65.21 (4.42)	140.37 (9.52)	75.16 (13.94)
	50%	74.71 (5.06)	127.55 (8.65)	52.84 (13.71)
Wetlands	10%	106.58 (7.23)	213.13 (14.45)	106.55 (21.67)
	25%	115.24 (7.81)	202.70 (13.70)	86.83 (21.51)
	50%	126.24 (8.56)	189.61 (12.85)	63.37 (21.41)
Croplands	10%	86.60 (5.87)	272.47 (18.47)	185.87 (24.34)
	25%	102.20 (6.93)	250.78 (17.00)	148.58 (23.93)
	50%	120.56 (8.17)	226.01 (15.32)	105.45 (23.49)

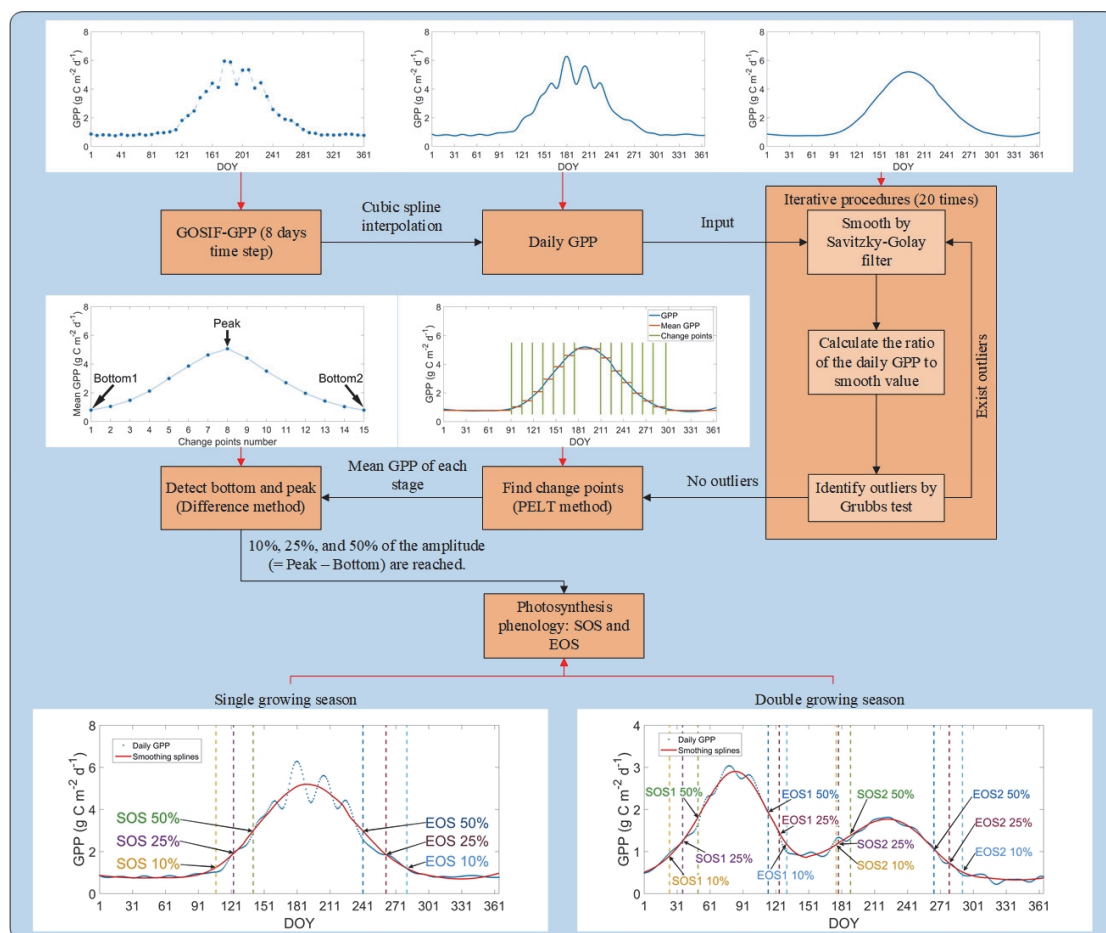


Fig. 1. Illustration of the method for identifying the transition dates of photosynthetic phenology. The method is based on three thresholds, 10%, 25%, and 50%. Bottom1: baseline for dormancy season before the growing season; Peak: the peak value in one single cycle; Bottom2: baseline for dormancy season after growing season. The example of the single growing season is from one forest site (latitude: 60.0° N, longitude: 15.5° E); the example of the double growing season is from one cropland site (latitude: 36.5° N, longitude: 36.0° E).

