**Prospective titles:**

Biological correlates and ecological drivers of seed oil content in alpine species

Unravelling seed oil content patterns in alpine species

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**Acknowledgements**

We are grateful to the Analysis Service Unit facilities of ICTAN for the seed oil analysis using Chromatography. We want to thank Giovanni Marchese and Francesco Cortese for their assistance in laboratory work. We also thank the Picos de Europa National for support in field sampling. This research is framed within the SEEDALP project (MCI-20-PID2019-108636GA-I00) funded by the Spanish Research Agency (AEI/10.13039/501100011033), and supported by Ayudas para Contratos Predoctorales para la Formación de Doctores del Ministerio de Ciencia e Innovación (FPI: MCI-21-PRE2020-092874); Ayuntamiento de Gijón/Xixón - Jardín Botánico Atlántico (SV-23-GIJON-JBA) and the grant “Laboratorio de Vegetación y Biodiversidad” (IDE/2024/0007020, Principality of Asturias-Sekuens-EU-FEDER).

**Funding**

Agencia Española de Investigación - Spanish Research Agency (AEI/10.13039/501100011033)

Ayuntamiento de Gijón/Xixón (SV-23-GIJON-JBA)

Grant “Laboratorio de Vegetación y Biodiversidad” (IDE/2024/0007020, Principality of Asturias-Sekuens-EU-FEDER).

**Data availability statement**

Raw data and R script for analysis are available in GitHub, once the manuscript is accepted Github repository will be public and stored in Zenodo.

**Author contributions**

Conceptualization: Clara Espinosa del Alba, Borja Jiménez-Alfaro; Investigation: Clara Espinosa del Alba, Silvia Baizan, Andrea Mondoni; Formal Analysis: Clara Espinosa del Alba, Andrea Mondoni; Writing – original draft: Clara Espinosa del Alba; Writing – Review & Editing: all authors; Supervision: Borja Jiménez-Alfaro, Eduardo Fernández Pascual, Andrea Mondoni; Funding acquisition: Borja Jiménez-Alfaro.

## Abstract (308 words)

Seed oil content, one of the crucial energy reserves in angiosperms, is important for multiple biological functions that impact plant fitness. Seed oil can be considered an adaptive trait with potential trade-offs involving other seed traits and displays latitudinal patterns as well as changes across life forms. Additionally, seed oil composition, namely the relative proportions of saturated and unsaturated fatty acids, has been proposed as a key trait for micro- and macro-evolutionary biogeographical patterns. However, the availability of oil data for wild species is minimal, and few studies have addressed the potential biological correlates and ecological drivers of seed oil content and composition.

Our study aims to explore these relationships with a unique data set of 47 European alpine species co-occurring in the Cantabrian Mountains (NW Spain) across altitudinal and microtopographical gradients. Seed oil content and composition analyses were conducted with fresh seeds collected in natural communities, successfully detecting 26 different fatty acids. We also measured seed traits with ecological and evolutionary relevance, such as seed mass, longevity, and germination timing. We calculated species' local ecological preferences along microenvironmental gradients to analyse ecological drivers based on 160 community plots sampled in the study area.

We found no relationships between seed oil content and oil composition with seed mass and germination timing, but we did find evidence that oil content significantly influences seed longevity. These results are essential for understanding the potential impacts of seed oil content on soil seed banks' persistence and adapting conservation strategies for seed banks and/or restoration programmes. Interestingly, and against expectations, we did not find patterns corroborating altitudinal or microtopographical gradients driving changes in seed oil content or composition. Alpine species showed strong phylogenetic constraints on seed oil content and composition. This study provides a meaningful advance to complete the seed ecological spectrum picture and a step forward to exploring seed trait functional ecology.

Keywords: Seed oil content; Seed oil composition; Germination; Longevity; Alpine species; altitudinal gradient; microenvironmental gradients.

## 1. Introduction (1542 words main text)

Seeds are energy reservoirs, mainly accumulating three macromolecules: proteins, lipids and carbohydrates (Levin, 1974; Baud and Lepiniec, 2010). The quantity and quality of these reserves are directly related to plant fitness, as they influence primary regeneration functions like seed persistence in the soil seed bank via seed longevity (Hoekstra, 2005), germination timing (Linder, 2000), and seedling establishment (Graham, 2008) having potential implications for the seed ecological spectrum (Saatkamp *et al.*, 2019). In angiosperms, the predominant reserve forms are lipids and carbohydrates, the firsts majorly in the form of triglycerides (an ester of glycerol plus three fatty acids) (Harwood, 1980; Voelker and Kinney, 2001), and the latter accumulated as starch (Levin, 1974). Carbons in fatty acids, through lipid oxidation, release more than twice the energy than the oxidation of starch on a per g basis of dry weight because their carbons are highly reduced (Levin, 1974; Baud and Lepiniec, 2010). Consequently, plants mainly rely on lipids (hereafter “seed oil”) for energy mobilization during germination (Harwood, 1980). In endospermic seeds, seed oil is mainly stored in the cotyledons and endosperm (Ellis, 2006) but also in the radicle and hypocotyl (Li *et al.*, 2006).

