



SHORT COMMUNICATION **OPEN ACCESS**

Lipid Phase Behaviour of the Curvature Region of Thylakoid Membranes of *Spinacia oleracea*

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ABSTRACT

Thylakoid membranes (TMs) of oxygenic photosynthetic organisms are flat membrane vesicles, which form highly organised, interconnected membrane networks. In vascular plants, they are differentiated into stacked and unstacked regions, the grana and stroma lamellae, respectively; they are densely packed with protein complexes performing the light reactions of photosynthesis and generating a proton motive force (pmf). The maintenance of pmf and its utilisation for ATP synthesis requires sealing the TMs at their highly curved regions (CRs). These regions are devoid of chlorophyll-containing proteins but contain the curvature-inducing CURVATURE THYLAKOID1 (CURT1) proteins and are enriched in lipids. Because of the highly curved nature of this region, at the margins of grana and stroma TMs, the molecular organisation of lipid molecules is likely to possess distinct features compared to those in the major TM domains. To clarify this question, we isolated CR fractions from *Spinacia oleracea* and, using BN-PAGE and western blot analysis, verified that they are enriched in CURT1 proteins and in lipids. The lipid phase behaviour of these fractions was fingerprinted with ³¹P-NMR spectroscopy, which revealed that the bulk lipid molecules assume a non-bilayer, isotropic lipid phase. This finding underpins the importance of the main, non-bilayer lipid species, monogalactosyldiacylglycerol, of TMs in their self-assembly and functional activity.

1 | Introduction

The thylakoid membranes (TMs) exhibit a complex architecture that accommodates the main protein components performing the light reactions of oxygenic photosynthesis. In chloroplasts, they form extended multilamellar networks of flattened vesicles, separating the inner and outer aqueous phases, the lumen and the stroma, respectively. In vascular

plants, TMs are organized into two main domains: the granum, consisting of tightly stacked membranes, and the stroma lamellae, which are unstacked membranes winding around the grana in a quasi-helical manner (Mustárdy and Garab 2003; Bussi et al. 2019). The protein composition in plant TMs displays lateral heterogeneity (Andersson and Anderson 1980; Dekker and Boekema 2005; Trotta et al. 2025). Photosystem II (PSII) and its associated light-harvesting antenna complex

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(LHCII) are predominantly located in the appressed membranes, while Photosystem I (PSI) and its light-harvesting antenna, LHCI, along with the ATP synthase, reside in the stroma lamellae. The cytochrome *b6f* complexes are distributed uniformly throughout the membrane system (Dekker and Boekema 2005).

In addition to these two basic building units of TMs, a third domain, the granum margin (GM), has been proposed to be a compositionally, structurally, and functionally distinct region (Albertsson 2001; Armbruster et al. 2013). However, different experimental techniques yielded varying interpretations of this membrane domain. On one hand, certain electron microscopy (EM) studies identified the GM as highly curved areas at the periphery of grana, comprising roughly 5%–7% of the TM area, while experiments employing mechanical fragmentation isolated a 60 nm wide margin annulus, revealing a significant contact area between PSII and PSI domains (Albertsson 2001). Recently, a detailed biochemical analysis using digitonin solubilization of *Arabidopsis* TMs has demonstrated that GM segregates in two distinct regions: (i) the connecting domain (CD), which physically connects grana with stroma lamellae and hosts both photosystems, and (ii) the domain containing curvature particles, which are deficient in photosystems but are enriched in CURT1 proteins (Trotta et al. 2025). This latter, highly purified CURT1-enriched domain, the curvature region (CR) fraction, has been identified using a low concentration of digitonin (0.4% or 0.25%); this fraction (aka loose pellet) is largely devoid of chlorophyll-binding protein complexes (Trotta et al. 2019). Hence, CR represents a subdomain of GM. It is noteworthy that CR possesses approximately three times more lipids relative to chlorophyll (Chl) and about twice as much relative to protein than TMs (Koochak et al. 2019). This suggests that the generation of the CR domain, besides CURT1, depends on the organization of the lipid molecules.