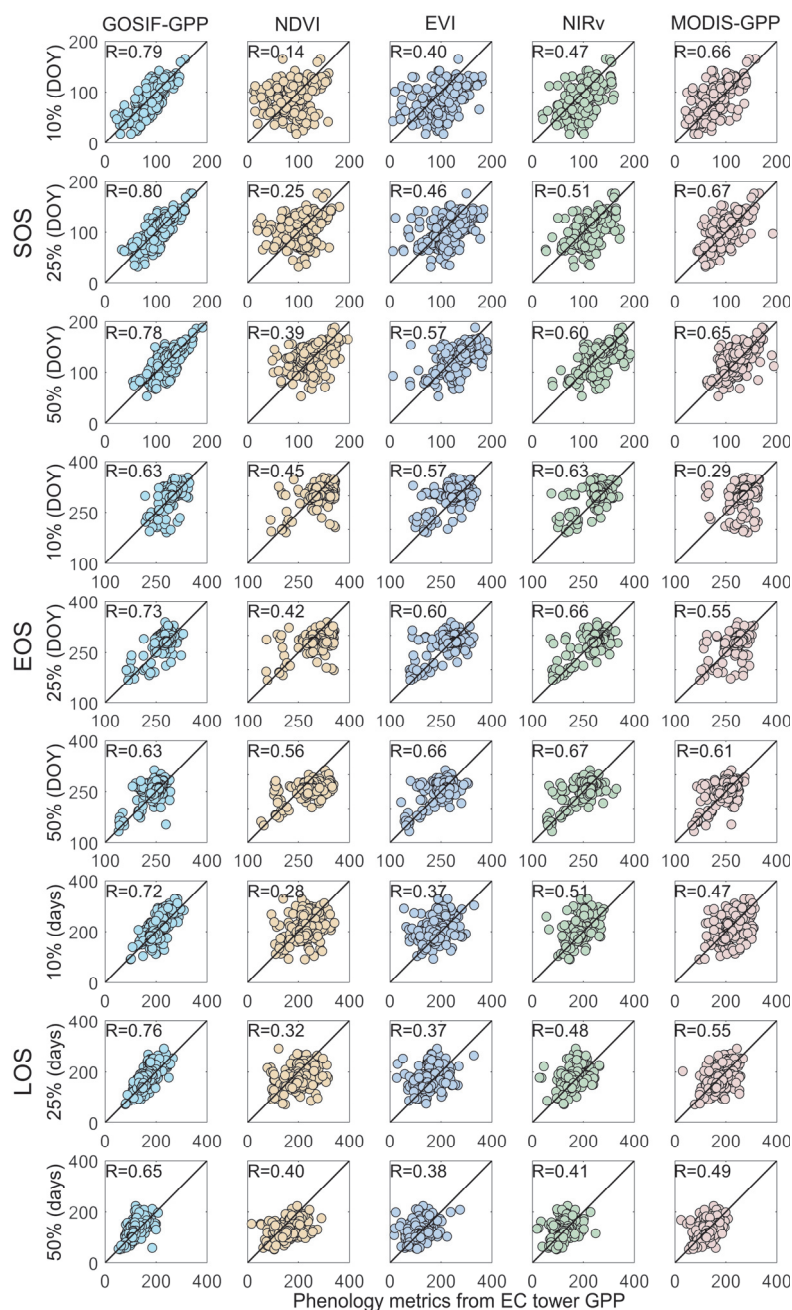


Fig. 2. The comparison of the phenology metrics retrieves from EC tower GPP and GOSIF-GPP, NDVI, EVI, NIRv, and MODIS-GPP. The dotted line represents a 1:1 line. DOY: day of the year; R: correlation coefficient.