In seeds, most fatty acids range from 10 to 22 carbons in length, and the carbons may be joined by single or double bonds, referred to as saturated (SFA) and unsaturated fatty acids (UFA), respectively (Ellis, 2006). The relative proportion of specific fatty acids and the relative proportions of unsaturated and saturated fatty acids constitute what we will refer to as “oil composition”. The most abundant fatty acids found in seeds are unsaturated oleic acid (18:1n-9), linoleic acid (18:2n-6), alpha-linolenic acid (18:3n-3), and saturated palmitic (16:0) and stearic (18:0) acids (Voelker and Kinney, 2001; Ellis, 2006; Baud and Lepiniec, 2010). Unsaturated and saturated fatty acids possess distinct biochemical properties (Lehninger, Nelson and Cox, 1993), influencing their functioning as an energy reserve. On a per-carbon basis, unsaturated fatty acids are more expensive to produce and yield less energy when oxidised than saturated fatty acids (Lehninger, Nelson and Cox, 1993; Linder, 2000). Moreover, unsaturated fatty acids have a higher potential for oxidative damage (Rael *et al.*, 2004), requiring antioxidant molecules to prevent damage. Hence, an optimal energy storage strategy for seeds should maximise saturated storage instead of unsaturated fatty acids (Linder, 2000). Although seed oil content is highly constrained by phylogeny and subject to evolutionary change (Levin, 1974), variation in oil content and oil composition exists within and among genera of the same family (Levin, 1974; Voelker and Kinney, 2001; Bretagnolle *et al.*, 2016). Contrary to expectations, many species synthesise a low amount of saturated fatty acids (Linder, 2000). A potential explanation for this anti-intuitive pattern is that saturated fatty acids have a higher melting point than unsaturated fatty acids (Knothe and Dunn, 2009; Sanyal and Decocq, 2016), making energy less accessible at low temperatures. Additionally, the biochemical properties of unsaturated fatty acids increase the unpalatability, serving as a defence against herbivory, and also function as anti-freezing compounds (Linder, 2000); see Figure 1 for a summary table.

Seed oil content and composition influence a wide range of biological functions and, as such, are crucial for plant adaptation (Levin, 1974; Sanyal and Decocq, 2016), especially when regeneration stages come into play. It is known that seed energy and oil content are positively correlated (Levin, 1974; Bretagnolle *et al.*, 2016). Such a relationship suggests that oil synthesis is energetically costly and could be advantageous only for small seeds, which can store more energy in a smaller volume (Bretagnolle *et al.*, 2016). Large seeds generally store less oil than small seeds; however, a tiny oily seed can release as much energy as a larger, starchy seed (Bretagnolle *et al.*, 2016). There is a consensus in the literature that seed mass and oil content are negatively correlated, with seed oil content contributing up to 60% of seed mass weight (Ellis, 2006; Theodoulou and Eastmond, 2012). In concordance with the correlations reported, and since seed mass influences among other things, dispersal biology (Westoby, Leishman and Lord, 1996) and germination rate (Bu *et al.*, 2007), there could be indirect correlations between oil content and other seed traits and functions. Thus, selection could act simultaneously on multiple seed traits (Sanyal and Decocq, 2016). Previous studies have claimed that seed oil content influences seed longevity (Hoekstra, 2005), meaning the ability of seeds to remain viable over specific storage periods, which partially determines the capacity of seeds to persist in the soil seed bank. Secondary lipid oxidation metabolites, needed for energy release, can cause detrimental effects on other macromolecules (Graham, 2008). Remarkably, the oxidation of unsaturated fatty acids contributes significantly to producing free radicals and subsequent attacks by these radicals (Harman and Mattick, 1976). To prevent deleterious effects, oily seeds also store antioxidants, and studies show a positive correlation between antioxidant levels and the relative proportion of unsaturated fatty acids (Sattler *et al.*, 2004). Accordingly, previous studies have found oily seeds to be more prone to ageing (Nagel and Börner, 2010; Neto *et al.*, 2019), and detailed studies, including oil composition, have also shown that a higher proportion of unsaturated fatty acids is associated with shorter longevity (Hoekstra, 2005). Thus, not only oil content but also oil composition affects the storage behaviour of seeds (Volk *et al.*, 2006). Further investigations on the effects of seed oil content and composition on longevity are desirable (Pritchard and Dickie, 2003) but are lacking for most wild species. Another key process in regeneration is germination, a complex and irreversible process which requires the use of the energy stored in the seed (Nonogaki, Bassel and Bewley, 2010). It is reasonable to think that the greater the energy reserve, i.e. oil content, the higher and faster germination (Gardarin, Dürr and Colbach, 2011), especially in conditions of higher competition for light (Levin, 1974). Additionally, it has also been reported that due to the different fatty acid melting points, under colder temperatures, seeds with a higher proportion of unsaturated fatty acids can germinate earlier (Linder, 2000).

In a pioneering study on wild species, Levin (1974) found significant seed oil content variations depending on habitat type (oil increased in woodlands) and life form (lower oil in herbs than in shrubs and trees) but no geographical differences, except within herbs where subtropical herbs showed higher oil values than temperate and tropical herbs. However, a more recent study found a general latitudinal gradient, with higher seed oil content in tropical plants compared to temperate plants (Sanyal and Decocq, 2016). In the tropics, seeds need higher energy to germinate fast and survive the first life stages with high competition for light (Levin, 1974). In the same study, they also found significant oil composition gradients, with saturated fatty acids decreasing with increasing latitude (Sanyal and Decocq, 2016), in concordance with previous studies (Linder, 2000). Higher growing temperatures have been correlated with higher oil content in several cultivars (Piper and Boote, 1999) and higher saturated fatty acid accumulation (Sanyal and Linder, 2013). Additionally, the different fatty acids' melting points can give a complementary explanation. At higher latitudes (cooler temperatures), seeds with higher unsaturated fatty acids proportions, which have a lower melting point, can access the energy stored and germinate earlier, providing a competitive advantage (Linder, 2000). At lower latitudes (higher temperatures), seeds do not have a temperature limitation to access higher saturated fatty acids energy resources with higher melting points (Sanyal and Decocq, 2016). Consequently, seed oil content and composition seem to be under strong selection by temperature at both micro- and macro-ecological levels (Sanyal and Linder, 2013; Sanyal and Decocq, 2016).