Using mainly ^{31}P -NMR spectroscopy, a technique for fingerprinting the lipid phase behaviour of phospholipid-containing assemblies in vivo and in vitro (Watts 2013), it has been documented that the bulk lipid molecules in plant TMs display marked polymorphism: in addition to the bilayer or lamellar (L) phase, they contain an inverted hexagonal (H_{II}), and at least two isotropic (I) lipid phases (Garab et al. 2022). The non-bilayer propensity of TMs is evidently due to the presence of their main (~50%) non-bilayer lipid species, monogalactosyldiacylglycerol (MGDG). It has also been clarified that the lipid polymorphism in isolated granum and stroma TMs is very similar to that in TMs, showing that the strikingly different protein composition of these domains does not bring about different lipid polymorphism (Dlouhý, Javornik, et al. 2021). Experiments using lipases and proteases have revealed that the non-bilayer lipid phases are found in different subdomains of TMs, distinct from the bilayer domains enriched in PSI and PSII supercomplexes. I phases were found in the luminal compartment, where they form VDE:lipid assemblies (Dlouhý et al. 2020), and were also proposed to be involved in membrane fusion (Böde et al. 2024). (VDE, violaxanthin de-epoxidase, a water-soluble lipocalin-like photoprotective enzyme, the activity of which has been shown to require a non-lamellar lipid phase (Goss and Latowski 2020).) Trypsin-digestion experiments led to the inference that lipids in the H_{II}

phase are associated with stroma-exposed proteins or polypeptides (Dlouhý et al. 2022). It is noteworthy that CURT1 proteins were shown to be susceptible to trypsin digestion (Armbruster et al. 2013).

The aim of this work was to clarify the lipid phase behaviour of the CR region isolated from *Spinacia oleracea*. To this end, we have established that our CR preparation is enriched in CURT1 protein, as previously reported for *Arabidopsis thaliana* (Trotta et al. 2025) and therefore, it is used here as a control for TM fractionation. Using ^{31}P -NMR spectroscopy, we showed that the bulk lipid molecules in CR do not form a bilayer (L) phase, but are assembled into isotropic (I) phases.

2 | Materials and Methods

2.1 | Isolation of the Curvature Region

To isolate the curvature region (CR) or loose pellet, we followed the step-by-step protocol described by (Suorsa et al. 2015). *Spinacia oleracea* (hereafter spinach) leaves were homogenized in ice-cold B1 buffer (50 mM HEPES/KOH (pH 7.5), 330 mM sorbitol, 5 mM MgCl_2 , 2.5 mM ascorbate, 0.05% (w/v) BSA), then filtered through two layers of nylon mesh and centrifuged at $5000\times g$ at 4°C for 4 min. Then, we suspended the resulting sediment in 30–40 mL of B2 buffer (50 mM HEPES/KOH (pH 7.5), 5 mM sorbitol, 5 mM MgCl_2) and centrifuged again at $5000\times g$ for 4 min. We suspended the sediment in 30–40 mL of B4 buffer (50 mM HEPES/KOH (pH 7.5), 100 mM sorbitol, 5 mM MgCl_2), which was followed by another centrifugation with the same parameters, then we suspended the sediment again in a small amount of B4 buffer. The volume was adjusted to $0.5\text{ mg Chl mL}^{-1}$ with B3 buffer (15 mM Tricin (pH 7.8), 100 mM sorbitol, 10 mM NaCl, 5 mM MgCl_2) and digitonin (Sigma-Aldrich) at a final concentration of 0.4% (w/v). The suspension was stirred for 8 min with a magnetic stirrer at room temperature and then centrifuged at $1000\times g$ for 3 min. The supernatant was centrifuged for 30 min at $10,000\times g$ to separate the core of grana; the supernatant was centrifuged at $40,000\times g$ for another 30 min and the pellet containing the CD was collected. Following these steps, the supernatant was centrifuged at $144,000\times g$ for another 1 h to separate the solid sediment (containing the stroma lamellae) and the loose pellet containing the CR. The fractions collected were suspended in B4 buffer, and the suspension was stored at -80°C until use. *Arabidopsis thaliana* (hereafter *Arabidopsis*) TM and CR, obtained with 0.4% DIG, were isolated using the same procedure as in (Trotta et al. 2025).

