Photoperiod and Vernalization Control of Flowering-Related Genes: A Case Study of the Narrow-Leafed Lupin (Lupinus angustifolius L.)

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Narrow-leafed lupin (Lupinus angustifolius L.) is a moderate-yielding legume crop known for its high grain protein content and contribution to soil improvement. It is cultivated under photoperiods ranging from 9 to 17 h, as a spring-sown (in colder locations) or as an autumn-sown crop (in warmer regions). Wild populations require a prolonged cold period, called vernalization, to induce flowering. The key achievement of L. angustifolius domestication was the discovery of two natural mutations (named Ku and Jul) conferring vernalization independence. These mutations are overlapping deletion variants in the promoter of LanFTc1, a homolog of the Arabidopsis thaliana FLOWERING LOCUS T (FT) gene. The third deletion, named here as Pal, was recently found in primitive germplasm. In this study, we genotyped L. angustifolius germplasm that differs in domestication status and geographical origin for LanFTc1 alleles, which we then phenotyped to establish flowering time and vernalization responsiveness. The Ku and Jul lines were vernalization-independent and early flowering, wild (ku) lines were vernalization-dependent and late flowering, whereas the Pal line conferred intermediate phenotype. Three lines representing Ku, Pal, and ku alleles were subjected to gene expression surveys under 8- and 16-h photoperiods.

FT homologs (LanFTa1, LanFTa2, LanFTc1, and LanFTc2) and some genes selected by recent expression quantitative trait loci mapping were analyzed. Expression profiles of LanFTc1 and LanAGL8 (AGAMOUS-like 8) matched observed differences in flowering time between genotypes, highlighted by high induction after vernalization in the ku line. Moreover, these genes revealed altered circadian clock control in Pal line under short days. LanFD (FD) and LanCRLK1 (CALCIUM/CALMODULIN-REGULATED RECEPTOR-LIKE KINASE 1) were negatively responsive to vernalization in Ku and Pal lines but positively responsive or variable in ku, whereas LanUGT85A2 (UDP-GLUCOSYL TRANSFERASE 85A2) was significantly suppressed by vernalization in all lines. Such a pattern suggests the opposite regulation of these gene pairs in the vernalization pathway. LanCRLK1 and LanUGT85A2 are homologs of A. thaliana genes involved in the FLOWERING LOCUS C (FLC) vernalization
pathway. Lupins, like many other legumes, do not have any FLC homologs. Therefore, candidate genes surveyed in this study, namely LanFTc1, LanAGL8, LanCRLK1, and LanUGT85A2, may constitute anchors for further elucidation of molecular components contributing to vernalization response in legumes.

Keywords: vernalization, photoperiod, flowering, expression, FLOWERING LOCUS T, duplication, deletion

INTRODUCTION

Narrow-leafed lupin (Lupinus angustifolius L.) is a legume plant that is cultivated as “green” manure or as a grain crop for animal feed or human consumption (Kurlovich, 2002). The exploitation of L. angustifolius as a crop is advantageous in many respects. First, L. angustifolius cultivation has a positive influence on soil fertility due to the mobilization of soil-bound phosphorus and diazotrophic nitrogen fixation (Evans et al., 1987; Lambers et al., 2013). Moreover, L. angustifolius grains are characterized by high protein content and the use of them in livestock farming systems has many benefits in terms of the economic and environmental impact (Abraham et al., 2019). As seed alkaloid content was reduced by breeding below 0.01% of seed dry weight, a bitter taste, a typical feature of lupins, was fully eliminated from modern cultivars (Kamel et al., 2016). Additionally, the postharvest stubble can be now safely grazed by livestock, because the risk of lupinosis disease was diminished by the introduction of Phomosipis stem blight resistance genes (Mulholland et al., 1976; Jago et al., 1982; Cowling et al., 1987; Williamson et al., 1994; Shankar et al., 2002; Książkiewicz et al., 2020). Furthermore, L. angustifolius is currently being promoted in human food markets as a result of its nutritional, metabolomic, and other health benefits (Foyer et al., 2016; Kouris-Blazos and Belski, 2016).

Cultivated L. angustifolius germplasm primarily originated from the western Mediterranean basin (Mousavi-Derazmohalleh et al., 2018b). Natural adaptation of wild L. angustifolius populations to the Mediterranean climate is the requirement of the prolonged cold period (i.e., vernalization) during germination or juvenile phase to induce flowering (Gladstones and Hill, 1969; Rahman and Gladstones, 1972; Landers, 1995; Adhikari et al., 2012). The selection of early phenology was based on the removal of the vernalization requirement, a major achievement in the domestication of L. angustifolius, enabling temperature-independent sowing (French and Buirchell, 2005). L. angustifolius is grown in various environments ranging from the subtropics (Australia) and Mediterranean regions (including Morocco, Spain, southern France, and Italy), through temperate oceanic climates (such as the United Kingdom, northern France, and Benelux), humid continental climates (for example in Germany, the Baltic countries, and Ukraine), though to the subarctic zone, localized as far north as 60° latitude (Russia). In warmer regions it is usually autumn-sown, whereas in colder regions it is exclusively spring-sown (Annicchiario and Carroni, 2009).

Lupin crops are cultivated in various photoperiod conditions, ranging from about 9 h (autumn sowing in Australia) to 16–17 h (spring sowing in Baltic countries and Russia) (Książkiewicz et al., 2017). The vernalization requirements of wild populations are so demanding that they can only be completely fulfilled by spring sowing in northern locations (Adhikari et al., 2012). Worldwide, L. angustifolius cultivation is based on vernalization independence selected from only two natural mutations (named Ku and Jul) that were discovered in domesticated germplasm just a little over a century ago (Mikołajczyk, 1966; Gladstones and Hill, 1969). The use of only two donors of early flowering in L. angustifolius breeding, followed by a strong selection of key agronomic traits, have resulted in a lack of phenological diversity in domesticated germplasm (Stefanova and Buirchell, 2010; Berger et al., 2012; Cowling, 2020). Therefore, further adaptation of this crop to new agronomic conditions resulting from a rapidly changing climate, may only be possible with the incorporation of novel genetic resources of intermediate phenology.

During recent years L. angustifolius was supplemented with numerous molecular resources, including bacterial artificial chromosome libraries carrying nuclear DNA inserts, consecutively improved linkage maps with sequence-defined markers, transcriptome assemblies, and a progressively updated genome sequence of the reference cultivar Tanjil (Kasprzak et al., 2006; Nelson et al., 2006; Gao et al., 2011; Yang et al., 2016; Nelson et al., 2017). Screening of germplasm resources with the LanFTc1 allele, in the form of a 5162 bp deletion in the promoter region, enabled novel genetic resources of intermediate phenology. Interestingly, the second L. angustifolius early phenology mutation, Jul, was recently revealed to be the third LanFTc1 allele, in the form of a 5162 bp deletion in the promoter region, fully encompassing that 1423 bp Ku deletion (Taylor et al., 2019). This functional mutation underlying the domesticated Ku allele in L. angustifolius was assigned to the 1423 bp deletion in the promoter region of LanFTc1, carrying potential binding sites for several transcription factors acting as FT gene repressors in Arabidopsis thaliana (Nelson et al., 2017). Interestingly, the second L. angustifolius early phenology mutation, Jul, was recently revealed to be the third LanFTc1 allele, in the form of a 5162 bp deletion in the promoter region, fully encompassing that 1423 bp Ku deletion (Taylor et al., 2019). Screening of germplasm resources with the LanFTc1 markers resulted in the identification of a fourth LanFTc1 allele in wild population line originating from Palestine (a country annotated as Israel in Australian collection or Egypt in Polish gene bank), carrying 1208 bp deletion partially overlapping with domesticated Ku deletion (Rychel, 2018; Taylor et al., 2019). This Palestinian allele was named here as Pal.
with the stability of CONSTANS (CO) protein and shows two
peaks, the first in the morning (lower) and the second in
the evening (higher) (Turck et al., 2008). This CO-dependent
regulation is facilitated by enhancer protein binding to specific
sites in the promoter sequence, located approximately 5 kbp
upstream of the first codon, which results in forming a DNA
loop bringing all the components together (Cao et al., 2014).
Interestingly, L. angustifolius allelic variants of the LanFTc1
promoter have the distal binding sequences (CCCAAT-boxes)
preserved (Książkiewicz et al., 2016; Nelson et al., 2017; Taylor
et al., 2019). This study sought to explore whether the altered
length of the LanFTc1 promoter is associated with the altered
circadian clock control of LanFTc1 and other related genes.

Taking into consideration the (i) role of FT genes in
integrating key environmentally responsive pathways, (ii)
demonstrated allelic variability of LanFTc1 gene associated
with flowering time, and (iii) wide range of environmental
variables occurring at lupin cultivation sites, we decided to
explore the photoperiod and vernalization responsiveness in
L. angustifolius. Here, the L. angustifolius germplasm differing
in domestication status and geographical origin was genotyped
for LanFTc1 alleles and phenotyped for flowering time and
vernalization responsiveness. Then, the transcriptomic response
of candidate genes from the vernalization pathway in early
(Ku), intermediate (Pal), and late flowering (ku) L. angustifolius
germplasm was assayed under contrasting photoperiod and
vernalization conditions, accounting the influence of circadian
rhythm control. This study discusses the hypothetical function
of these genes in terms of flowering time regulation and response
to vernalization.

MATERIALS AND METHODS

Plant Material
Ninety-two L. angustifolius lines were subjected to genotyping
and phenotyping. Lines were derived from the European Lupin
Gene Resources Database maintained by the Poznań Plant
Breeding Ltd. station located in Wiatrowo. The lines originated
from 13 countries, including Spain (32 lines), Poland (15),
Australia (11), Russia (11), Germany (7), Italy (4), Belarus (3),
Israel (2), Algeria, Morocco, Palestine, Portugal and Republic
of South Africa (1). Accession numbers and information on
the domestication status and country of origin are provided in
Supplementary Table 1.