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Figure 1. A) Fatty acid biochemical properties. B) Altitudinal oil content and unsaturated UFA/SFA ratio (Unsaturated Fatty Acids /Saturated Fatty Acids) patterns expected in our study.

In this study, we investigate biological correlates and ecological drivers of seed oil content and composition in a regional alpine flora. Alpine ecosystems are unique laboratories to explore seed oils because they are subjected to strong ecological filters towards similar life forms (Körner, 2021b), but at the same time, support high evolutionary diversity (Rahbek *et al.*, 2019) and functional variation across microtopographical gradients (Scherrer and Körner, 2011). Also, alpine plants have adapted to multiple reproductive strategies (through different flowering, seed and clonal traits); they may have responded to elevation and microclimatic snow gradients with a specific combination of seed oil content and composition. Temperate alpine species are generally known for having average-sized seeds (Körner, 2021b), short longevity (Mondoni *et al.*, 2011), and delayed germination phenology due to physiological dormancy (Fernández-Pascual *et al.*, 2021; Espinosa del Alba, Fernández-Pascual and Jiménez-Alfaro, 2024). Our first objective is characterising seed oil content and seed oil composition in the alpine temperate flora. Second, we test the existence of a regional altitudinal seed oil pattern potentially caused by the differential temperatures experienced. Third, we explore potential correlations between seed oil content and composition with three seed traits related to important regeneration functions: (1) seed mass, (2) seed longevity (p50), calculated as the time for viability to drop below 50% and (3) germination timing (EHS), meaning how many environmental degrees seeds need to experience to reach 50% germination, early germination is indicated by lower accumulated degrees. Finally, we test potential ecological drivers of seed oil content and composition at a local microtopographical scale, measuring three different bioclimatic indices to quantify warmth (Growing Degree Days), freezing (Freezing Degree Days), and snow cover (snow days) species preferences, see Table 1 for detailed expectations.

Table 1. Study expectations relationships between seed oil content and composition with biological correlates and ecological drivers at a microtopographical scale.

|  |  |  |  |
| --- | --- | --- | --- |
|  | | Oil content | Ratio UFA/SFA |
| Biological correlates | | | |
|  | Seed mass | Decreasing seed mass with increasing oil content | Decreasing seed mass with increasing ratio |
| Seed longevity (p50) | Decreasing p50 with increasing oil content | Decreasing p50 with increasing ratio |
| Germination timing (EHS) | Decreasing EHS with increasing oil content | Decreasing EHS with increasing ratio |
| Ecological drivers | | | |
|  | Growing Degree Days (GDD) | Increasing oil content with increasing GDD | Decreasing ratio with increasing GDD |
| Freezing Degree Days (FDD) | Decreasing oil content with increasing FDD | Increasing ratio with increasing FDD |
| Snow Days (Snow) | Decreasing oil content with increasing snow | Increasing ratio with increasing FDD |

## 2. Materials and Methods (1717 words main text)

### Study system

We focused on herbaceous alpine species from grassland communities in the Cantabrian Mountains (north-western Spain), a transitional mountain system between southern Europe's temperate and Mediterranean regions (García-Gutiérrez *et al.*, 2018). These grassland communities are continuously distributed along the mountain range, occupying reduced areas above the treeline and around mountaintops between 1580 and 2600 m a.s.l (González Le Barbier et al., 2025). Grassland communities are predominantly dominated by Poaceae and Cyperaceae, and the main life forms are hemicryptophytes and dwarf chamaephytes. Climatic conditions follow a north-south temperature and precipitation gradient, with colder and wetter conditions on northern slopes compared to warmer and drier conditions on southern slopes. Mean annual temperatures in the study area range from 2.5 to 4.5 ºC, and mean summer precipitation ranges from 160 to 260mm (values extracted from Chelsa 2.1 bio1 and bio17, Karger *et al.*, 2017). We established eight sampling sites in the alpine grassland zone, four on the northern slopes and four on the southern slopes of the Cantabrian Mountains, ensuring representation of the flora from the two biogeographical regions. We collected floristic and community composition data for all vascular plants at each sampling site and classified species as strict alpine or generalist according to their altitudinal distribution in the study region. We identified plant specialists as those significantly associated with alpine grasslands, using the Indicator Values (IndVal) in the indicspecies R package (De Cáceres and Legendre, 2009). The calculations were based on 12,000 vegetation plots of grasslands stored in the SIVIM database (www.sivim.org) for the ecoregion in which our system occurs (The WWF Cantabrian Mixed Forest ecoregion). Seed collection permits were granted for the sampling sites inside Picos de Europa National Park: expedient numbers CO/09/203/2021 and CO/09/192/2023.