2.2 | Enzymatic Treatments With Wheat Germ Lipase and Trypsin

Isolated CR particles were treated with wheat germ lipase (WGL)—a substrate nonspecific general tri-, di-, and monoglyceride hydrolase/lipase (Kublicki et al. 2021). WGL was purchased from Sigma-Aldrich (Burlington) and was applied at a concentration of 20 U mL^{-1} ; the treatments and incubations were performed at 5°C . Earlier, using thin-layer chromatography, we verified that WGL digests the main TM lipid species, MGDG (Dlouhý et al. 2022; Böde et al. 2024).

For trypsin (T8003; trypsin from bovine pancreas; Sigma-Aldrich) treatment, a stock of 300 mg mL⁻¹ was dissolved in demineralised water, and 33 μ L per mL of sample was added for a total concentration of 10 mg mL⁻¹. The suspension was thoroughly mixed and kept at 5°C until the start of the measurements.

2.3 | IpBN-PAGE and Western Blotting

Analysis of the pattern of TM protein complexes was performed by large pore Blue-Native Polyacrylamide Gel Electrophoresis (IpBN-PAGE) as described in (Sandoval-Ibáñez et al. 2021). Analysis of the amount of PSI, PSII, and CURT1 proteins was performed by immunoblotting as described in (Trotta et al. 2025), by using PsaB antibody (for PSI, Agrisera www.agrisera.com catalogue numbers AS10695), CP47 antibody (kind gift of Prof. Roberto Barbato) and CURT1A antibody (Armbruster et al. 2013). Samples were loaded on a chlorophyll basis (5 μ g for IpBN-PAGE and 1 μ g for immunoblotting).

2.4 | Scanning Electron Microscopy

The membrane fractions were fixed in 2.5% glutaraldehyde for 2 h, settled on poly-L-lysine-coated polycarbonate filter for 45 min. After post-fixation in 1% OsO₄ for 50 min, the samples were dehydrated in aqueous solutions of increasing ethanol concentrations, critical point dried, covered with 5 nm gold by a Quorum Q150T ES (Quorum Technologies) sputter, and observed in a JEOL JSM-7100F/LV (JEOL SAS) scanning electron microscope.

2.5 | ³¹P-NMR Spectroscopy

³¹P-NMR spectroscopy tracks the motion of bulk phosphatidylglycerol (PG) molecules in the TM, serving as a sensitive internal probe of the lipid phase behaviour. This is justified by the fact that PG displays no lateral heterogeneity among the bulk phase of TM domains (Duchene and Siegenthaler 2000) and is homogeneously distributed in TM lipid mixtures (van Eerden et al. 2015).

³¹P-NMR spectroscopy was performed as described in (Dlouhý, Javorník, et al. 2021). Briefly, the spectra were recorded at 5°C, using an Avance Neo 600 MHz NMR spectrometer (Bruker) with a BBFO SmartProbe that was tuned to the P-atom frequency. The sample was loaded into 5 mm diameter NMR tubes. For spectra acquisition, 40° RF pulses with an inter-pulse time of 0.5 s were applied without ¹H-decoupling, as in earlier experiments (Krumova et al. 2008). As an external chemical-shift reference, an 85% solution of H₃PO₄ in water was used. In the saturation transfer (ST) experiments, we employed RF pulses with low power at the designated frequency for a duration of 0.3 s, followed by 40° pulses with an acquisition time of 0.2 s and a repetition time of 0.5 s. The intensity of the pre-saturation pulse was adjusted based on the intensity of the saturated peak. For the pre-saturation RF pulses, the field strength was set to 40 Hz.

CR samples from two batches, with very similar features, were averaged to improve the signal-to-noise ratio. For ³¹P-NMR data

processing, TopSpin software (Bruker) was used. The figures were plotted using MATLAB R2020b (MathWorks Inc.).