Identification of LanFTc1 Alleles
Young leaf tissue from three biological replicates per plant was
sampled. Frozen (−80°C) plant tissue (50 mg) was homogenized
using TissueLyser II (Qiagen, Hilden, Germany) and two
stainless steel beads (ø 5 mm) in 2 ml tubes (Eppendorf,
Hamburg, Germany). DNA was isolated using DNeasy Plant
Mini Kit (Qiagen). PCR was performed using GoTaq Long PCR
Master Mix (Promega, Mannheim, Germany) and published
LanFTc1_INDEL2 primers (Taylor et al., 2019), provided here
for reference in Supplementary Table 2. PCR conditions were
as follows: initial denaturation (94°C for 2 min), then 35 cycles
composed of denaturation (94°C for 30 s), annealing (62°C
for 30 s), and elongation (72°C for 5 min), followed by the
final extension (72°C for 10 min). Products were resolved
by agarose gel electrophoresis and SYBR Safe DNA staining
(Invitrogen, Carlsbad, CA, United States) and visualized on UV-
transilluminator (Uvitec, Thermo Fisher Scientific, Waltham,
MA, United States). Wild P27255 allele (ku, without deletion) was
encoded as “A,” Palestinian allele (Pal, 1208 bp deletion) as “B,”
83A:476 allele (Ku, 1423 bp deletion) as “C,” and Krasnolistsyn
allele (Jul, 5162 bp deletion) as “D.”

Evaluation of Vernalization
Responsiveness in Greenhouse
Vernalization was carried out by placing imbibed seeds for
21 days at 5°C in darkness on moist filter paper in Petri dishes.
Non-vernalized control plants were sown five days before the end
of the vernalization procedure and kept at ~21°C to maintain a
similar thermal time (Huyghe, 1991). Plants were cultivated in
a greenhouse maintained by the Institute of Plant Genetics, the
Polish Academy of Sciences, Poznań, Poland (52°26′N 16°54′E)
during growing seasons of 2014 (sowing of vernalized plants on
14.05) and 2015 (sowing of vernalized plants on 25.03) under
ambient long day photoperiod (12−17 h). The greenhouse was
equipped with automatic heating to keep the minimum air
temperature above 18°C. Passive cooling was maintained by a
temperature-dependent ventilation system (activated at 22°C).
Air temperature (daily mean and maximum) and daily sunshine
hours recorded by the nearby localized meteorological station
(Poznań-Lawica, 5.1 km) as well as the theoretical photoperiod
hours calculated for this latitude (covering 120 days from
sowing date for both years) were provided for reference in
Supplementary Tables 3, 4. Flowering time was recorded as the
number of days from sowing date of vernalized plants until the
first fully colored petal was observed. The average number of
plants sampled in 2014 was 5.6 for the non-vernalized variant
(min. 3, max. 6) and 6.9 for the vernalized variant (min. 4, max.
7), whereas in 2015 it was 4.7 for the non-vernalized variant (min.
4, max. 5) and 5.0 for the vernalized variant.

Controlled Environment Experiment for
Gene Expression Profiling
Based on the results of LanFTc1 allele genotyping and
vernalization responsiveness phenotyping, three accesses were
selected for gene expression profiling: P27255 from Morocco
(96,234, carrying wild allele ku), 83A:476 from Australia (96,233,
carrying domesticated allele Ku), and Palestyna from Palestine
(95,799, carrying intermediate allele Pal). The vernalization of
lines was performed as described above. Non-vernalized control
plants were sown five days before the end of the vernalization
procedure. Plants from both variants were cultivated in climatic
chambers with controlled humidity (40−50% day, 70−80%
night) and temperature (22°C day, 18°C night). Two types of
photoperiods were studied, short day (SD, 8 h, from 8 AM to
4 PM) and long day (LD, 16 h, from 4 AM to 8 PM). Young
leaves from five biological replicates were sampled every week
at two times of day to follow a circadian rhythm, at 9 AM, and
3 PM under SD or at 7 AM and 6 PM under LD. Plant material was immediately frozen in liquid nitrogen and stored at −80°C. Three replicates with similar plant phenology (growth rate and time to flowering) were subjected to gene expression profiling. Taking into consideration observed flowering time, from 2 to 4 terms were selected for gene expression, representing the period from about 2 weeks before flowering to the flowering date and/or few days after flowering (Supplementary Table 5).

RNA Isolation and cDNA Synthesis
Frozen young leaf tissue (50 mg) was homogenized using TissueLyser II (Qiagen) and two stainless steel beads (ø 5 mm) in 2 ml tubes (Eppendorf). RNA isolation was performed using the SV Total RNA Isolation System (Promega) without any alterations to the protocol. RNA concentration and purity were measured using NanoDrop 2000 (Thermo Fisher Scientific) and A260/A280 ratio. RNA integrity was visualized by 1% agarose gel electrophoresis of denatured samples in 1 × TAE buffer. RNA concentration was equalized to 1000 ng/µl in nuclease-free water. First-strand cDNA synthesis was performed using GoScript (TM) Reverse Transcription System (Promega) and 5 µg of total RNA per sample.

Selection of Genes for Quantitative PCR
The set of genes selected for quantitative PCR included, among others, four L. angustifolius representatives of FT clade, namely LanFTa1 (Lup021189, XM_019571501.1), LanFTa2 (XM_019596455.1), LanFTc1 (Lup015264, XM_019601808.1), and LanFTc2 (Lup005674, XM_019565316.1) (Książkiewicz et al., 2016; Nelson et al., 2017). First names in parentheses correspond to gene names provided in the L. angustifolius pseudochromosome assembly paper (Hane et al., 2017), whereas the second names address L. angustifolius NCBI Reference Sequences LupAngTanji_l_v1.0. Moreover, our assay endeavored also some candidate genes revealed by recent L. angustifolius (eQTL mapping) study (Plewiński et al., 2019) to be associated with the vernalization response, as follows: LanUGT85A2 (Lup002110, XM_019574900.1), LanCRLK1 (Lup011808, XM_019603391.1), LanAGL8 (Lup018485, XM_019583439.1), and LanFD (Lup018024, XM_019567853.1). Alignment of coding sequences to the L. angustifolius genome assembly provided evidence that LanAGL8 and LanFD are present in single copies.

Moreover, some major A. thaliana components of the flowering induction pathway were considered, as follows: LEAFY (LFY, AT5G61850), APETALA1 (API, AT1G69120), VERNALIZATION INSENSITIVE 3 (VIN3, AT3G57380) and VERNALIZATION 5 (VRN5, AT3G24440). Multiple sequence alignment revealed that all these genes have putatively three copies in the L. angustifolius genome, namely Lup006312 (XM_019602464.1), Lup012189 (XM_019558971.1) and Lup027481 (XM_019607325.1) for LFY; Lup021855 (XM_019605469.1), Lup023438 (XM_019588203.1) and Lup066876 (XM_019572960.1) for API; Lup069440 (XM_019586878.1), Lup013437 (XM_019598860.1) and Lup062125 (XM_019608083.1) for VIN3; and Lup009144 (XM_019591910.1), Lup018692 (XM_019567058.1) and Lup032778 (XM_019590742.1) for VRN5. Analysis of leaf transcriptome data obtained for the L. angustifolius linkage mapping population, covering developmental phases from juvenile to generative growth after partial vernalization (Plewiński et al., 2019), revealed negligible expression of all LFY and API homologs (~0.07 reads per kilobase million, RPKM), very low expression of VRN5 (~0.56 RPKM) and high expression of VIN3 copies (~25.07 RPKM) (Supplementary Table 6). Therefore, VRN5 and VIN3 homologs were selected for the expression assay, whereas LFY and API homologs were discarded. Reference genes validated in previous L. angustifolius quantitative gene expression studies were selected for this assay, namely LanDexH7 (Lup023733, XM_019579367.1), and LanTUB6 (Lup032899, XM_019581544.1) (Taylor et al., 2016, 2019; Nelson et al., 2017). Primers were designed in Geneious Prime (Auckland, New Zealand) using Primer3 (Kearse et al., 2012; Untergasser et al., 2012). Due to the high similarity between particular copies, all three VRN5 homologs were analyzed together using one primer pair. The remaining genes were profiled on one by one basis. Designed primers and expected product sizes are provided in Supplementary Table 2.

Quantitative Gene Expression Analysis
In prior experiments, a CFX Connect Real-Time PCR Detection System (Bio-Rad Polska, Warsaw, Poland) was calibrated using Melt Calibration Kit (Bio-Rad Polska) according to the manufacturer's protocol. A standard curve was developed to assess the performance of the quantitative PCR assay and its dynamic range following recent recommendations (Svec et al., 2015). Analyzed genes were amplified using GoTaq G2 Flexi DNA Polymerase (Promega) and subjected to 1% agarose gel electrophoresis. Amplicons were excised from a gel, extracted with the aid of QIAquick Gel Extraction Kit (Qiagen), quantified using NanoDrop 2000 (Thermo Fisher Scientific), and its dynamic range following recent recommendations (Svec et al., 2015). Analogous primers were amplified using GoTaq G2 Flexi DNA Polymerase (Promega) and subjected to 1% agarose gel electrophoresis. Amplicons were excised from a gel, extracted with the aid of QIAquick Gel Extraction Kit (Qiagen), quantified using NanoDrop 2000 (Thermo Fisher Scientific), and outsourcing (Genomed Ltd., Warsaw, Poland) for direct Sanger sequencing on ABI PRISM 3130 Genetic Analyzer XL (Applied Biosystems, Hitachi). A series of dilutions in concentrations ranging from 1 to 10−10 of the original templates were prepared for every gene using an initial volume of 20 µl to reduce the sampling error. 6 replicates per each concentration were performed using the iTaq Universal SYBR Green Supermix (Bio-Rad Polska). A two-step PCR protocol was exploited according to the protocol. A calculation of R² and PCR efficiency values was done in Bio-Rad CFX Manager 3.1. The values obtained are provided in Supplementary Table 7.