### Seed oil and seed mass

In a preliminary exploratory analysis, we tested a destructive method to calculate seed oil content and composition for three seed lots from five species and different taxonomic families. Since we did not find significant differences within the seed lots, we focused our seed collections on a single sample per species. Out of the 119 initially recorded species, we collected enough material for 47 species (19 plant families), a minimum of 200 mg of dry seeds per sample. Oil analyses were performed at the USTA-CSIC laboratory using a gas chromatographer with a Flame Ionization Detector (Agilent 7820A, EZChrom Elite software). Samples were manually ground with liquid nitrogen; then, fatty acids were transformed into fatty acid methyl esters for detection following the procedure described by Lee et al. (2012) section 2.2.4, potentially detecting up to 45 different fatty acids (supplementary Table S1). As a result, we obtained the absolute value of the oil content on a dry-weight basis and the percentage of each fatty acids type relative to the total oil content. The analysis identified 26 different fatty acid types in the studied species (see supplementary Table S2). This data set was used to conduct the local alpine species analysis of the variation in seed oil content and composition, together with seed mass data measured the 47 species, averaging the weight of five replicates of 50 dry seeds.

To contextualize our study at the regional level and for comparison with non-alpine species, we retrieved seed oil and seed mass information for another 33 species native to Europe, congeneric with the species we collected from the Seed Information Database (SID) (SER, INSR and RBGK, 2023) and classified as either generalist or strict lowland. In addition to our own seed oil data, we used these data to study regional altitudinal gradients, obtaining oil content and seed mass data for 80 species (29 strict alpines, 31 generalists and 20 strict lowlands) from 19 plant families and 12 orders.

### Seed germination and longevity

To estimate the germination timing, we used the Environmental Heat Sum trait (EHS), calculated as the amount of degrees Celsius (°C) accumulated for each species until reaching 50% germination, obtained from a previous phenology germination experiment conducted in the same system (Espinosa del Alba, Fernández-Pascual and Jiménez-Alfaro, 2024) and available for 34 of our study species. Seed longevity was experimentally calculated for 35 species in the laboratory using a standard comparative longevity protocol (Probert, Daws and Hay, 2009; Davies *et al.*, 2016) optimised for short-lived species (Davies *et al.*, 2016) and successfully applied to alpine species by Mondoni et *al.*, (2011). The artificially accelerated ageing protocol allows the measurement of the p50 value (amount of time for seed viability to drop to 50%). The initial RH value was 30-35% (Hygropalm 3 display unit; Rotronic Instrument UK Ltd, Crawley, UK). Before the start of the ageing protocol, all species were tested to ensure germination above 85% (see specific germination conditions below) and species from Cistaceae were scarified with sandpaper to break physical dormancy. Seed samples (200 seeds/ species) were first rehydrated to 47% RH at 20ºC for five days in a non-saturated LiCl solution (Hay *et al.*, 2008) in crystal Petri dishes and kept in a 300 x 300 x 130 mm sealed electric enclosure box (Ensto UK Ltd, Southampton, UK) before moving the Petri dishes into the ageing conditions. The ageing conditions comprised a temperature of 45°C, 60% RH and darkness. Subsamples of 42 seeds were withdrawn after 2, 10, 15 and 30 days in ageing conditions and consequently sowed in Petri dishes 1% agar with GA3 following Kew standard procedure (Davies, Sacco and Newton, 2015). After the ageing period, those species requiring a cold stratification period (e.g., *Saxifraga oppositifolia*) were stored at 5ºC in darkness for a month before germination experiments. After ageing, germination conditions for the species were set with a 12/12 h photoperiod and two alternating temperatures: warmer conditions at 22-12ºC or 15/5ºC for those germinations which required colder temperatures (see supplementary Table S3). After sowing, seeds were checked once a week for four weeks. Germination was scored when there was a visible root (>1.5 mm) and then removed from the Petri dishes. After the 28-day germination test ended, the ungerminated seeds were cut-tested under the binocular stereoscope to visually assess the embryo's state. We considered seeds with white and firm embryos viable, i.e. potentially germinable (Baskin and Baskin, 2014) and removed empty or infected seeds from further analysis.

Probit analysis was carried out on the data using GenStat Release 11.1 (VSN International Ltd, Oxford, UK) to estimate the time for viability to fall to 50% (p50) by fitting the viability equation (Ellis and Roberts, 1980):

*v = Ki – (p/σ)*

where v is the viability (in normal equivalent deviates, NED) of the seed lot after p days in storage, Ki is the initial viability (NED) of the seed lot, and *σ* is the time (d) for viability to fall by 1 NED (i.e. the standard deviation of the normal distribution of seed deaths over time).

For the complete seed oil and seed traits dataset, analyses and measurements were done in two or three batches, corresponding with the seed availability after each year's collection campaign. Nevertheless, we are confident that we have collected fully mature seeds, following the ENSCONET protocol (2009), and have applied a consistent methodology for all experiments and analyses.

### Species preferences along local micro topographical gradients

To calculate species preferences, we established 20 additional plots (1 m²) per site (following the methodology of Jiménez-Alfaro *et al.*, 2024) where we recorded all vascular species and buried a temperature datalogger at a depth of 5 cm (Thermochron, iButton, Newbury, UK; accuracy: ± 0.5 °C from -10 °C to +65 °C, resolution: 0.5 °C, records every four hours), which recorded temperatures over 11 months. From the microenvironmental data recorded, we calculated three bioclimatic indices per plot following Jiménez-Alfaro *et al.*, (2024): (1) Growing Degree Days (GDD) as the sum of daily mean temperatures for days in which the soil mean temperature at five cm deep was above 5 ºC (Körner, 2021); (2) Freezing Degree Days (FDD) as the sum of daily mean temperatures for days in which the mean temperature was below 0 ºC (Choler, 2018); and (3) days with snow cover, based on the period in which the maximum temperatures were < 0.5 ºC and the minimum temperatures were > -0.5 ºC (Zhang *et al.*, 2005). Then, we used the bioclimatic indices to calculate the species' local preferences, averaging the climatic variables of the plots where the species was present and weighting them by their coverage (only considering those plots where the species had more than 10% relative coverage), assuming species would have more coverage in those plots with climatic conditions closer to their local ecological optimum.