2.6 | Fourier-Transform Infrared Spectroscopy

To determine the lipid-to-protein ratios in TMs and CR, we used FTIR spectroscopy (see [Supporting Information](#)). Membrane fraction suspensions were washed in D₂O-based PBS (phosphate-buffered saline) solution for complete H₂O to D₂O exchange. The sample was layered between CaF₂ windows, separated by an aluminium spacer, and placed in a Bruker Vertex70 FTIR spectrometer. Spectra were recorded at 5°C between 4000 and 900 cm⁻¹, 512 interferograms were collected for each spectrum, with a spectral resolution of 2 cm⁻¹. The infrared absorption spectrum of the samples was calculated from the background and the sample of single beam spectra with Opus software of Bruker. For the analysis of the structural properties of the membrane, the ester C=O plus the amide I region, which in this paper will be referred to as 'Ester + Amide I' region, was used between 1800 and 1595 cm⁻¹. To obtain the relative intensities of the 'ester' and the 'amide I' bands in the 'Ester + Amide I' region, a 3rd order polynomial was fitted and subtracted as baseline. The area under the curves was calculated by using built-in MATLAB functions, after trimming the spectral regions corresponding to the 'ester' and 'amide I' bands.

3 | Results

3.1 | Pattern of Protein Complexes in TM Fractions Obtained by Digitonin Solubilization

The pattern of the protein complexes in spinach TM and the fractions obtained by 0.4% digitonin solubilization were analyzed by IpBN-PAGE (Figure 1A). The typical pattern observed in fractions obtained with the same procedure in Arabidopsis was observed (Trotta et al. 2025). In particular, the 10 k fraction, enriched in grana core, contained the majority of the PSII-LHCII supercomplexes (sc) and LHCII complexes (M-, L- and monomeric LHCII). On the other hand, the 144 k fraction showed the typical pattern of stroma lamellae, being enriched in PSI-LHCI and other PSI complexes, and depleted in LHCII. The 40 k fraction, enriched in CD, had somewhat intermediate features. Considering the three strong pellet fractions (i.e., 10, 40 and 144 k), PSII monomer and Cyt b₆f were mostly enriched in the 10 and 40 k fractions. However, the highest amount of PSII monomer and Cyt b₆f, together with ATP synthase, was present in the CR fraction, which was instead strongly depleted of Chl-binding protein complexes (Figure 1A). A similar pattern was visible in the Arabidopsis CR, which, in line with expectations using 0.4% digitonin, contained Chl-binding protein complexes to some extent (compare with (Trotta et al. 2025)).

The analysis by immunoblotting confirmed the results obtained with IpBN-PAGE that is, enrichment of PSII in 10 k and of PSI in 144 k (Figure 1B). Consistent with previous results (Trotta et al. 2025; Trotta et al. 2019) the CR fraction had the highest enrichment in CURT1, indicating that, besides other proteins collected in

the loose pellet due to digitonin solubilization, this was the fraction where most of the curvature domain of the GM was collected.

3.2 | Morphology of Curvature Region

For the structural characterisation of CR, scanning electron microscopy (SEM) was performed (Figure 2). The CR fraction consisted of small particles with a tendency to create large self-aggregates and/or smaller vesicle-like assemblies.

The observed particles are likely to originate from the highly curved regions of the TMs, which are enriched in CURT1 proteins. These structures are distinct from those observed in the grana and stroma fractions; in the latter, predominantly spherical membrane vesicles appear, as reported in previous studies (Koochak et al. 2019; Dlouhý, Karlický, et al. 2021). Notably, the

morphology of these particles does not appear to contradict the observations of (Koochak et al. 2019) using TEM either.