The quantitative PCR analysis of gene expression was performed using 96-well PCR plates and two reference genes (LanDexH7 and LanTUB6). Inter-run calibration sample (LanTUB6) and no template control were used on all plates. Three biological replicates were analyzed for each time point, and all samples were run in 5 technical repeats. High-resolution PCR product melting in the range of temperature from 65 to 85°C was performed after PCR to control the specificity of amplification. Melt profiles were inspected for the amplification of unspecific products, highlighted by the presence of melting peaks at different temperatures than those obtained during standard curve
Indel variation in the promoter region of the major flowering time gene LanFTc1 (Figure 1) was recently revealed to be associated with flowering time and vernalization responsiveness in L. angustifolius (Nelson et al., 2017; Taylor et al., 2019). To evaluate allelic composition in germplasm exploited by European lupin breeders, ninety-two L. angustifolius accessions, encompassing 43 primitive populations or landraces, 23 derivative or breeding lines, 25 cultivars, and one mutant, derived from the European Lupin Gene Resources Database, were screened with primers flanking polymorphic LanFTc1 promoter region. As expected, wild LanFTc1 allele (ku) was found mainly in primitive accessions collected in the Mediterranean Basin as well as in a few old domesticated materials originating from Russia, Poland, Germany, and the Republic of South Africa. One wild population line, Palestyna originating from Palestine, was found to carry the shortest variant of deletion (1208 bp), named here as Pal. Alleles Ku (1423 bp deletion) and Jul (5162 bp deletion) were found only in domesticated germplasm – the first one was present mostly in released cultivars, whereas the second one was typically in breeding materials at a different stage of improvement. Accessions carrying Ku originated primarily from Australia, Germany, and Poland, whereas those carrying Jul from Poland and Russia. Results of marker screening are provided in Supplementary Table 8.

Palestinian LanFTc1 Allele Confers Intermediate Flowering Time Moderately Responsive to Vernalization

Ninety-two L. angustifolius accessions with known LanFTc1 promoter allele composition were evaluated for phenotypic response to vernalization in two consecutive years, 2014 and

FIGURE 1 | Indel variation in the promoter sequence of LanFTc1 gene controlling flowering induction in L. angustifolius. Allele ku is typical for wild populations, alleles Jul and Ku are present only in domesticated germplasm, whereas allele Pal was found only in wild germplasm from Palestine. The position is given in relation to the first nucleotide of CCAAT-box (Książkiewicz et al., 2016; Nelson et al., 2017; Taylor et al., 2019).
TABLE 2 | Comparison of flowering time and vernalization responsiveness of *L. angustifolius* germplasm carrying *Jul*, *Ku*, *Pal*, and *ku* *LanFTc1* alleles, cultivated in a greenhouse under natural long days (LD).

| *LanFTc1* allele | Days to flowering ‘2014 | Vernalization response ‘2014 | Days to flowering ‘2015 | Vernalization response ‘2015 |
|------------------|------------------------|-----------------------------|------------------------|-----------------------------|
| *Ku*, without deletion | 88.8 ± 15.6 | −41.1 ± 13.8 | 88.0 ± 5.9 | −21.8 ± 6.0 |
| *Pal*, 1208 bp deletion | 53.5 ± 1.9 | −14.5 ± 2.7 | 68.2 ± 5.6 | −14.8 ± 3.4 |
| *Ku*, 1423 bp deletion | 40.5 ± 2.6 | −3.3 ± 2.6 | 56.7 ± 2.1 | −3.0 ± 2.5 |
| *Jul*, 5162 bp deletion | 38.8 ± 1.1 | −0.2 ± 2.2 | 55.6 ± 1.9 | −2.2 ± 2.2 |

| 2015 (Table 2). Lines carrying wild *Ku* allele revealed the longest vegetative phase, ranging from 61.2 ± 0.9 to 116.2 ± 3.0 days in ’2014 and from 79.0 ± 5.8 to 101.5 ± 3.9 days in ’2015. These lines demonstrated also high vernalization responsiveness, highlighted by the acceleration of flowering after vernalization by 13.9 to 66.4 days in 2014, and by 11.6 to 35.6 days in 2015. The line carrying *Pal* allele showed intermediate phenology and flowered about 20-35 days quicker than the average *Ku* genotype, namely after ~53 days from sowing in ’2014 and ~68 days in ’2015. The vernalization procedure advanced flowering induction in this line by about two weeks. Accessions carrying domesticated *Ku* allele were very early and started flowering after 37.2 ± 0.4 to 46.2 ± 8.9 days in 2014, and after 54.0 ± 0.0 to 59.6 ± 2.6 days in 2015. This set contained germplasm low-responsive to vernalization, which accelerated flowering time by up to ~8.6 days, as well as some truly thermoneutral accessions, which flowered at the same time regardless of the vernalization procedure. Lines carrying domesticated European *Jul* allele also revealed very early phenology, manifested by the onset of flowering after 37.0 ± 1.3 to 41.0 ± 5.9 days in 2014, and after 52.2 ± 1.5 to 60.8 ± 1.8 days in 2015. Most *Jul* lines were fully thermoneural. On average, *Jul* lines flowered earlier than *Ku* lines in both years, however taking into consideration standard errors resulting from variability between biological replicates, differences between mean values were not statistically significant. Results of time to flowering and vernalization responsiveness in relation to *LanFTc1* INDEL2 marker polymorphism are provided in Supplementary Table 8 (2014) and Supplementary Table 9 (2015).

Based on the results of *LanFTc1* INDEL2 marker screening and vernalization responsiveness, three lines were selected for vernalization response phenotyping under 8-h (SD) and 16-h (LD) photoperiods: 84A:476 carrying domesticated *Ku* allele (early flowering, thermoneutral), Palestyna carrying wild *Pal* allele (moderately flowering and responsive to vernalization), and P27255 carrying wild *Ku* allele (late flowering and highly responsive to vernalization). 83A:476 was revealed to be the earliest line in both photoperiods, followed by Palestyna (Table 3). Under SD, 83A:476 accelerated transition between phases in response to vernalization by about 5 days, whereas under LD by about 3 days. These responses were higher in Palestyna, amounting to about 12-19 days and 3-5 days, respectively. P27255 did not flower during the experiment (90 days) in all variants except vernalized plants under LD.

TABLE 3 | Number of days to first bud, flower, and pod in *L. angustifolius* germplasm carrying *Ku*, *Pal*, and *Ku* *LanFTc1* alleles, cultivated under 8- and 16-h photoperiods.

| *LanFTc1* allele | Vernalization variant | Days to first bud | Days to first flower | Days to first pod |
|------------------|-----------------------|------------------|---------------------|------------------|
| 8-h photoperiod   |                       |                  |                     |                  |
| *Ku*             | −                     | 51.3 ± 1.6     | 57.2 ± 4.3       | 63.3 ± 9.8      |
| *Ku*             | +                     | 45.7 ± 2.9     | 52.0 ± 4.2       | 58.1 ± 2.5      |
| *Pal*            | -                     | 65.3 ± 3.7     | 71.2 ± 3.8       | 83.5 ± 2.7      |
| *Pal*            | +                     | 52.9 ± 3.0     | 55.9 ± 2.6       | 64.7 ± 4.6      |
| *Ku*             | −                     | –                | –                   | –               |
| *Ku*             | +                     | –                | –                   | –               |
| 16-h photoperiod |                       |                  |                     |                  |
| *Ku*             | −                     | 33.6 ± 1.1     | 38.7 ± 1.3       | 44.1 ± 1.4      |
| *Ku*             | +                     | 29.8 ± 2.7     | 35.3 ± 1.0       | 41.2 ± 1.0      |
| *Pal*            | -                     | 36.7 ± 0.5     | 43.8 ± 0.4       | 49.0 ± 0.5      |
| *Pal*            | +                     | 40.0 ± 0.0     | 48.4 ± 1.4       | 52.0 ± 0.8      |
| *Ku*             | −                     | –                | –                   | –               |
| *Ku*             | +                     | –                | –                   | –               |

| *LanFTc1 Expression Was High and Thermoneutral in *Ku*, High and Partially Vernalization-Independent in *Pal*, Whereas Low and Positively Responsive to Vernalization in *Ku* |

83A:476 (*Ku*), Palestyna (*Pal*), and P27255 (*Ku*) grown under SD and LD conditions were used for gene expression profiling (Supplementary Data Sheet 1). The vernalization responsiveness of analyzed genes was calculated as a mean fold change of expression in vernalized plants compared to non-vernalized control (averaged across all day terms and dates measured for a particular line), as summarized in Table 4. The circadian clock responsiveness were calculated as a mean fold change of expression occurring between the morning and the evening terms (averaged across all dates measured for a particular line) and are provided in Table 5. The trend in expression level during plant growth was calculated as a fold change of expression occurring between the first and the last term (based on maximum daily values) and is provided in Table 6.