### Statistical analysis

We analyzed the seed trait data by fitting Markov Chain Monte Carlo generalized linear mixed models (MCMCglmm) with Bayesian estimation using the R package MCMCGLMM (Hadfield, 2010). For biological correlates, we investigated how seed oil content and composition may explain other seed biological traits (seed mass, longevity and germination timing). Thus, oil traits are used as the explanatory variable. We applied a log transformation to seed mass and EHS to meet the normal distribution of the data, and model assumptions were visually checked and fulfilled. For the ecological drivers (altitudinal distribution and local ecological gradients, i.e. GDD, FDD and Snow), we wanted to investigate how seed oil and composition are regulated by microclimatic gradients (across local snow-related factors). Thus, oil traits are used as the response variable, both log-transformed. We used weakly informative priors in all models, with parameter-expanded priors for the random effects. Each model was run for 1,000,000 iterations, with an initial burn of 100,000 and a thinning interval of 100. We calculated mean parameter estimates and 95% credible intervals (CI) from the resulting posterior distributions. Phylogeny was included as a random factor using a reconstructed tree (Supporting information Figure S1 and Figure S2 for regional and alpine species pool, respectively), created with V.PHYLOMAKER R package (Jin and Qian, 2019) using phylosignal and phylobase R packages (Keck *et al.*, 2016; R Hackaton et *al*., 2024). To work with the phylogeny and calculate the phylogenetic signal (lambda), we used the ape (Paradis and Schliep, 2019) and phylosignal packages (Keck *et al.*, 2016). To assess the significance of the models, we use the posterior mean from posterior probabilities and the credible intervals (CI). The posterior mean is an analogue of model effect sizes, and when the CI values do not overlap with 0, there is a significant effect of the explanatory variable.

## 3.Results (907 words main text)

### Seed oil content and oil composition

Within our local alpine species pool (n=47), oil content varies from 1.2% in *Avenella flexuosa* (Poaceae) to 34.2% in *Jasione cavanillesii* (Campanulaceae) (Figure 2A). The most abundant fatty acids in alpine seeds were the unsaturated linoleic acid (C18:2n-6c, 42.4%±2.4, mean ± se), oleic acid (C18:1n-9c, 22.6%±1.9), and alpha-linolenic acid (C18:3n3, 15.1%±2.7); as well as saturated palmitic (C16:0, 10.8%±0.8) (details in supplementary Table S2). These four averages 94.8% of seed oil content (se=23). Additionally, in our data set, erucic acid (C22:1n9) had high values only in Brassicaceae and gamma-linolenic acid (C18:3n6) in some Caryophyllaceae (Figure 2B). The mean frequency of saturated fatty acids is 14.2%±0.95, and the mean ratio between unsaturated and saturated fatty acids is 7.2 (se=0.5), ranging from 1.3 in *A. flexuosa* up to 20.1 in *Iberis carnosa* (Brassicaceae). Poaceae, Salicaceae, and Cistaceae generally had higher proportion of saturated fatty acids synthesis while Brassicaceae and Lamiaceae were characterized by higher unsaturated fatty acids proportion (Figure 2C).

We conducted a Principal Component Analysis using fatty acid values with >3% relative proportion and correlations below 0.7 to reduce dimensionality (Figures 3D and 3E). The UFA/SFA ratio contributed the most to PC1 (26.4%) followed by palmitic acid (C16:0, 18.6%), while eicosenoic acid (C20:1n9) and erucic acid (C22:1n9) mainly contributed to PC2 (21.9% and 16.3%, respectively). Additionally, PCA revealed that the oil content percentage is not highly correlated with any specific fatty acid (details in supplementary Table S4).