3.3 | Lipid Phase Behaviour of CR Particles

As evidenced by the ^{31}P -NMR spectra shown in Figure 3, the CR particles do not contain L and H_{II} phases, which are present in intact TMs and the granum and stroma subchloroplast membrane particles isolated with the same procedure (Dlouhý, Javornik, et al. 2021) (see also the ^{31}P -NMR spectrum of stroma lamellae in the Supporting Information). Instead, the spectra of CR preparations are dominated by well-discernible I phases with peak positions at around 0.15 and 2.7 ppm. To probe the spectral shapes and resolve overlapping phases, we applied low power radiofrequency pulses at selected chemical shifts (Krumova et al. 2008; Dlouhý, Javornik, et al. 2021). These

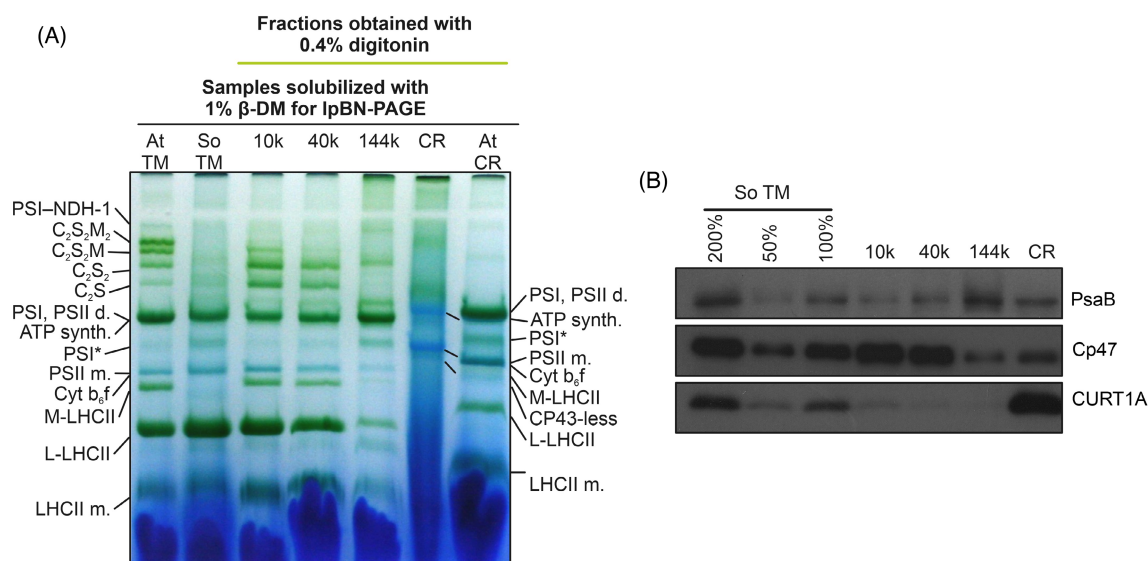


FIGURE 1 | (A) IpBN-PAGE analysis of TM and fractions after solubilization with 0.4% digitonin of spinach (So) TM. At: *Arabidopsis thaliana*. (B) immunoblot with PsaB, Cp47 and CURT1A antibodies of the same TM and fractions as in (A). A cross-reactivity scale with So TM is shown on the left.