First, we analyzed the expression of the *LanFTc1* gene, which is considered as the major controller of vernalization responsiveness and early flowering in *L. angustifolius* (Nelson et al., 2017; Taylor et al., 2019). The studied genotypes showed different patterns of *LanFTc1* expression in response to photoperiod, vernalization, and circadian rhythm. Under SD, *LanFTc1* expression in non-vernalized plants was the highest in 83A:476 and the lowest in P27255 (Figure 2A). Indeed, *LanFTc1* expression in 83A:476 was up to 2.4 times higher than in Palestyna (*P* = 0.0016) and 460-2213 times higher than in P27255 (*P* = 0.0000). After vernalization, *LanFTc1* expression in 83A:476 was up to 4.5 times higher than in Palestyna (*P* = 0.0064) and up to 686 times higher than in P27255 (*P* = 0.0029). The difference of *LanFTc1* expression between...
83A:476 and P27255 increased during plant development in both vernalization variants, whereas for the pair 83A:476 and Palestyna it was decreasing. Under LD, LanFTc1 expression was the highest in Palestyna and the lowest in P27255 (Figure 2B). Namely, in non-vernalized plants, expression in Palestyna was 1.8-5.4 times higher than in 83A:476 (P = 0.0125) and 121-2642 times higher than in P27255 (P = 0.0085), whereas in vernalized plants it was 1.5-13.8 times higher in Palestyna than in 83A:476 (not significant, NS) and 24-764 times higher than in P27255 (P = 0.0094). The difference to Palestyna was increasing during plant growth for 83A:476 in both vernalization variants and P27255 in the absence of vernalization.

Vernalization influence on LanFTc1 expression in P27255 was manifested by up to a 7.2-fold increase under SD (NS) (Figure 2C) and up to 427-fold increase under LD (P = 0.0125) (Figure 2D). The vernalization effect under SD was changing in 83A:476 from a 0.5-fold decrease (P = 0.0115) to a 3.2-fold increase (P = 0.0094) and similarly in Palestyna from a 0.5-fold decrease (P = 0.0291) to 1.6-fold increase (NS). This effect under LD was neutral in 83A:476 and positive in Palestyna, up to 5.1-fold increase (P = 0.0351).

The circadian clock regulation differed between genotypes and partially between environments. In 83A:476, LanFTc1 expression was generally higher in the evening than in the morning in all combinations of the photoperiod and vernalization (up to 18.4-fold increase, P = 0.0008). In Palestyna, LanFTc1 expression under SD was usually higher in the morning (up to 2.9-fold increase, P = 0.0081), whereas under LD this effect was variable. In P27255, LanFTc1 expression was higher in the evening, especially under LD after vernalization (P = 0.0046).

### Table 4: Vernalization responsiveness of analyzed genes in L. angustifolius germplasm carrying Ku, Pal, and ku alleles, cultivated under 8-h (SD) and 16-h (LD) photoperiods.

| Gene name | Mean change of expression after vernalization under SD | Mean change of expression after vernalization under LD |
|-----------|--------------------------------------------------------|-----------------------------------------------------|
|           | Ku                      | Pal                      | ku                      | Ku                      | Pal                      | ku                      |
| LanAGL8   | 1.6^a                    | 2.2                      | 13.0                    | 1.8                      | 6.1                      | 138.6                   |
| LanCRLK1  | 0.7                      | 0.6                      | 2.6                    | 0.6                      | 1.3                      | 1.0                     |
| LanFD     | 0.5                      | 0.5                      | 3.6                    | 0.3                      | 0.7                      | 1.0                     |
| LanFTA1   | 1.1                      | 1.3                      | 0.9                    | 1.8                      | 2.1                      | 14.7                    |
| LanFTA2   | 1.1                      | 0.3                      | 13.3                   | 3.2                      | 1.1                      | 1.2                     |
| LanFTc1   | 1.3                      | 0.9                      | 4.3                    | 1.3                      | 3.6                      | 144.4                   |
| LanFTc2   | 0.7                      | 0.7                      | 3.5                    | 1.1                      | 3.2                      | 1.4                     |
| LanUGT85A2| 0.3                      | 0.2                      | 0.4                    | 0.2                      | 0.2                      | 0.4                     |
| LanVIN3-1 | 0.9                      | 0.9                      | 1.9                    | 0.8                      | 1.0                      | 1.1                     |
| LanVIN3-2 | 0.7                      | 1.1                      | 2.3                    | 0.7                      | 1.3                      | 1.6                     |
| LanVIN3-3 | 0.9                      | 1.7                      | 2.3                    | 0.6                      | 1.3                      | 1.0                     |
| LanVRN5   | 0.8                      | 1.0                      | 2.3                    | 0.9                      | 1.0                      | 0.8                     |

^a Fold change of expression occurring in response to vernalization, averaged across all data points (day terms and dates).

### Table 5: Circadian clock responsiveness of analyzed genes in L. angustifolius germplasm carrying Ku, Pal, and ku alleles, cultivated under 8-h (SD) and 16-h (LD) photoperiods without vernalization and with vernalization.

| Gene name | Mean change of expression during light phase under SD | Mean change of expression during light phase under LD |
|-----------|--------------------------------------------------------|-----------------------------------------------------|
|           | Ku                      | Pal                      | ku                      | Ku                      | Pal                      | ku                      |
| LanAGL8   | 1.6 | 1.0^a                        | 0.3 | 0.7        | 0.8 | 0.7        | 2.6 | 1.6        | 1.3 | 3.6        | 2.0 | 0.5        |
| LanCRLK1  | 0.9 | 1.0                  | 0.7 | 1.0        | 0.8 | 0.7        | 0.8 | 0.5        | 0.8 | 1.3        | 1.0 | 1.0        |
| LanFD     | 0.3 | 0.8                      | 1.7 | 3.1        | 1.2 | 1.6        | 0.3 | 0.4        | 1.6 | 2.4        | 1.9 | 1.3        |
| LanFTA1   | 1.0 | 1.7                      | 0.6 | 0.6        | 0.7 | 0.8        | 0.8 | 2.7        | 0.8 | 4.5        | 0.8 | 0.7        |
| LanFTA2   | 0.6 | 0.9                      | 0.9 | 0.4        | 0.4 | 0.4        | 0.4 | 2.2        | 0.7 | 0.4        | 0.6 | 0.6        |
| LanFTc1   | 4.8 | 3.1                      | 0.5 | 0.8        | 1.2 | 2.4        | 1.8 | 2.9        | 1.2 | 1.1        | 2.7 | 4.3        |
| LanFTc2   | 2.0 | 4.3                      | 2.0 | 11.1       | 5.6 | 24.3       | 2.7 | 2.7        | 2.0 | 0.5        | 3.4 | 4.2        |
| LanUGT85A2| 2.5 | 5.4                      | 2.6 | 2.1        | 2.4 | 2.4        | 3.2 | 4.6        | 3.8 | 2.2        | 3.3 | 2.9        |
| LanVIN3-1 | 1.2 | 1.5                      | 0.7 | 1.1        | 1.4 | 0.9        | 1.4 | 1.4        | 1.9 | 2.7        | 1.5 | 1.5        |
| LanVIN3-2 | 1.9 | 1.2                      | 1.0 | 1.2        | 1.3 | 0.9        | 1.7 | 1.4        | 1.5 | 2.4        | 1.4 | 1.4        |
| LanVIN3-3 | 1.4 | 1.3                      | 0.8 | 1.0        | 1.2 | 0.9        | 2.1 | 1.5        | 1.5 | 2.0        | 1.4 | 1.7        |
| LanVRN5   | 0.7 | 0.8                      | 0.7 | 0.7        | 1.3 | 0.9        | 1.3 | 0.9        | 1.1 | 0.9        | 1.0 | 0.9        |

^a Fold change of expression occurring between the morning and the evening terms, averaged across all dates; the first number corresponds to the experiment without vernalization, the second number to the experiment with vernalization.
**Table 6** Change of expression level of analyzed genes in *L. angustifolius* germplasm carrying *Ku*, *Pal*, and *ku* alleles, during plant growth under 8-h (SD) and 16-h (LD) photoperiods.

| Gene name | Change of expression during experiment under SD | Change of expression during experiment under LD |
|-----------|-----------------------------------------------|-----------------------------------------------|
|           | Ku                        | Pal                        | ku                          | Ku                        | Pal                        | ku                        |
| LanAGL8   | 1.5 | 2.3a | 3.5 | 6.3 | 0.4 | 6.4 |
| LanCRLK1  | 0.5 | 1.2 | 0.9 | 1.2 | 0.6 | 0.5 |
| LanFD     | 0.2 | 0.6 | 0.7 | 1.2 | 0.4 | 1.9 |
| LanFTA1   | 0.7 | 1.3 | 1.7 | 5.6 | 0.5 | 1.4 |
| LanFTA2   | 0.2 | 0.4 | 0.7 | 1.6 | 0.09 | 9.2 |
| LanFTc1   | 0.4 | 1.2 | 0.9 | 3.0 | 0.08 | 0.3 |
| LanFTc2   | 0.4 | 1.5 | 1.2 | 2.5 | 0.5 | -  |
| LanUGT85A2| 0.1 | 0.1 | 0.3 | 0.3 | 0.4 | 1.2 |
| LanWTN3-1 | 0.3 | 3.0 | 1.0 | 1.2 | 0.8 | 1.4 |
| LanWTN3-2 | 0.4 | 2.6 | 1.3 | 1.3 | 0.9 | 1.9 |
| LanWTN3-3 | 0.2 | 3.5 | 1.2 | 3.2 | 0.4 | 1.3 |
| LanVRN5   | 0.3 | 1.5 | 1.4 | 1.1 | 0.3 | 0.9 |

*aFold change in expression occurring between the first and the last term, based on the maximum daily values; the first number corresponds to the experiment without vernalization, the second number to the experiment with vernalization.