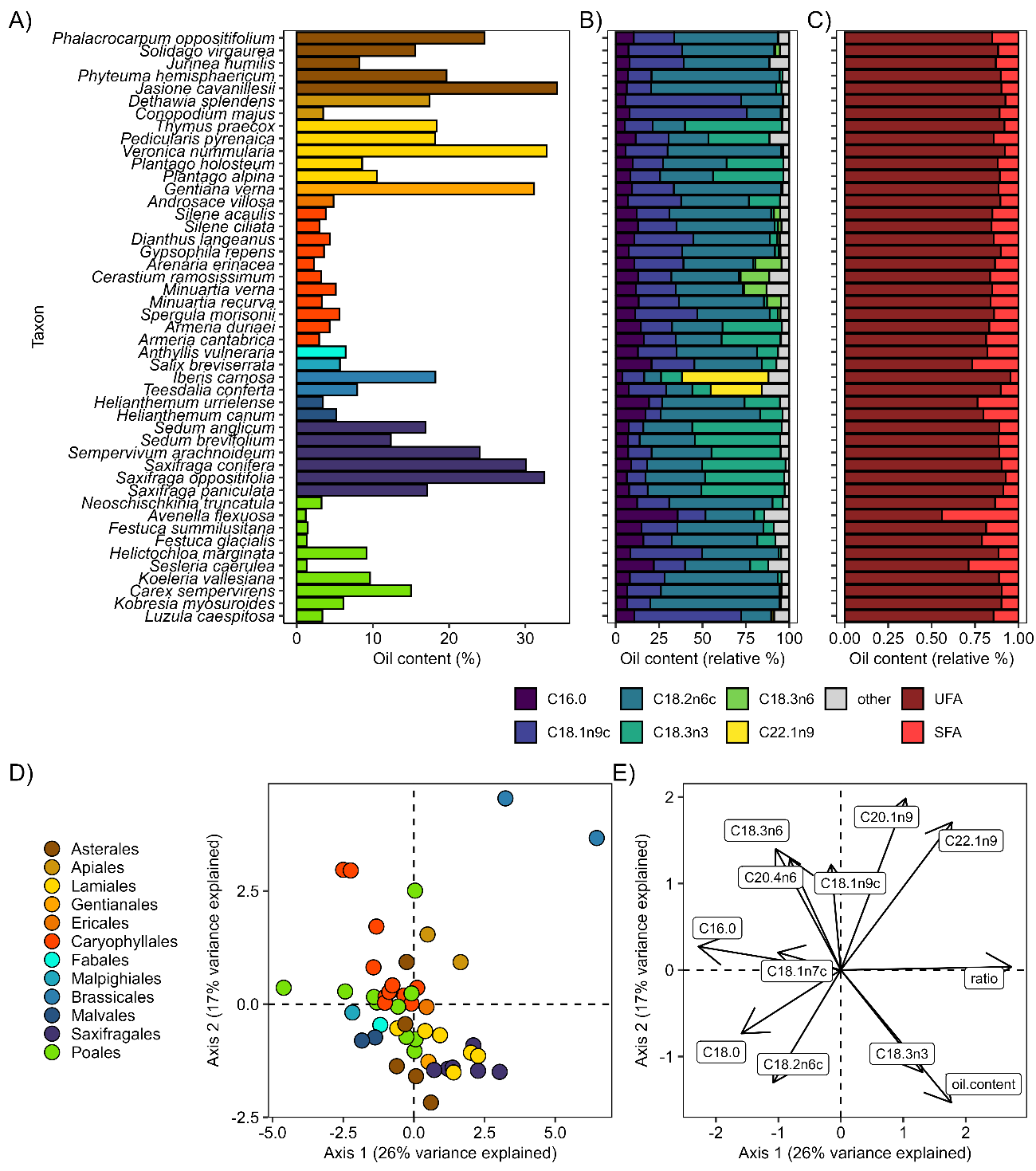


Figure 2. Local species oil content and composition data exploration (n=47). A) Oil content (%) per species, colours representing orders as panel D. B) Seed oil composition of fatty acids with more than 10% relative proportion on average across species. C) Seed oil composition per species with fatty acids divided between Unsaturated Fatty Acids (UFA) and Saturated Fatty Acids (SFA). D) Exploratory PCA with species as points and E) variables directions and contributions.

In our regional dataset with 80 species analysed (Figure 3A), we found no significant oil content differences between strict alpine, generalist and strict lowland species (model details in supplementary Table S5), although the relationship was significantly modulated by phylogeny (posterior mean= 0.86, CI: [0.74|0.97]), random factors model details in supplementary Table S6. Similarly to the patterns found in the local alpine species, oil content values ranged from 1.19% to 38%, with a mean value of 11.5%. Additionally, we found no significant correlation between seed mass (mg) and oil content (%) (posterior mean=-0.02, CI: [-0.06|0.01]); again, the relationship between both variables was significantly modulated by phylogeny (posterior mean=0.73, CI: [0.5|0.93]). We observed a considerable variation of seed oil content in smaller seeds, while variation was reduced with increasing seed mass (Figure 3B). Seed mass values ranged from 0.64 mg to 335 mg, with a mean value of 51 mg. When we calculated the phylogenetic signal of both traits, lambda values were 0.77 and 0.21 for seed oil content and seed mass, respectively, and significant only in seed oil content (p-value = 0.001).

Gráfico, Gráfico de dispersión

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Figure 3. Regional oil content patterns (n=80 species). A) Seed oil content (%) between regional altitudinal distribution showed by different colours. B) Seed oil content (%) and seed mass (mg) non-significant relationship (from MCMC-GLMMs models with gaussian family) showed by posterior mean (post. mean) and credible intervals (CI); if CI values do not overlap with 0, they indicate a significant relationship

### Biological correlates

Seed mass values ranged from 0.6 mg (*Sedum brevifolium*) to 268 mg (*Jurinea humilis*), with a mean of 45.4 mg (se=7.84). Although we observed a general trend of decreasing seed mass with a higher proportion of seed oil content, the relationship was not significant (posterior mean=-0.02, CI: [-0.06|0.02], Figure 4A, left panel). Seed oil composition, calculated as the UFA/SFA ratio, also did not significantly correlate with seed mass (posterior mean=-0.01, CI: [-0.11|0.09], Figure 4A right panel). However, in both cases, the relationship between variables was significantly modulated by phylogeny (posterior mean=0.94, CI: [0.84|0.99] for seed mass – oil content and posterior mean=0.95, CI: [0.87|0.99] for seed mass – oil composition). Additionally, all three seed traits had a high lambda for the local alpine species pool: 0.8 for seed mass, 0.71 for seed oil content and 0.62 for the UFA/SFA ratio. The p50 values ranged from 3 to 47 days, with a mean of 21 days (se=2). Higher oil content significantly reduced seed longevity (posterior mean=-0.72, CI: [-1.12|-0.27], Figure 4B left panel). The relationship with the ratio was less noticeable; seeds with a higher UFA/SFA ratio (i.e. more unsaturated fatty acids) showed a trend of decrease in longevity, but it was not significant (posterior mean=-0.96, CI: [-2.52|0.79], Figure 4B right panel). Interestingly, here, phylogeny does not appear to have a significant effect either in the trait p50 (lambda = 0.0006) or in the relationship between variables (posterior mean=0.1, CI: [0|0.41] for p50 – oil content and posterior mean=0.23, CI: [0|0.82] for p50 – oil composition). Germination timing, estimated as EHS, ranged from 63.2 to 1245 degrees Celsius, with a mean of 430°C (se=48.9). We observed no clear trends between oil content or UFA/SFA ratio and germination timing and the relationships were not significant (for oil content posterior mean=0.02, CI: [-0.01|0.05] and for UFA/SFA ratio posterior mean=-0.04, CI: [-0.03|0.14], Figure 4C). Here again, we observed a lack of phylogeny constraints in the relationship between the variables (posterior mean=0.12, CI: [0|0.41] for seed mass – oil content and posterior mean=0.11, CI: [0|0.4]) or in the trait EHS (lambda= 0.0007).

