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Enhancement of Electron Spin Echo Envelope Modulation Spectroscopic Methods to Investigate the Secondary Structure of Membrane Proteins

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Abstract

This paper reports on a significant improvement of a new structural biology approach designed to probe the secondary structure of membrane proteins using the pulsed EPR technique of Electron Spin Echo Envelope Modulation (ESEEM) spectroscopy. Previously, we showed that we could characterize an α -helical secondary structure with ESEEM spectroscopy using a ^2H -labeled Val side chain coupled with site-directed spin-labeling (SDSL). In order to further develop this new approach, molecular dynamic (MD) simulations were conducted on several different hydrophobic residues that are commonly found in membrane proteins. ^2H -SL distance distributions from the MD results indicated that ^2H -labeled Leu was a very strong candidate to significantly improve this ESEEM approach. In order to test this hypothesis, the secondary structure of the α -helical M26 peptide of the acetylcholine receptor (AChR) incorporated into a bicelle was investigated with ^2H -labeled Leu d_{10} at position 10 (i) and nitroxide spin labels positioned 1, 2, 3 and 4 residues away (denoted $i+1$ to $i+4$) with ESEEM spectroscopy. The ESEEM data reveal a unique pattern that is characteristic of an α -helix (3.6 residues per turn). Strong ^2H modulation was detected for the $i+3$ and $i+4$ samples, but not for the $i+2$ sample. The ^2H modulation depth observed for ^2H -labeled d_{10} Leu was significantly enhanced (x4) when compared to previous ESEEM measurements that used ^2H -labeled d_8 Val. Computational studies indicate that deuterium nuclei on the Leu sidechain are closer to the spin label when compared to Val. The enhancement of ^2H modulation and the corresponding Fourier Transform (FT) peak intensity for ^2H -labeled Leu significantly reduces the ESEEM data acquisition time for Leu when compared to Val. This research demonstrates that a different ^2H -labeled amino acid residue can be used as an efficient ESEEM probe further substantiating this important biophysical technique. Finally, this new method can provide pertinent qualitative structural information on membrane proteins in a short time (few minutes) at low sample concentrations ($\sim 50 \mu\text{M}$).

Keywords

Electron Spin Echo Envelope Modulation; Membrane Protein; Spin Labeling; EPR

INTRODUCTION

A novel approach is being developed to characterize the secondary structure of membrane proteins utilizing Electron Spin Echo Envelope Modulation (ESEEM) spectroscopy coupled with site-directed spin-labeling (SDSL) ¹. Despite the importance and large number of

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membrane proteins, relatively limited structural information is known about them^{2,3}. New biophysical techniques are needed to probe their structural properties. Their hydrophobic nature and low expression yields cause difficulties for traditional structural techniques such as x-ray crystallography and solution NMR spectroscopy^{3,4}.

ESEEM spectroscopy indirectly observes NMR transitions through an electron spin coupled to a nearby NMR active nucleus⁵⁻⁷. ESEEM can detect weak dipolar interactions between a ²H atom and a spin label out to a maximum distance of approximately 8 Å^{7,8}. The modulation depth for weakly coupled nuclei is proportional to 1/*r*⁶. Previously, we demonstrated using ²H-labeled Val as a probe and a strategically placed MTSL (*S*- (2,2,5,5-tetramethyl-2, 5-dihydro-1H-pyrrol-3-yl) methyl methanesulfonothioate), α-helical content could be detected with ESEEM spectroscopy¹. The unique structure of an α-helix consists of 3.6 residues per turn with a vertical distance of 5.4 Å separating the turn. When a Cys mutated nitroxide spin label is positioned 1, 2, 3 and 4 residues away (denoted *i*+1 to *i*+4 respectively) from a ²H-labeled residue such as the d₈ Val (*i*) side chain, the periodicity of the α-helical structure gives rise to a unique pattern in the collective ESEEM data. The *i*+3 and *i*+4 samples reveal ²H modulation from the dipolar coupling between the ²H nuclei of the Val side chain and spin label (< 8 Å). However, in the *i*+1 and *i*+2 samples, the ²H nuclei are beyond the detection limit (> 8 Å). Due to the structure and dynamics of different amino acid side chains, a variety of patterns and signal intensities can be obtained using this approach^{9,10}. The aim of this approach is to provide additional tools for probing the structure of membrane proteins using SDSL and EPR spectroscopy.