**LanFTA1 Expression Was Low and Uniform in All Lines Under SD, However, It Was Highly Induced Just Before Flowering, Especially in *Pal* and *ku* Genotypes Under LD**

Besides *LanFTc1*, three other *FT* genes, namely *LanFTA1*, *LanFTA2*, and *LanFTc2*, were analyzed to complement our perspective on *FT* clade transcriptional activity in *L. angustifolius* response to major environmental cues. *LanFTA1* gene expression under SD did not reveal any significant trend during plant growth and differences between genotypes were also usually not significant (Figure 3A). The relative level of expression was rather low (mean 0.38). A different pattern was observed under LD, when the expression was highly induced before flowering compared to the first term, namely 24-fold (*P* = 0.045) in non-vernalyzed 83A:476 (7-fold in vernalized, NS), 405-fold (*P* = 0.0051) in non-vernalized Palestyna (340-fold in vernalized, *P* = 0.0015), and 765-fold in vernalized P27255 (Figure 3B). In non-vernalyzed P27255 *LanFTA1* expression remained at a low level but this genotype did not flower in such conditions. The vernalization effect on *LanFTA1* expression under LD in 83A:476 and Palestyna was slightly positive, up to 2.5-fold (NS) and 2.8-fold (*P* = 0.0011) increase, respectively (Figures 3C,D). In P27255, the vernalization influence was initially moderately negative (0.14-fold and 0.35-fold decrease, *P* = 0.0002) but became highly positive just before flowering (43.6-fold increase). The circadian regulation was not stable across terms and genotypes, however, when *LanFTA1* was induced before flowering in Palestyna under LD, its expression was significantly higher in the evening than in the morning (up to 25.1-fold increase, *P* = 0.0004).

**LanFTA2 and LanFTc2 Expression Was Low in Both Photoperiods in All Lines**

*LanFTA2* gene expression was very low in both photoperiods, amounting to mean values of 0.095 under SD and 0.004 under LD (Supplementary Figures 1A,B). There was remarkable induction of expression in P27255 after vernalization under SD (up to 38-fold, *P* = 0.0002), however, the relative expression values achieved were much lower than those observed for *LanFTc1* and *LanFTA1* (Supplementary Figures 1C,D). This pattern was not recreated under LD.

***LanFTc2*** expression was on a low level in all lines under both photoperiods (Supplementary Figures 2A,B). A moderate induction by vernalization in P27255 under SD (up to 7.3-fold increase, *P* = 0.0187) and in Palestyna under LD (up to 4.7-fold increase, *P* = 0.0023) was observed but obtained levels were less than half of the mean expression obtained for control genes (Supplementary Figures 2C,D).

**LanAGL8 Expression Profile Reflected Observed Differences in Plant Phenology and Was Similar to LanFTc1**

Besides homologs constituting the *L. angustifolius FT* clade (Książkiewicz et al., 2016; Nelson et al., 2017), four novel candidate genes (*LanAGL8, LanFD, LanUGT85A2*, and *LanCRLK1*), recently considered to be putatively involved in *Ku*-based response (Plewiński et al., 2019), were profiled in this study. *LanAGL8* is an *L. angustifolius* homolog of the *A. thaliana* FRUITFULL gene participating in flowering time control, meristem identity, and fruit development (Mandel and Yanofsky, 1995; Gu et al., 1998). *LanAGL8* expression was consecutively increasing during plant growth for all studied combinations of lines, photoperiod, and vernalization variants except non-vernalyzed P27255 under SD. However, there were considerable differences in the observed expression levels between genotypes. Under SD without vernalization, expression of the *LanAGL8* gene in 83A:476 (Ku) was approximately 7–17 times higher (*P* = 0.0028) than in Palestyna (Pal) and 1420–6090 times higher (*P* = 0.0035) than in P27255 (ku) (Figure 4A). The influence of vernalization resulted in a reduction of differences in *LanAGL8* expression between 83A:476 and Palestyna by
FIGURE 2 | Gene expression profile of the LanFTc1 gene in response to photoperiod and vernalization in three lines (83A:476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku). (A) expression under an 8-h photoperiod, (B) expression under 16-h photoperiod, (C) vernalization response under 8-h photoperiod, (D) vernalization response under 16-h photoperiod. T1-T4 stands for sampling terms (Supplementary Table 5), V for vernalized plants, and N for non-vernalized plants. Timespan of photoperiods: 8-h from 4 AM to 8 PM, 16-h from 4 AM to 8 PM. Two references were used for normalization (LanDExH7 and LanTUB6) and one sample (LanTUB6) for inter-run calibration. Error bars indicate a standard deviation of 3 biological replicates, each representing a mean of 3 technical replicates. A logarithmic scale was used to accommodate observed large differences in gene expression values.

11–23%, and between 83A:476 and P27255 by 30–92%. Under LD without vernalization, LanAGL8 expression in 83A:476 was higher up to 12-fold ($P = 0.0086$) than in Palestyna and up to 1805-fold ($P = 0.0001$) than in P27255 (Figure 4B). The application of vernalization reduced these differences considerably. To summarize, Palestyna revealed an intermediate LanAGL8 expression profile, however, it was much more like 83A:476 than P27255.

The LanAGL8 gene revealed high positive responsiveness to vernalization in P27255 in both photoperiods, and this effect was consecutively increasing during plant growth until flowering. Namely, the change of LanAGL8 expression in P27255 after vernalization was raised from 2.1-fold (NS) to 29.8-fold ($P = 0.0175$) under SD (Figure 4C), and from 4.3-fold ($P = 0.0121$) to 398.2-fold ($P = 0.0004$) under LD (Figure 4D). Vernalization was also inductive in Palestyna, providing an up to 2.6-fold increase ($P = 0.0329$) of LanAGL8 expression under SD (effect stable across sampling terms) and up to 11.8-fold increase ($P = 0.0297$) under LD (effect consecutively decreasing). In the 83A:476 line, the influence of vernalization was the lowest, yielding up to a 2.9-fold increase of LanAGL8 expression under SD ($P = 0.0066$) and up to a 2.0-fold increase under LD ($P = 0.0315$).

The LanAGL8 gene revealed diversified circadian clock regulation between genotypes and environments. Under SD, its expression was higher in the morning than in the evening in Palestyna and P27255, whereas in 83A:476 this relation was the opposite. Moreover, vernalization partially diminished these differences. Under LD, LanAGL8 expression was usually higher in the evening terms for all lines, however, during the growth of vernalized P27255 this trend reversed.
**LanCRLK1 Expression Was Higher Under SD in All Lines and Negatively Responsive to Vernalization in Domesticated Ku Line**

*LanCRLK1* is an *L. angustifolius* homolog of the *A. thaliana* CRLK1 gene which participates in response to low temperature in a calcium-dependent manner (Yang et al., 2010). Changes in *LanCRLK1* expression level were usually not related to the progress of plant growth, except P27255 cultivated under SD, which revealed a stable decrease of *LanCRLK1* both in non-vernalized and vernalized variants. The expression of *LanCRLK1* was higher under SD than under LD. Moreover, under SD without vernalization, *LanCRLK1* expression was the highest in Palestyna (up to 2.7-fold increase compared to 83A:476, $P = 0.0002$) and the lowest in P27255 (about 0.4-fold decrease, $P = 0.0099$). However, after vernalization, expression was highest in P27255 (up to a 3.5-fold increase compared to 83A:476, $P = 0.0005$) and lowest in 83A:476 (Figure 5A). The differences in the *LanCRLK1* expression between genotypes under LD were generally not significant (Figure 5B).

The effect of vernalization on *LanCRLK1* expression under SD was negative (from 0.4-fold to 0.7-fold decrease, $P = 0.0009$) or neutral in 83A:476, negative in Palestyna (from 0.6-fold to 0.8-fold decrease, $P = 0.0000$), and positive in P27255 (up to 3.0 fold increase, $P = 0.0000$) (Figure 5C). Under LD, the vernalization effect was also negative in 83A:476 (0.6-fold decrease, NS), however, in the remaining two lines, the effect changed during plant growth from negative to positive: from a 0.7-fold decrease to 1.9-fold increase in Palestyna (all NS) and from 0.6-fold decrease ($P = 0.0001$) to 1.4-fold increase in P27255.

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**FIGURE 3** | Gene expression profile of the *LanFtA1* gene in response to photoperiod and vernalization in three lines (83A:476, Palestyna, and P27255) carrying different *LanFtc1* alleles (Ku, Pal, and ku). (A) expression under 8-h photoperiod, (B) expression under 16-h photoperiod, (C) vernalization response under 8-h photoperiod, (D) vernalization response under 16-h photoperiod. T1-T4 stands for sampling terms ([Supplementary Table 5](#)), V for vernalized plants, and N for non-vernalized plants. Timespan of photoperiods: 8-h from 4 AM to 8 PM, 16-h from 4 AM to 8 PM. Two references were used for normalization (*LanDExH7* and *LanTUB6*) and one sample (*LanTUB6*) for inter-run calibration. Error bars indicate a standard deviation of 3 biological replicates, each representing a mean of 3 technical replicates.
FIGURE 4 | Gene expression profile of the LanAGL8 gene in response to photoperiod and vernalization in three lines (83A:476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku). (A) expression under 8-h photoperiod, (B) expression under 16-h photoperiod, (C) vernalization response under 8-h photoperiod, (D) vernalization response under 16-h photoperiod. T1-T4 stands for sampling terms (Supplementary Table 5), V for vernalized plants, and N for non-vernalized plants. Timespan of photoperiods: 8-h from 4 AM to 8 PM, 16-h from 4 AM to 8 PM. Two references were used for normalization (LanDExH7 and LanTUB6) and one sample (LanTUB6) for inter-run calibration. Error bars indicate a standard deviation of 3 biological replicates, each representing a mean of 3 technical replicates. A logarithmic scale was used to accommodate observed large differences in gene expression values.

(P = 0.0107) (Figure 5D). The influence of the circadian clock on LanCRLK1 expression was variable in both photoperiods and usually not significant.