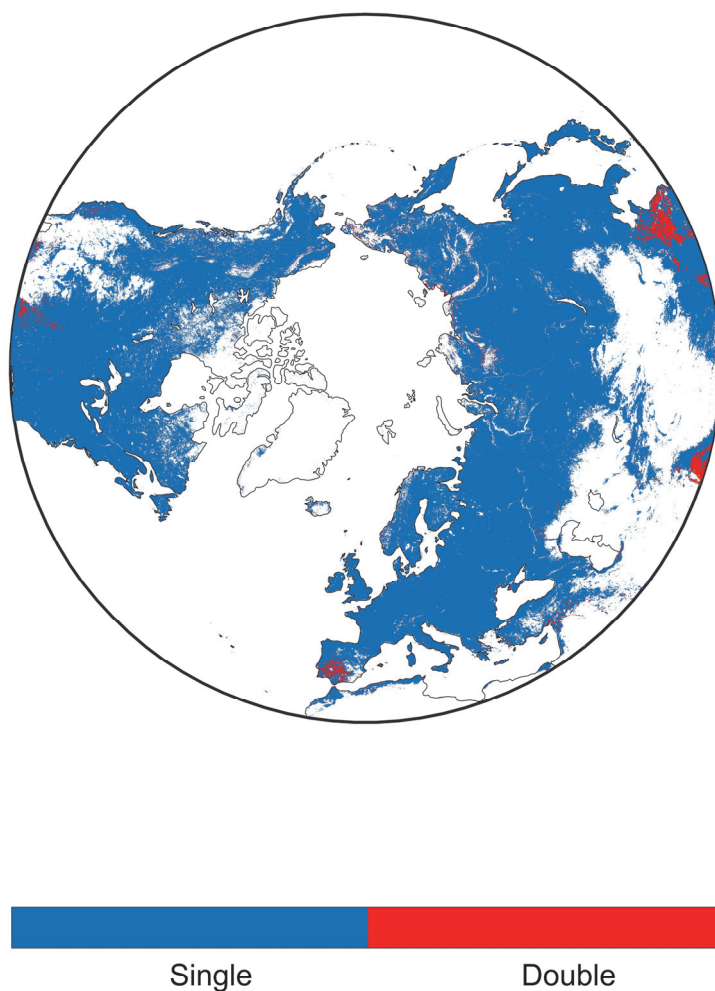


Fig. 3. The spatial distribution of the number of growing seasons in the Northern Hemisphere (0.05° spatial resolution).