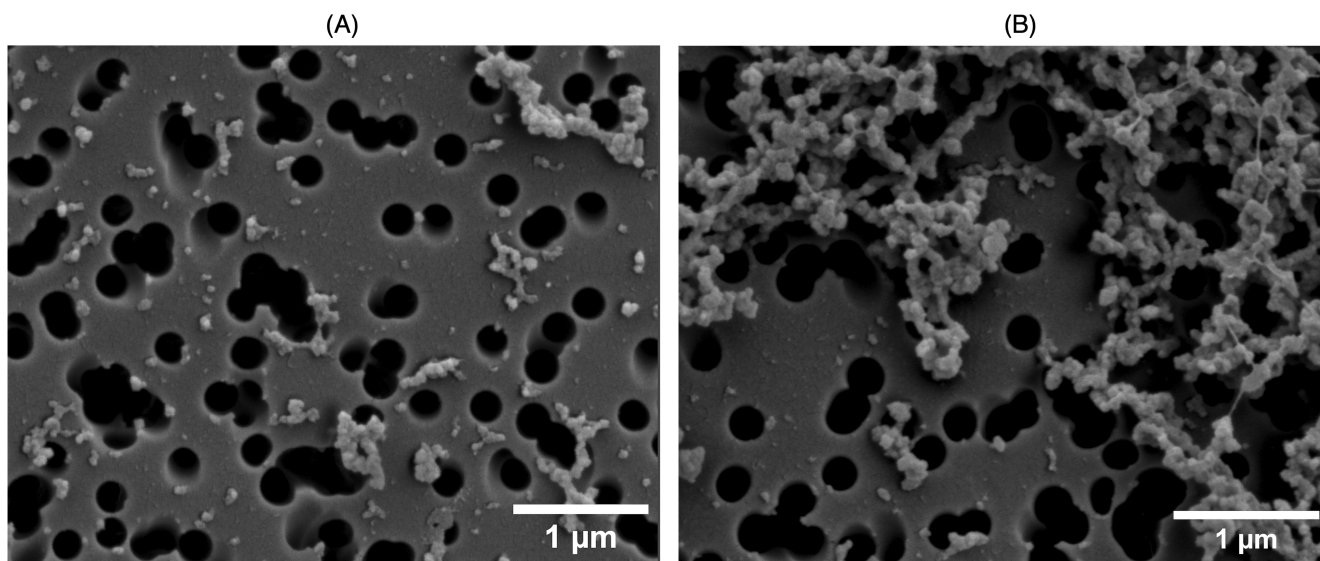


FIGURE 2 | SEM micrographs of CR fractions showing bent membrane particles (A) which often coagulate (B).

saturation transfer experiments confirmed the presence of these two strongly overlapping phases, although other resonances with smaller amplitudes could also be found in the band structure.

The resonance bands that originate from the lipid phases were sensitive to WGL digestion (Figure 4A), similarly to what has been observed in TMs and the granum and stroma subchloroplast particles (Dlouhý et al. 2022; Dlouhý, Javornik, et al. 2021). This shows that the character of the I phases in CR closely resembles those in intact TM and in the digitonin-fragmented grana and stroma lamellae.

In contrast, trypsin treatment did not elicit any sizeable effect on the spectra (Figure 4B). As we have previously shown (Dlouhý

et al. 2022), trypsin treatment selectively degraded the ^{31}P -NMR detectable H_{II} phase of TMs, which revealed that this lipid phase is associated with proteins or extrinsic protein domains protruding in the stromal aqueous side of TMs. Although CURT1 has been shown to be susceptible to trypsin digestion, the absence of the H_{II} phase and the trypsin-insensitive nature of the detected non-bilayer lipid phases show that these proteins do not contribute to the formation of the H_{II} phase.

It is to be noted here that while plastoglobuli might contribute to the composition of the CR fraction (Trota et al. 2025), they display no sizeable ^{31}P -NMR signal (Dlouhý et al. 2022); hence, they are highly unlikely to participate in the isotropic organisation of the lipid molecules around CURT1 proteins.