LanFD Expression Was Negatively Responsive to Vernalization in Ku and Pal Lines but Positively Responsive or Variable in ku

LanFD is an L. angustifolius homolog of the A. thaliana FD gene which triggers flowering based on FT-mediated signaling (Abe et al., 2005). LanFD expression in non-vernalized plants was usually the highest at the first sampling date and decreased during plant growth in both photoperiods, except in P27255 cultivated under LD. However, in vernalized plants, this decreasing trend was diminished or even reversed, as observed for P27255. Under SD, LanFD expression was the highest in Palestyna (1.1–5.6 times higher than in 83A:476, P = 0.0024) and the lowest in P27255 (0.3–0.9 times lower than in 83A:476, P = 0.0123) (Figure 6A). Interestingly, these relations were changed after vernalization as follows: LanFD expression in P27255 compared to 83A:476 was 3.2–12.2 times higher (P = 0.0011), whereas in Palestyna 2.1–7.4 times higher than in 83A:476 (P = 0.0001). Under LD without vernalization, differences in LanFD expression between genotypes were rather low, accounting for up to a 2.3-fold increase in Palestyna (P = 0.0205) and up to a 5.0-fold increase in P27255 (P = 0.0000) compared to 83A:476. However, vernalization exaggerated these contrasts and observed LanFD expression in P27255 was up to 25.7 times higher than in 83A:476 (P = 0.0078) and up to 5.5 times higher than in Palestyna (P = 0.0032) (Figure 6B).
The effect of vernalization on LanFD expression was negative in 83A:476 and Palestyna under both photoperiods (0.17-fold to 0.85-fold decrease), positive in P27255 under SD (up to 6.2-fold increase), and unstable in P27255 under LD (from 0.5-fold decrease to 1.4-fold increase) (Figures 6C,D). Genotypes differed in the circadian clock control of LanFD expression. Generally, in 83A:476 the levels were higher in the morning (up to 4.4 fold-increase), whereas in Palestine and P27255 in the evening (up to 4.5-fold and 3.1-fold increase, respectively).

LanUGT85A2 Expression Revealed a Growth-Dependent Decreasing Trend, Strong Circadian Clock Control, and Negative Response to Vernalization in All Lines Under Both Photoperiods

LanUGT85A2 is considered as an L. angustifolius homolog of the A. thaliana UDP-glycosyltransferase 85A2 gene. The expression of this gene was the highest at the first sampling date in all lines and consecutively decreased during the experiment. Palestyna was revealed to have the biggest decrease of LanUGT85A2 during the transition from juvenile to generative phase, whereas P27255 had the lowest. Indeed, this decrease of LanUGT85A2 expression during plant growth under SD reached approximately 0.1-fold for 83A:476 (P = 0.0114), 0.3-fold for Palestyna (P = 0.0033) and 0.4-fold for P27255 (P = 0.0003), whereas under LD about 0.3-fold for 83A:476 (P = 0.0501), 0.02-fold for Palestyna (P = 0.0182) and 0.5-fold for P27255 (P = 0.0196).

Under SD, the highest first-term LanUGT85A2 expression was observed in 83A:476, about 1.7-fold higher than in Palestyna (P = 0.0448) and P27255 (NS) (Figure 7A). These differences were doubled by vernalization. Contrary, under LD, the highest first-term expression was in P27255, 1.1-fold higher than in Palestyna (NS) and 3.7-fold higher than in 83A:476 (P = 0.0006), and these differences were tripled by vernalization (Figure 7B).
FIGURE 6 | Gene expression profile of the LanFD gene in response to photoperiod and vernalization in three lines (83A:476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku). (A) expression under 8-h photoperiod, (B) expression under 16-h photoperiod, (C) vernalization response under 8-h photoperiod, (D) vernalization response under 16-h photoperiod. T1-T4 stands for sampling terms (Supplementary Table 5), V for vernalized plants, and N for non-vernalized plants. Timespan of photoperiods: 8-h from 4 AM to 8 PM, 16-h from 4 AM to 8 PM. Two references were used for normalization (LanDExH7 and LanTUB6) and one sample (LanTUB6) for inter-run calibration. Error bars indicate a standard deviation of 3 biological replicates, each representing a mean of 3 technical replicates. A logarithmic scale was used to accommodate observed large differences in gene expression values.

Vernalization was observed to have a stable negative effect on LanUGT85A2 gene expression, highlighted by an average 0.29-fold decrease under SD (Figure 7C) and a 0.28-fold decrease under LD (Figure 7D). Response to vernalization was the weakest in P27255 (0.38-fold) and the strongest in Palestyna (0.21-fold). Such a disproportion in vernalization response, combined with the observed difference in decreasing trend during plant growth, resulted in the highest LanUGT85A2 expression in the P27255 line at the end of the experiment under both photoperiods (up to 31-fold increase compared to 83A:476 under LD, $P = 0.0233$).

The direction of circadian clock control was coherent across all genotypes and environments. LanUGT85A2 expression was higher in the evening than in the morning for all terms, both with and without vernalization. Under SD, it increased during the day up to 13.9-fold in 83A:476 ($P = 0.0165$), 8.3-fold in Palestyna ($P = 0.0075$) and 28.2-fold in P27255 ($P = 0.0000$). Under LD, it increased up to 7.4-fold in 83A:476 ($P = 0.0326$), 12.8-fold in Palestyna ($P = 0.0005$) and 4.4-fold in P27255 ($P = 0.002$).

**LanVIN3-1, LanVIN3-2, and LanVIN3-3 Genes Revealed High Expression in Ku Without Vernalization**

The *L. angustifolius* genome contains three homologs (named here as LanVIN3-1, LanVIN3-2, and LanVIN3-3) a VIN3 gene that is involved in the vernalization response in *A. thaliana* (Sung and Amasino, 2004). Therefore, the transcriptional activity of these genes was also profiled. In general, LanVIN3-1 revealed the highest expression, whereas LanVIN3-3 the lowest (see Supplementary Figures 3–5).

All three copies revealed differences in expression levels between genotypes, usually in descending order, 83A:476 – Palestyna – P27255. Maximum differences between 83A:476
and P27255 expression under LD reached 5.9-fold (LanVIN3-1, \( P = 0.0016 \)), 25.6-fold (LanVIN3-2, \( P = 0.0061 \)) and 8.4-fold (LanVIN3-3, \( P = 0.0059 \)), whereas under SD these values were as follows: 4.8-fold (LanVIN3-1, \( P = 0.0135 \)), 10.8-fold (LanVIN3-2, \( P = 0.0036 \)) and 5.5-fold (LanVIN3-3, \( P = 0.0072 \)). The expression levels of all three genes under LD were usually significantly higher in the evening than in the morning in all lines, whereas under SD, diurnal variations were frequently not significant or variable. Photoperiod conditions had a significant influence on expression changes during plant growth, especially for 83A:476. The expression levels of these genes in non-vernalized 83A:476 under LD increased by 2.3-fold (LanVIN3-1, \( P = 0.0008 \)), 3.3-fold (LanVIN3-2, \( P = 0.0048 \)) and 2.5-fold (LanVIN3-3, \( P = 0.0036 \)) whereas under SD they decreased by up to 0.32-fold (LanVIN3-1, \( P = 0.0000 \)), 0.37-fold (LanVIN3-2, \( P = 0.0110 \)), and 0.24-fold (LanVIN3-3, \( P = 0.0009 \)). The vernalization effect differed between genotypes and photoperiods. Under LD it was significant for 83A:476 (repression of all three genes, up to 0.43-fold) and Palestyna (induction of LanVIN3-2 and LanVIN3-3, up to 2.0-fold), whereas under SD it was significant for 83A:476 (changing from repression, up to 0.18-fold, to induction, up to 2.3-fold) and P27255 (induction, up to 3.6-fold). To summarize, all LanVIN3 homologs revealed different transcriptomic responses to vernalization than the LanFTc1 and LanAGL8 genes.

Besides VIN3, VRN5 protein is also considered to participate in the vernalization-induced epigenetic silencing in *A. thaliana* (Greb et al., 2007). Therefore, the expression of the latter gene was also profiled to supplement the analysis. All three *L. angustifolius* copies, LanVRN5-1, LanVRN5-2, and LanVRN5-3, were analyzed using one universal primer pair (Supplementary Figure 6). LanVRN5 clade revealed an expression profile relatively similar to LanVIN3, especially to LanVIN3-2. Like LanVIN3 genes, LanVRN5 in 83A:476 revealed an increase of expression during plant growth under LD and decrease under SD. All lines showed a significant decrease of expression during the light phase under SD and usually not a significant diurnal trend under LD. In

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**FIGURE 7** | Gene expression profile of the LanUGT85A2 gene in response to photoperiod and vernalization in three lines (83A:476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku). (A) expression under 8-h photoperiod, (B) expression under 16-h photoperiod, (C) vernalization response under 8-h photoperiod, (D) vernalization response under 16-h photoperiod. T1-T4 stands for sampling terms (Supplementary Table 5), V for vernalized plants, and N for non-vernalized plants. Timespan of photoperiods: 8-h from 4 AM to 8 PM, 16-h from 4 AM to 8 PM. Two references were used for normalization (LanDExH7 and LanTUB6) and one sample (LanTUB6) for inter-run calibration. Error bars indicate a standard deviation of 3 biological replicates, each representing a mean of 3 technical replicates.
both photoperiods without pre-sowing vernalization, 83A:476 revealed a significantly higher expression of LanVRN5 than Palestyna and P27255. However, the vernalization responsiveness of LanVRN5 considerably differed between photoperiods and genotypes. Under LD, only P27255 was responsive (showing repression), whereas under SD all lines responded but in different ways and with changing response during plant growth: namely from repression to induction in 83A:476, from induction to repression in Palestyna, and by continuously increasing induction in P27255. In vernalized plants under SD, LanVRN5 expression was the highest in P27255.