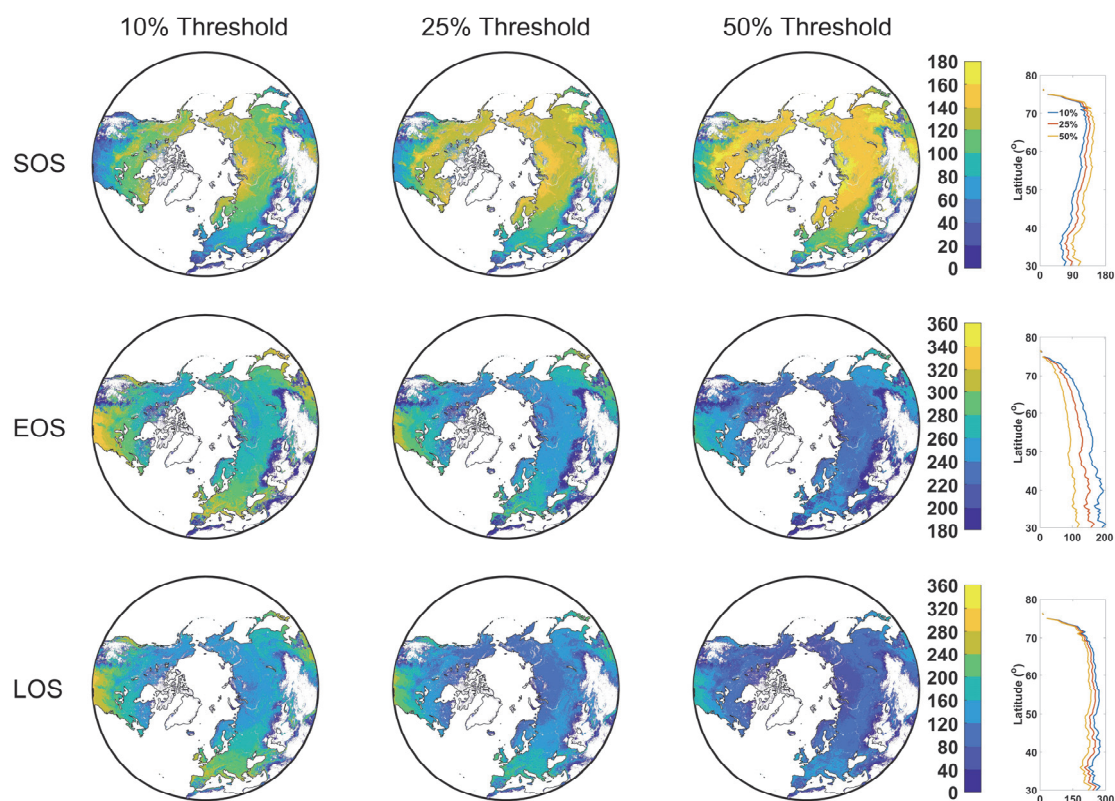


Fig. 4. The spatial distribution of the mean photosynthetic phenology metrics (first growing season) in the Northern Hemisphere of 2001-2020 (0.05° spatial resolution). The right parts are the latitudinal pattern.