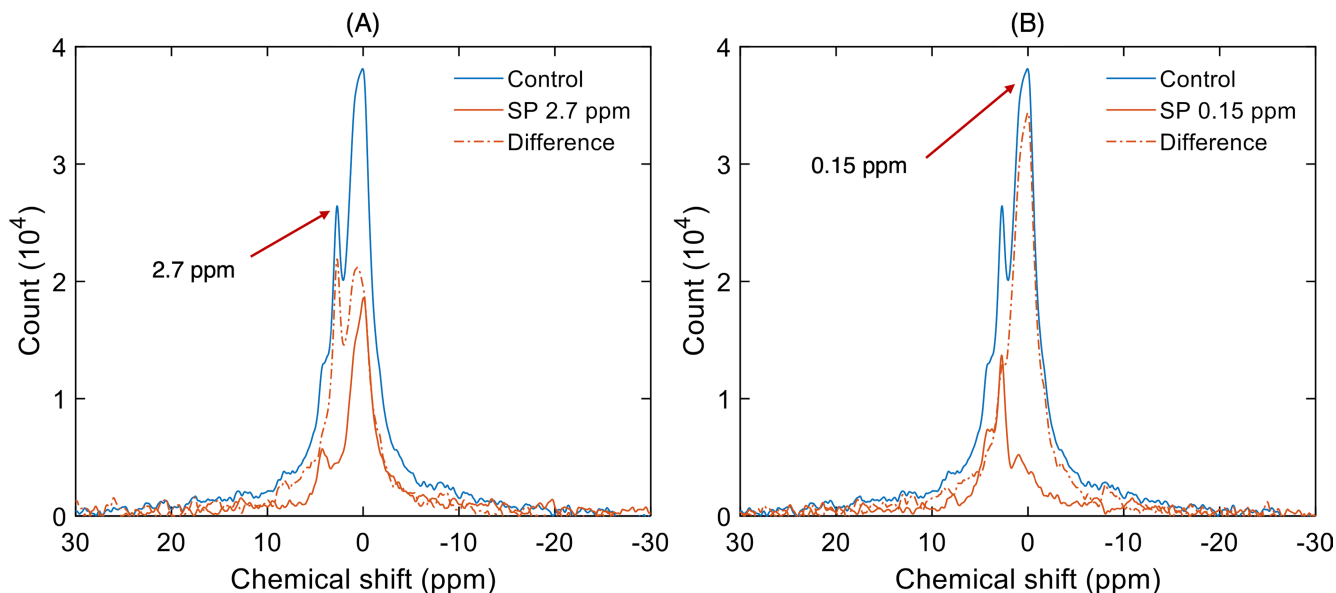


FIGURE 3 | ^{31}P -NMR spectra of CR under control conditions, displaying also the effects of saturation pulses (SP) at 2.7 ppm (A) and 0.15 ppm (B). The number of scans was 12,800.