**DISCUSSION**

**INDEL Polymorphism in the LanFTc1 Promoter as a Legume Model for Functional Studies Aiming Vernalization Responsiveness**

*FT* is a major flowering time integrator gene, gathering signals from several pathways that detect environmental conditions (Turck et al., 2008). First studies on the conservation of *A. thaliana* genes in legumes have revealed the presence of the well-preserved homologs of many flowering time regulatory genes, including some representatives of *FT* clade found in *Pisum sativum* L., *Medicago truncatula* L., and *Glycine max* (L.), Merrill (Hecht et al., 2005). When the first gene-based linkage map of *L. angustifolius* was published, it highlighted a conserved collinear block shared between the fragment of *M. truncatula* chromosome 7 and the so-called linkage group LG01 (currently NLL-10), carrying major early flowering locus Ku (Nelson et al., 2006). It was later revealed that the syntenic region in *M. truncatula* contained several *FT* homologs (Young et al., 2011). The development of a bacterial artificial chromosome library for the *L. angustifolius* nuclear genome opened a possibility of gene cloning by DNA hybridization and the sequencing of selected clones (Kasprzak et al., 2006). Such an approach, combined with novel high-throughput sequencing techniques and gene expression profiling, resulted in the identification of a candidate gene for Ku and enabled the formulation of the hypothesis that a 1423 bp deletion in the promoter region of this gene is a causal mutation conferring early flowering phenotype (Nelson et al., 2017). Then, two other overlapping deletion variants, covering 1208 bp (*Pal*) and 5162 bp (*Jul*) were identified (Taylor et al., 2019). In the *A. thaliana*, a variation in promoter length was found to modulate the photoperiodic response of *FT*, and some important regulatory blocks contributing to this response were identified (Adrian et al., 2010; Liu et al., 2014). However, comparative mapping revealed a low sequence similarity between LanFTc1 and *A. thaliana* *FT* promoters, except RE-alpha and CCAAT boxes which were found at expected positions and indicated that *FT* promoter length in *L. angustifolius* may be at least as big as in *A. thaliana* (up to 7 kb) (Książkiewicz et al., 2016). All recognized LanFTc1 deletions conferring *Ku*, *Pal*, and *Jul* alleles are located downstream of the pair of CCAAT boxes marking a putative beginning of the functional promoter (Książkiewicz et al., 2016; Nelson et al., 2017; Taylor et al., 2019). Thus, *L. angustifolius Ku, Pal*, and *Jul* alleles have both distal and proximal regions preserved (Taylor et al., 2019). However, deletion sequences encompassed candidate binding sites for many transcription factors, including those already evidenced to be involved in *FT* regulation in model plants (Nelson et al., 2017; Taylor et al., 2019). In *A. thaliana*, similarly to *L. angustifolius*, the functional *FT* promoter indels also retained distal and proximal regions (Adrian et al., 2010; Liu et al., 2014). On the contrary, the study involving the *FT* promoter from cotton provided evidence that the proximal region might play an important role in this species (Sang et al., 2019). To our knowledge, a model revealed for *L. angustifolius* is the only known legume example of *FT* promoter indel variation. However, in many legumes, *FT* genes were found to be associated with flowering traits. In the economically most important legume crop worldwide, a soybean, *Glycine max*, mutations in *FT* genes are responsible for at least three loci conferring early/late flowering, namely *E9* (*GmFT2a*), *E10* (*GmFT4*), and *qDTF-J1* (*GmFT5a*) (Takeshima et al., 2016; Zhao et al., 2016; Samanfar et al., 2017). Moreover, natural variations of the *GmFT2b* sequence are associated with soybean adaption to high—latitude regions (Chen et al., 2020). In *M. truncatula*, vernalization responsiveness and early flowering are conferred by the *FTai* gene, whereas photoperiod response by the *FTb* gene (Laurie et al., 2011; Putterill et al., 2013). In pea (*Pisum sativum*), the *FTai* gene corresponds to the pea **GIGAS** locus, which is essential for flowering under LD and promoting flowering under SD (Hecht et al., 2011). In chickpea, a major quantitative trait locus (QTL) for the flowering time under SD conditions was mapped in the region containing a cluster of three *FT* genes (*FTa1-FTa2-FTc*), which collectively showed upregulated expression in domesticated germplasm (Ortega et al., 2019). In the sister lupin crop species, white lupin (*L. albus* L.), one of the four major QTLs conferring early flowering and partial vernalization independence was found associated with the *FTai* gene (Rychel et al., 2019). *L. angustifolius* genome contains two *FTa* and two *FTc* genes, which putatively arose from single copies by lineage-specific duplication, whereas the whole *FTb* subclade is absent (Książkiewicz et al., 2016). Indeed, *L. angustifolius* was recently used as a reference species in several phylogenetic studies addressing the influence of whole-genome and local duplications on the evolutionary fate of selected legume-specific and plant-wide gene clades (Przysecka et al., 2015; Narożna et al., 2017; Szczepaniak et al., 2018; Czyż et al., 2020). The differences in the expression profiles for *FTa* and *FTc* genes, as established in the present study, provided novel evidence supporting the hypothesis on a functional divergence of particular duplicates.

**Pal Allele Carrying Intermediate Phenology and Light Vernalization Responsiveness Provides Worldwide Opportunities for *L. angustifolius* Breeding**

In the present study, based on experiments performed in a greenhouse under natural LD (12–17 h photoperiod), *Ku* and *Jul* alleles were found to be associated with early
flowering and vernalization independence, *Pal* with slightly delayed flowering and partial vernalization responsiveness, and *ku* with late flowering and high vernalization dependence. The same phenomenon was reported for the recent study performed in phytotron under natural 10–12 h photoperiod (intermediate between SD and LD) in Australia (Taylor et al., 2019). However, we observed relatively high differences in flowering time for early lines (*Ku, Pal*, and *Jul*) in a greenhouse between 2014 and 2015 repeats, accounting for about two weeks on average (earlier in 2014). These differences can be attributed to the variation in temperature which occurred during these experiments. During the first 70 days of the 2014 experiment, we observed 31 days with maximum temperature ≥25°C and 9 days with ≥30°C, whereas in 2015 these numbers were much lower, 3 and 0 days, respectively. The higher temperature could advance flowering because it is shown to strongly accelerate flowering in model plants as well as in many other plant species (Parmesan and Yohe, 2003; Thines et al., 2014). Moreover, the average photoperiod in 2014 was about 2 h longer than in 2015 due to differences in sowing times. Indeed, our subsequent controlled-environment study revealed that even early *L. angustifolius* germplasm is responsive to LD conditions and accelerated transition between particular developmental phases by about 18–25 days compared to SD. The observed photophenology of the *Pal* allele can be very beneficial for *L. angustifolius* cultivation in the era of changing climate, especially in Europe and Australia where the majority of worldwide lupin production occurs. Thus, the European land climate experienced rapid warming in recent decades, resulting in the mean year temperature surge to approximately 2°C above the 1910–1960 average (NOAA, 2020). Climate warming raised multiple challenging issues for grain legume breeders, including higher water deficits and severe drought periods, propagation of pests and diseases as well as de-regulation of temperature-based control of growth and development processes (Vadez et al., 2012; Scheelbeek et al., 2018; Lippmann et al., 2019). Affected regulatory pathways include, among others, the flowering time control (Nelson et al., 2010). The rapid flowering of domesticated germplasm may favor drought escape and adaptation for spring sowing in higher latitudes (Annicchiarico et al., 2010, 2018; Berger et al., 2017). However, the observed extension of the vegetation period has raised the demand for germplasm with intermediate photophenology and cross-environment adaptation. Such research was recently initiated in *L. albus* in three European locations contrasting sowing time (autumn or spring) and climate type (Annicchiarico et al., 2019). Climatic variables were also addressed in an *L. angustifolius* genome-wide association study, providing some candidate polymorphisms that await further exploitation (Mousavi-Derazmahalleh et al., 2018a). *L. angustifolius* *Pal* allele confers flowering time and vernalization responsiveness phenotype intermediating between domesticated and wild lines. As this phenotype is consistent within the large range of photoperiod conditions (8, 10–12, and 16–17 h), it may be found applicable for all regions where lupins are currently cultivated.

**LanFTc1, LanAGL8, LanCRLK1, and LanUGT85A2 Are Candidate Genes Involved in the Vernalization Responsiveness of L. angustifolius**

Previous studies have highlighted the negative association of *LanFTc1* and *LanAGL8* gene expression with the number of days to flowering in *L. angustifolius*, mapping population and the positive direction of such association for *LanFD*, *LanCRLK1*, and *LanUGT85A2* genes (Nelson et al., 2017; Plewiński et al., 2019). The present study revealed that these genes differ in vernalization responsiveness between genotypes and photoperiods. *LanFTc1* and *LanAGL8* genes were found to be highly induced by vernalization in wild germplasm, whereas *LanUGT85A2* was found to be significantly suppressed (Table 4).

*LanAGL8* protein sequence revealed the highest similarity to *A. thaliana FRUITFULL* (*FUL*, AGAMOUS-LIKE 8, AT5G60910) and *APETALA1* (*API*, AGAMOUS-LIKE 7, AT1G69120) genes. Both *API* and *FUL* play a role in floral meristem identity but have different functions. *API* controls the formation of sepals and petals whereas *FUL* is involved in cauline leaf and fruit development (Irish and Sussex, 1990; Gu et al., 1998). These genes revealed tissue-specific expression during generative organ development (Irish and Sussex, 1990; Mandel and Yanofsky, 1995; Klepikova et al., 2016). In the present study, high levels of *LanAGL8* expression were revealed in leaf tissue. Moreover, the expression profiles and vernalization responsiveness of *LanAGL8* and *LanFTc1* were very similar. Both genes revealed comparable circadian clock control, i.e., morning induction in *Pal* line under short days. Confronting these observations with the information on a 100% association between the *LanFTc1* genotype and flowering time phenotype in a large germplasm collection (Nelson et al., 2017; Taylor et al., 2019), the conclusion can be raised that *LanAGL8* acts putatively downstream of *LanFTc1* in *L. angustifolius*, in which *LanAGL8* may perform a similar function, like its homolog in cereals, an *API*-like gene called VRN1, which regulates the transition from vegetative to generative phase in response to vernalization and is expressed in many organs, including leaves (Treviskis et al., 2003; Yan et al., 2003). Indeed, the wheat homolog of *FT*, (VRN3), activates expression of VRN1 in leaves and shoot apical meristem, promoting flowering under inductive long days (Li et al., 2015). As a MADS box transcription factor, VRN1 binds to many targets in the genome and may regulate many genes, linking vernalization and photoperiod pathways (Deng et al., 2015). Moreover, the allelic diversity of VRN1 copies provides wide plasticity of temperature-based responses in winter wheat (Dixon et al., 2019).