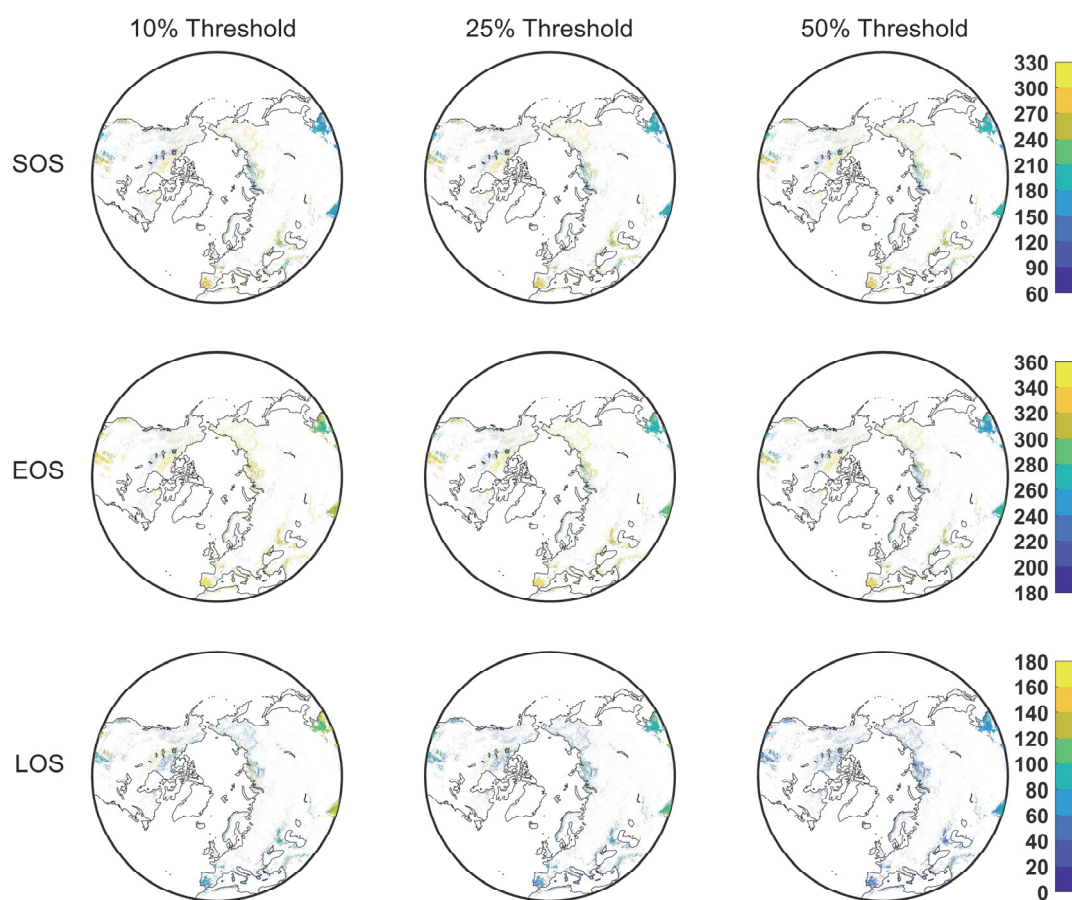


Fig. 5. The spatial distribution of the mean photosynthetic phenology metrics of the second growing season in the Northern Hemisphere of 2001-2020 (0.05° spatial resolution).