Gráfico, Gráfico de dispersión

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Figure 4. Seed oil content and ratio UFA/SFA biological correlates, significances from MCMC-GLMM models (Gaussian family) showed by posterior mean (post. mean) and credible intervals (CI); if CI values do not overlap with 0, they indicate a significant relationship. A) Correlation between seed mass, oil content, and UFA/SFA ratio. B) Correlation between seed longevity (P50, from GENSTAT probit analysis), oil content, and UFA/SFA ratio. C) Correlation between germination timing (EHS, from Espinosa del Alba, Fernández-Pascual and Jiménez-Alfaro, (2024), oil content and UFA/SFA ratio. Colours represent the different plant orders.

### Local ecological drivers

GDD values ranged from 650 to 2295 °C, averaging 1421 °C (se = 67.9), while FDD values ranged from 0.05 to 170 °C, with a mean of 31.8 °C (se = 4.7). We found a general trend of species with lower oil content and a lower UFA/SFA ratio placed in warmer conditions (high GDD and low FDD) along local microclimatic gradients (Figure 5), although these effects were not significant. The trends were more consistent for GDD, since the effect of FDD was mainly driven by a few species with more extreme oil content and composition values. In addition, snow values ranged from 3 to 157 days, with an average of 62 days of snow per year. The data showed a trend where species that preferred snowier sites showed higher oil content and UFA/SFA ratios but with no significant effects. Details of the model results are shown in supplementary Table S4.

Gráfico, Gráfico de dispersión

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Figure 5. Seed oil content ecological trade-offs, oil content and ratio UFA/SFA log-transformed for MCMC-GLMM models (family=Gaussian, n=46), significances showed by posterior mean and credible intervals (CI); if CI values does not overlap with 0 indicate a significant relationship. A) Relationship with GDD (Growing Degree Days °C); B) Relationship with FDD (Freezing Degree Days °C) and C) Relationship with Snow days.

## 4. Discussion (1515words main text)

This study aimed to explore biological correlates and ecological drivers of seed oil content and seed oil composition in a local pool of 47 alpine species. We found strong indications that seed oil content significantly influences seed longevity, potentially impacting soil seed banks and conservation strategies for seed bank managers or restoration programmes. Interestingly, we found little evidence of temperature-related patterns pointing to macroevolutionary processes driving seed oil content and composition at the regional or local alpine scales.

While oil content is known to vary globally from 1% in *Musa paradisiaca* to 76% in *Chrysobalanus icaco* (Matthäus, 2012), our alpine species showed a reduced but still relevant range of seed oil percentages (1-34%). This variation is similar to the results of other studies focused on herbaceous plants, like the one exhibited by weed species in France (Bretagnolle *et al.*, 2016). These patterns align with the general variation in seed oil predicted for herbaceous species in the pioneer studies of Levin (1974). The seed oil composition we found in our alpine species was also comparable to what Bretagnolle *et al.* (2016) reported for weed species, except for erucic acid (C22:1n9) in Brassicaceae. Previous reports from other commercial Brassicaceae species confirmed high proportions of elongated acyl chains from C20 to C24 in this family (Baud and Lepiniec, 2010). The high levels of erucic acid are surprising due to its “high” melting point (33.5°C) (Sanyal and Linder, 2013), which would hinder energy release; however, erucic acid has also been associated with lower seed predation (Jong *et al.*, 2016).

Within our alpine system, we could not corroborate the expected negative relationships between seed mass and seed oil content described in the literature (Nagel and Börner, 2010; Hamilton *et al.*, 2013; Bretagnolle *et al.*, 2016). A plausible explanation for this result is that most studies reporting significant relationships include different vegetation types and multiple comparisons among herbs, shrubs, and trees. In contrast, our study focused on similar life forms, mostly herbaceous hemicryptophytes and dwarf chamaephytes, where seed mass and other biological traits are expected to be more constant (Díaz *et al.*, 2016). Similar non-significant trends were reported in another study focused on trees (Finkelstein and Grubb, 2002), suggesting that life forms constrain global variation in seed oil content. Nevertheless, more studies across different life forms and vegetation types are still necessary to confirm this idea. In our local alpine species pool, we also found a negative trend between seed mass and UFA/SFA ratio. However, this trend contradicts the expectation of small seeds storing more saturated fatty acids (Linder, 2000), i.e. lower ratio values. We found a considerable variation of oil content in small seeds, although such variation strongly decreases towards low oil values in large seeds, supporting the results reported by Bretagnolle *et al.* (2016). We found strong indications of a strong phylogenetic signal of seed oil content and oil composition, confirming previous results (Sanyal and Decocq, 2016), with species from the same family mostly grouped together and a strong phylogenetic signal of seed mass. Interestingly, with our regional dataset, we consistently found a strong phylogenetic signal of seed oil content but not of seed mass, suggesting that alpine species are filtered towards small seed mass independently of their phylogeny.