In this study, we also revealed differences in the vernalization responsiveness of a *LanFD* gene between early and late flowering germplasm. In wheat, FD—like, VRN3, and 14–3–3 proteins form together a florigen activation complex which can bind the VRN1 promoter, therefore a variation in *FD* expression may modulate the effect of mobile florigen signal (Li et al., 2015). Results obtained in this study, are supported by a significant correlation between the *LanFD* gene expression profile and vernalization responsiveness in the *L. angustifolius* mapping...
population (Plewiński et al., 2019), which indicates that *LanFD* may contribute to *FT* regulatory function, especially in the wild, vernalization-responsive germplasm.

Our recent expression quantitative trait loci (eQTL) mapping study provided transcriptomic evidence for the contribution of several genes acting in C-repeat binding factor (CBF) cold responsiveness (*LanCRLK1*), and in UDP-glycosyltransferases (*LanUGT85A2*) pathways in the vernalization response via *LanFT1* in *L. angustifolius* (Plewiński et al., 2019). *LanCRLK1* is a homolog of *A. thaliana* CALCIUM/CALMODULIN-REGULATED RECEPTOR-LIKE KINASE 1, which is the first component in the cold responsiveness pathway (Yang et al., 2010). Downstream genes in this pathway, the C-repeat binding factors (CBF) and INDUCER OF CBF EXPRESSION 1 (ICE1), provide regulatory links to FLOWERING LOCUS C (FLC) (Kim et al., 2004; Lee et al., 2015). The present study has highlighted the positive vernalization responsiveness of the *LanCRLK1* gene but only in wild germplasm under SD. In other genotype × environment combinations, the response was quasi thermoneutral. This finding is coherent with the general observation that CBF cold responsiveness pathway is downregulated and less effective under LD conditions than under SD (Lee and Thomashow, 2012). The expression profile of *LanCRLK1* did not provide convincing evidence on the contribution of this gene in the vernalization responsiveness of *L. angustifolius*. Nevertheless, the reduction of *LanCRLK1* expression in the evening, combined with the decreasing trend during development that was revealed in this study for vernalized *ku* line under LD, may explain the direction of the association between the *LanCRLK1* expression pattern and the vernalization responsiveness observed in the *L. angustifolius* mapping population e-QTL study, which was also performed under LD with partial vernalization (Plewiński et al., 2019). The question arises as to whether these differences in the *LanCRLK1* gene expression profile between the *Ku/Pal* and the *ku* lines, may have consequences in terms of cold acclimation and freezing tolerance of early flowering lines. A negative correlation between early phenology and cold acclimation could be a very undesirable trait hampering the autumn sowing of *L. angustifolius* in many regions of Southern Europe.

This study evidenced the negative response of *LanUGT85A2* to vernalization in all genotypes under both photoperiods. The genotypes explored in this study revealed different responses to photoperiod, and under SD, the *LanUGT85A2* expression was highest in the early flowering line, whereas under LD, it was in late flowering. Indeed, during the *L. angustifolius* mapping population e-QTL assay, which was performed under LD with mild vernalization, *LanUGT85A2* expression revealed a significant positive correlation with the late-flowering phenotype (Plewiński et al., 2019). *LanUGT85A2* is a representative of the UDP-glycosyltransferases protein family. In *A. thaliana*, a relatively close homolog of this gene, *UGT87A2*, promotes flowering in the vernalization and gibberellin pathways by repression of *FLC* (Wang B. et al., 2012). Similarly, ectopic over-expression in tobacco of a putative glycosyltransferase gene 1, *PtGT1*, derived from poplar (*Populus tomentosa* Carr.), resulted in an early flowering phenotype (Wang Y.-W. et al., 2012). Contrary, another *A. thaliana* homolog, *UGT84A2*, delays flowering by activation of the indole-3-butyric acid (IBA) pathway, leading to down-regulation of AUXIN RESPONSE FACTOR 6 (ARF6) and ARF8 genes, and, consequently *FT* (Zhang et al., 2017). Taking into consideration the direction of the association between *LanUGT85A2* expression and time to flowering, the latter mechanism seems to be more probable in *L. angustifolius* than those of *UGT87A2* and *PtGT1*.

This research highlighted the hypothetical involvement of *FLC*-related genes in *L. angustifolius* vernalization-dependent flowering time regulation. However, legume genomes, except for soybean, generally do not have *FLC* homologs (Hecht et al., 2005; Lyu et al., 2020). Nevertheless, other genes from the vernalization pathway, including some potential activators and repressors of *FLC*, are present (Hane et al., 2017). Interestingly, soybean has retained one *FLC* copy within its genome and does not require vernalization to initiate flowering. *FLC* in this species is involved in long-term low temperature-triggered late flowering by inhibiting *FT* gene expression (Lyu et al., 2020).

In *Arabidopsis*, vernalization-dependent silencing of the *FLC* gene is performed by VIN3 and VRN5 proteins which contribute to H3K27me3 and H3K9me2 enrichments during cold periods (Sung and Amasino, 2004; Greb et al., 2007; Kim and Sung, 2013). Despite their similar function in *A. thaliana*, VIN3 and VRN5 genes differ in expression profiles and vernalization responsiveness. Namely, VIN3 expression in *A. thaliana* is very low and highly induced during vernalization, but soon after the end of the cold period, it decreases again to pre-vernalization level; whereas the VRN5 gene is constitutively expressed at much higher levels than VIN3 with additional high upregulation after the end of vernalization (Kim and Sung, 2013). In our study, *LanVIN3* genes revealed significantly higher relative expression than *LanVRN5*. Apart from this difference, expression profiles of *LanVRN5* and *LanVIN3* homologs were similar to each other. Moreover, this research provided lines of evidence for the high expression of three *LanVIN3* homologs in early flowering 83A:476 lines without vernalization, highlighting the opposite effects of vernalization for particular genotypes. The observed differences in vernalization responses and expression profiles of *LanVIN3* and *LanVRN5* genes did not match observed differences in phenotypes (time to flowering). This may indicate that *LanVIN3* and *LanVRN5* genes are not involved in the vernalization response in *L. angustifolius*.

**DATA AVAILABILITY STATEMENT**

All datasets presented in this study are included in the article/Supplementary Material.

**AUTHOR CONTRIBUTIONS**

SR-B designed experiments and performed plant phenotyping for time to flowering and vernalization responsiveness, plant genotyping with molecular markers, and gene expression profiling by quantitative PCR. PP performed RNA isolation
and quantification. RG participated in the concept of the study and the interpretation of the results. BK contributed to data analysis and statistics. MK analyzed the data, interpreted the results, prepared the figures, and drafted the manuscript. All authors approved the final manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2020.572135/full#supplementary-material

Supplementary Table 1 | Accession numbers and information on domestication status and country of origin of L. angustifolius lines used in the study.

Supplementary Table 2 | Primers used in the study.

Supplementary Table 3 | Air temperature (daily mean and maximum) and daily sunshine hours recorded by the local/nearest meteorological station (Poznań-Lawica, 5.1 km) and theoretical photoperiod hours calculated for this location during the 2014 experiment.

Supplementary Table 4 | Air temperature (daily mean and maximum) and daily sunshine hours recorded by the local/nearest meteorological station (Poznań-Lawica, 5.1 km) and theoretical photoperiod hours calculated for this location during the 2015 experiment.

Supplementary Table 5 | Sampling terms selected for gene expression profiling.

Supplementary Table 6 | Leaf expression of selected flowering-related genes (reads per kilobase million, RPKM) derived from the transcriptome sequencing of L. angustifolius mapping population and correlation with vernalization responsiveness (ku).

Supplementary Table 7 | R2 and PCR efficiency values obtained for standard curves.

Supplementary Table 8 | Comparison of LanFTc1_INDEL2 marker scores and time to flowering in response to vernalization observed during ’2014 experiment.

Supplementary Table 9 | Comparison of LanFTc1_INDEL2 marker scores and time to flowering in response to vernalization observed during ’2015 experiment.

Supplementary Figure 1 | Gene expression profile of the LanFTa2 gene in response to photoperiod and vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

Supplementary Figure 2 | Gene expression profile of the LanFTc2 gene in response to photoperiod and vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

Supplementary Figure 3 | Gene expression profile of the LanVIN3-1 gene in response to photoperiod and vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

Supplementary Figure 4 | Gene expression profile of the LanVIN3-2 gene in response to photoperiod and vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

Supplementary Figure 5 | Gene expression profile of the LanVIN3-3 gene in response to photoperiod and vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

Supplementary Figure 6 | Gene expression profile of LanVIN5 genes in response to photoperiod and vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

Supplementary Data Sheet 1 | Gene expression data and calculations for LanAGL8, LanCALX1, LanFD, LanFTa1, LanFTa2, LanFTc1, LanFTc2, and LanFTTSA2 genes, obtained for two photoperiod variants (8- and 16-h) in response to vernalization in three lines (83A-476, Palestyna, and P27255) carrying different LanFTc1 alleles (Ku, Pal, and ku).

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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