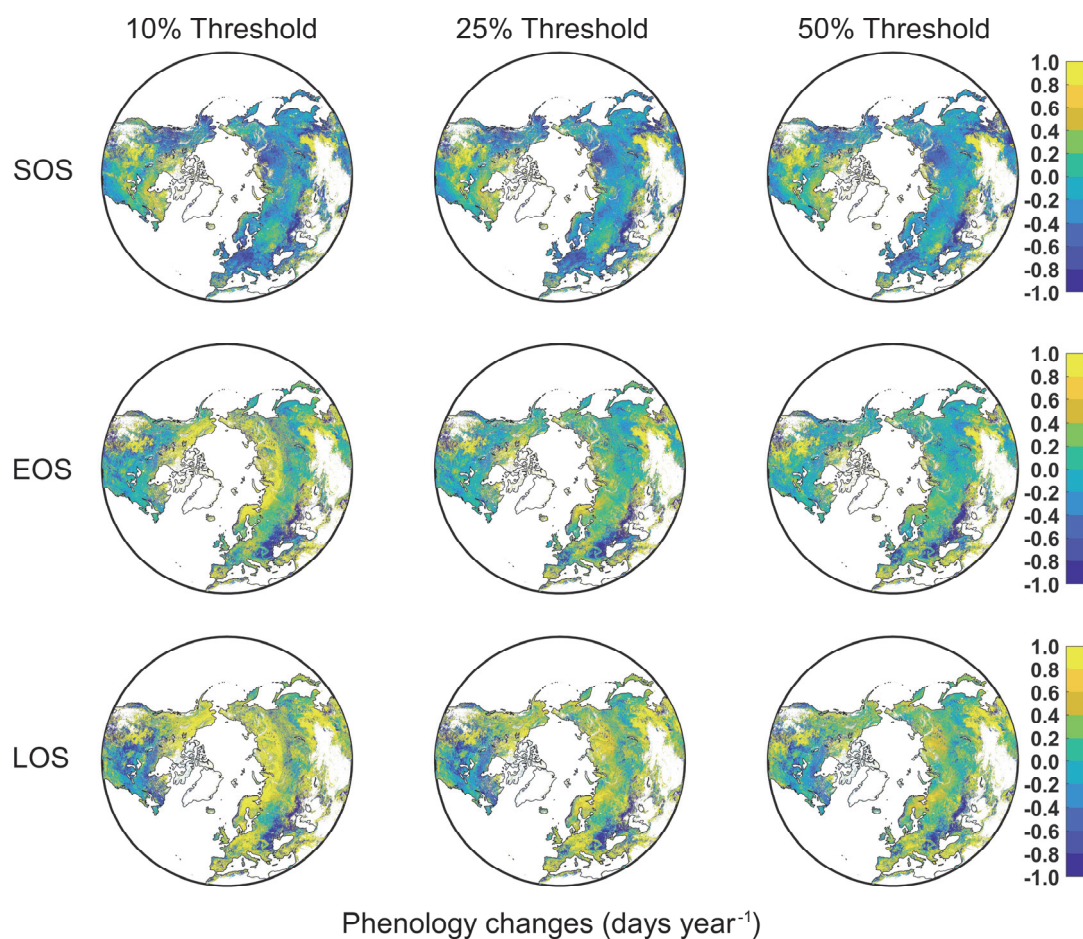


Fig. 6. Changes in photosynthetic phenology metrics in the Northern Hemisphere over the period 2001-2020.

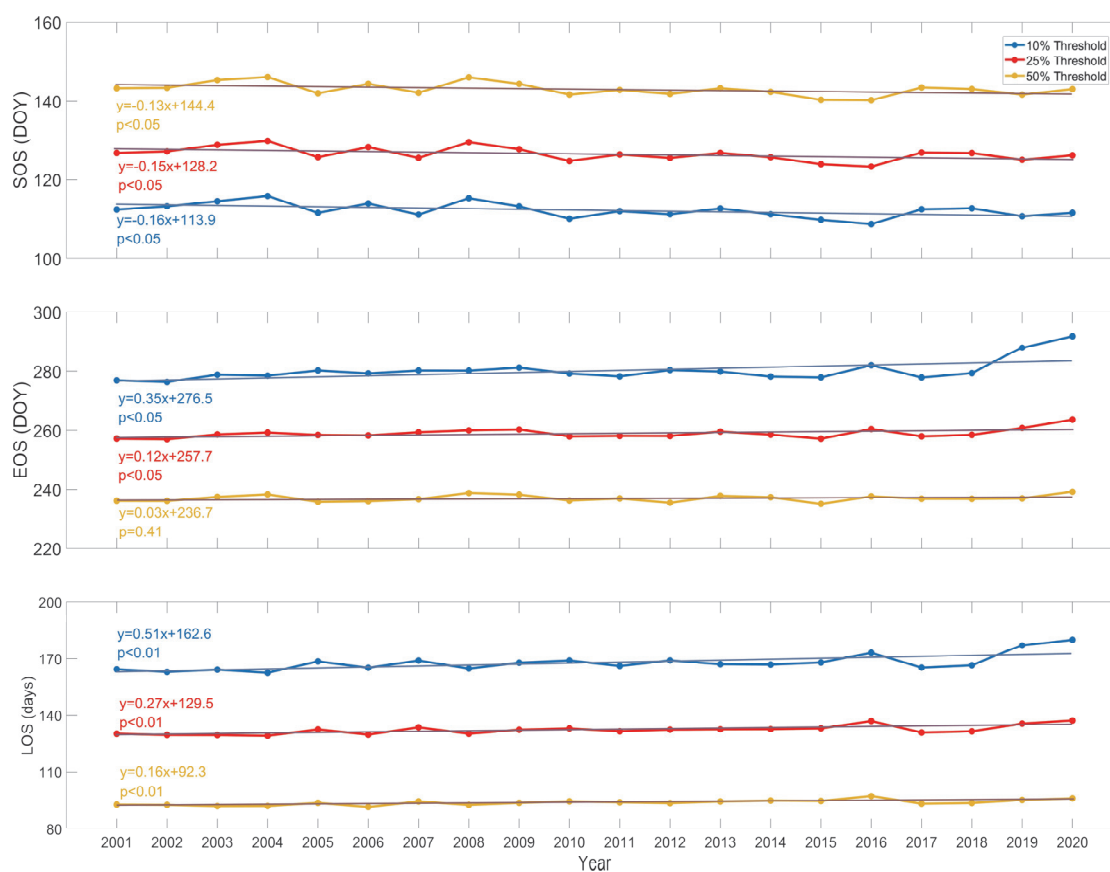


Fig. 7. Annual photosynthetic phenology metrics in the Northern Hemisphere during 2001-2020. The straight lines represent the change trends.