In concordance with our expectations, we found a significant negative relationship between seed longevity and seed oil content. Alpine seeds, generally known for their short longevity (Mondoni *et al.*, 2011), showed a consistent and significant decrease in seed longevity with increasing oil content. The UFA/SFA ratio also showed a decreasing trend of longevity with more unsaturated fatty acids, although not significant. This finding supports the hypothesis that oily seeds have lower longevity (Nagel and Börner, 2010; Neto *et al.*, 2019) and that a higher proportion of unsaturated fatty acids could be associated with shorter longevity (Hoekstra, 2005; Probert, Daws and Hay, 2009). Our results are consistent with previously described adverse effects observed in seed longevity, possibly due to lipid oxidation (Bailly *et al.*, 1998; Tammela *et al.*, 2003) and contrast with other studies reporting no relationship between seed longevity and oil content (Walters, Wheeler and Grotenhuis, 2005; Probert, Daws and Hay, 2009; Gardarin *et al.*, 2010). With the current biodiversity crisis, plant conservation is a global priority, especially ex-situ conservation in seed banks. Having the necessary knowledge and appropriate methods to store and maintain seed viability in these facilities is vital. The results presented here suggest that, for alpine species, seed oil content can be an important factor to control for in seed banks. Measuring seed oil content and composition is a destructive yet fast method that can be used to separate those alpine species with particularly low seed longevity, which can then be included in more frequent germination screenings and seed lot renewals.

Based on previous studies by Sanyal and Decocq (2016) and Linder (2000), we predicted that alpine species, adapted to live and germinate under colder temperatures, would show earlier germination with increasing oil content and an increasing UFA/SFA ratio. Surprisingly, the negative relationship between oil content and germination timing contradicted our initial expectations. The lack of patterns with oil content is not completely surprising in wild species, as has been reported in rainforest species in Australia (Hamilton *et al.*, 2013), whereas weeds (Gardarin, Dürr and Colbach, 2011) and crops (Gu *et al.*, 2019; Cheng *et al.*, 2024) have shown positive correlations. Observing this pattern between wild and crop-related species (cultivars or weeds) stresses the importance of human selection of varieties or cultivars exhibiting high oil content and earlier germination (Gardarin, Dürr and Colbach, 2011; Gu *et al.*, 2019). This artificial selection has encouraged a lot of research focused on increasing oil content (Miao *et al.*, 2020) or improving oil composition for nutritional purposes, even for the production of biofuels (Dierig *et al.*, 2006). Nevertheless, these data are unrepresentative of natural patterns in wild species and more research is needed to fill this knowledge gap. We also observed the opposite expected trend with species with higher unsaturated fatty acids germinating later against Linder (2000) results. A plausible explanation could be that alpine species' physiological seed dormancy constraints (Schwienbacher *et al.*, 2011) drive germination patterns more strongly than oil content and composition.

Against our expectations, we found no significant differences along the altitudinal gradient at the regional scale and did not find any significant relationship with the species' local ecological preferences. It is important to remember that here, we focused on species-level traits, with the expectation that species with different ecological niches along an altitudinal and microtopographical gradient have suffered selection for oil content. Previous research involving different populations of a few cultivars species also failed to detect a significant positive correlation between seed oil and elevation (Dierig *et al.*, 2006; Cai *et al.*, 2012). These results might indicate a low effect of elevation in seed oil content (Dierig *et al.*, 2006), although there are reports of a positive correlation between oil content and mean temperatures (Piper and Boote, 1999). Parallelly, Cai *et al.* (2012) did find a positive correlation between elevation and percentage of unsaturated fatty acids within a single species grown in a broad altitudinal range, possibly due to plasticity or differential physiological functioning. None of the local microenvironmental gradients studied (GDD, FDD and snow) showed any significant relationship with oil content or composition. However, all three gradients show a consistent trend with species living preferentially colder places (low GDD, high GDD and higher snow days) tended to have higher oil content and higher proportion of unsaturated fatty acids, which could potentially be consistent with the anti-freezing properties that unsaturated fatty acids can provide (Linder, 2000). The lack of regional and local alpine patterns seems to limit Sanyal and Linder's claims (Linder, 2000; Sanyal and Linder, 2013; Sanyal and Decocq, 2016) of strong temperature selection at the macroevolutionary level, constraining seed oil patterns. In the present study, we are working with a cold-adapted alpine flora, with different degrees of cold preference/tolerance. Life in cold-season regions may impose some constraints on seed oil properties, such as temperatures during seed maturation, and these constraints may be so tight that they leave no room for within-region variation. However, providing a more complete picture is difficult due to the limited information on oil content and fatty acid composition available for wild species (Levin, 1974), especially involving other habitats and biomes.

The reported results add new insights to the seed ecological spectrum and provide a small step forward to comprehending seed trait ecology in line with the framework described by Saatkamp *et al.* (2019). Specifically, in the understudied topic of seed oil and composition and their correlates and drivers in wild species. Seed oil content strongly influences seed longevity in alpine species, which supplies relevant information for seed bank managers, while we found no significant patterns of correlation with seed mass and germination timing. Ecological drivers appear to have a limited effect on seed oil and composition along regional altitudinal and local microtopographical gradients. Nevertheless, collective and global efforts are necessary to complement the few data available for wild species in order to disentangle seed oil patterns across species and vegetation types.

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