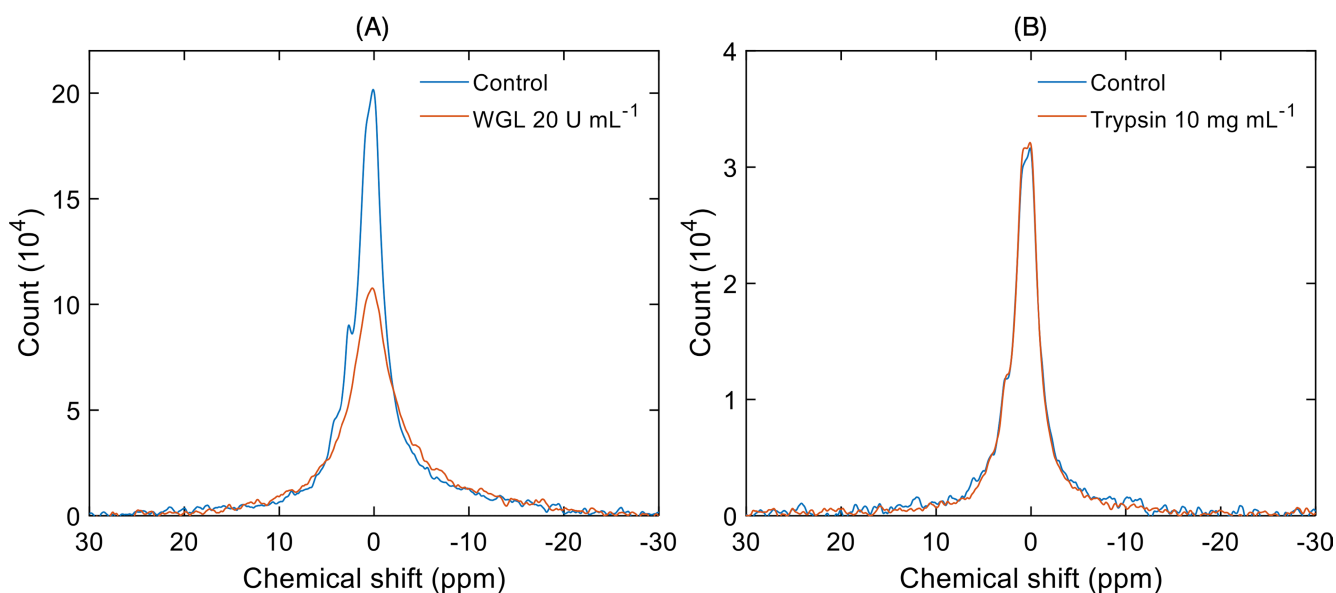


FIGURE 4 | ^{31}P -NMR spectra of untreated (Control) and WGL-treated (20U mL^{-1} , 25,600 scans) (A) and trypsin-treated (10mg mL^{-1} , 19,200 scans) (B) CR samples.

4 | Discussion

In this paper, we confirmed the presence of CURT1 in the CR fraction of *Spinacia oleracea*, which corroborates the notion that this protein plays the main role in generating membrane curvature in various species (Armbruster et al. 2013; Heinz et al. 2016; Sandoval-Ibáñez et al. 2021). Notably, the CURT1 protein was detected using an antibody originally developed for *Arabidopsis thaliana*, demonstrating its cross-reactivity and reinforcing the evolutionarily conserved nature of CURT1 across taxa.

³¹P-NMR spectroscopy—which tracks the motion of bulk PG molecules in different lipid phase environments—has unveiled that the CR fraction of TMs displays merely an intense and a weaker isotropic (I) lipid phase. The lack of the lamellar (L) phase in CR preparations is evidently the consequence of the deficiency of PSII and PSI supercomplexes, which in TMs stabilize the bilayer, similarly to that shown for LHCII:MGDG macroassemblies (Simidjiev et al. 2000). Note that the absence of the L phase is unlikely to be caused by the usage of digitonin since both the granum and the stroma TM subdomains—with the same isolation procedure—preserve the bilayer (L) phase (Dlouhý, Javornik, et al. 2021).

The presence of I phases in the CR might be attributed to curved bilayers (Cullis and DE Kruijff 1979). However, the observed inverse relationship between the insulation efficiency of TMs and the intensity of I phase(s) renders this explanation unlikely; with increasing I phase contributions at the expense of L phase, the permeability of the membrane increases (Ughy et al. 2019). Hence, the organisation of lipid molecules in CR evidently deviates from a typical “well-sealed” bilayer arrangement. It is interesting to note here that, as argued by (Garab et al. 2025), the lipid polymorphism of TMs does not necessarily conflict with the efficient energisation of TMs and utilisation of pmf for ATP synthesis. Also, as pointed out by (Goni 2014), “a membrane composed solely of lamellar lipids would be an optimum insulator, only non-compatible with cell function, that is, life.”

Overall, these findings strongly indicate that the lipids surrounding CURT1 proteins are organized into isotropic arrays. This means that the lipids surrounding the CURT1 protein lack the inherent tendency to adopt a bilayer arrangement in the highly curved CR of TMs. These data underline the role of the major non-bilayer lipid, MGDG, of plant TMs in their self-assembly, and in general, corroborate the fundamental importance of lipid polymorphism in these major, energy-converting membranes.

Author Contributions

The study was conceptualized by G.G., K.B., A.T., E.-M.A., and V.Š.; TM and CR preparations were isolated by O.Z. and K.B.; ³¹P-NMR spectroscopy measurements were performed by U.J., supervised by J.P.; data analyses were performed by K.B. and O.D., with the help of U.J., supervised by G.G.; spectral characterizations of the membranes were performed by K.B., O.D., supervised by V.Š.; FTIR measurements were carried out and analyzed by K.B.; SEM experiments were performed by I.D.; lpBN-PAGE and immunoblotting was performed by V.P. and H.F., supervised by A.T. and E.-M.A. The paper was written by G.G., K.B., and A.T., with all authors contributing to the writing.

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Conflicts of Interest

The authors declare no conflicts of interest.

Data Availability Statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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Supporting Information

Additional supporting information can be found online in the Supporting Information section.