In this work, a molecular modeling study was conducted to obtain distance distributions between ²H nuclei on different amino acid side chains and a spin label. Four different ²H-labeled hydrophobic amino acids (Leu, Val, Phe, Ala) that are commonly found in membrane proteins and commercially available were compared. The Leu side chain was found to a very strong candidate for conducting the present study based on the molecular modeling studies carried on different amino acid residues in the sequence of AChR M26 peptide. ²H-labeled d₁₀ Leu was used to demonstrate the practicality of this method instead of ²H-labeled d₈ valine. The observed experimental ESEEM data have similar patterns for α-helical segments with a significantly enhanced ²H modulation depth. The distance distributions obtained from the modeling studies between ²H nuclei on the Leu side chain and the spin label are consistent with the experimental ESEEM data. These experiments demonstrate that another ²H-labeled amino acid can be used as probe for secondary structure determination with significantly improved sensitivity; thus, making this method applicable to more proteins that contain Leu.

MATERIALS AND METHODS

The M26 subunit of the acetylcholine receptor (AChR) was used as an α-helical model for transmembrane peptides and proteins¹¹⁻¹³. All peptides were synthesized using Fmoc solid-phase peptide synthesis chemistry on a CEM microwave solid phase peptide synthesizer¹⁴. Four different peptides were designed by positioning the ²H-labeled d₁₀ leucine at position 10 (*i*) and the cysteine (X) at four successive positions (*i*+1 to *i*+4). For one complete set of samples, the sequences of peptides are as follows: *i*+1 (EKMSTAISXLLAQAVFLLLSQR), *i*+2 (EKMSTAIXVLLAQAVFLLLSQR), *i*+3 (EKMSTAXSVLLAQAVFLLLSQR), and *i*+4 (EKMSTXISVLLAQAVFLLLSQR), where X represents the position of the cysteine for spin labeling. Additionally, one *i*+3 control sample was prepared without ²H labeling at the Leu10 position.

After the peptides were cleaved from their solid support, reverse-phase HPLC was used for purification with a C4 preparation column and a gradient of 5% to 95% solvent B (90%

acetonitrile)^{15,16}. Purified peptides were labeled with MTSL (Toronto Research Chemicals) in DMSO for 20 hours and excess MTSL was removed by HPLC¹⁶. MALDI-TOF was utilized to confirm the molecular weight and purity of the target peptides. MTSL-labeled M26 peptides were integrated into DMPC/DHPC (3.5/1) bicelles at 1:1000 molar ratio¹⁶. For these experiments, bicelles were used as membrane mimic system and yielded high quality ESEEM data^{17,18}. Comparable ESEEM data could be obtained with liposomes.

X-band CW-EPR (~9 GHz) spectroscopy was used to measure spin concentration (~150 μ M) of all bicelle samples. Three-pulse ESEEM measurements were performed on a Bruker ELEXSYS E580 with an ER 4118X MS3 resonator using a 200 ns tau value with a microwave frequency of ~9.269 GHz at 80 K¹. For all samples, a starting T of 386 ns and 512 points in 12 ns increments were used to collect the spectra¹. All ESEEM data were obtained with 40 OI of bicelle samples and in 30 scans.

Molecular modeling and molecular dynamics studies for each sample were performed using nanoscale molecular dynamics (NAMD)¹⁹ with the molecular graphics software VMD²⁰. The structure of the AChR M26 peptide was obtained from the solution NMR coordinates (PDB entry: 1EQ8). Cysteine mutants were created at *i*+1, *i*+2, *i*+3, and *i*+4 positions using VMD, a MTSL nitroxide spin probe was attached by using CHARMM force-field topology files incorporated in NAMD. The resultant assembly of spin labeled peptides was solvated in a water box. Further equilibration and energy minimization were performed using NAMD simulations. Molecular dynamic simulations were collected out to 1 ns at room temperature using Langevin dynamics under NAMD. This timescale corresponds to the MTSL and leucine sidechain dynamics. The trajectory data were recorded in 1 ps increments. The possible distance distribution for each deuterium and SL was obtained from the analysis of the trajectory data file using VMD. All molecular dynamics simulations were run on a home-built 24-node Linux Beowulf style cluster in our lab.

RESULTS

Molecular modeling and molecular dynamics studies were performed on different hydrophobic amino acids (Ala, Leu, Val, and Phe) in which deuterium labeled side chains are commercially available. Distance distributions were determined between ²H nuclei on the side chain (*i*) and the N-O bond for the SL at the *i*+3 and *i*+4 positions. ²H-SL distance distributions were found to be within the 8 Å ESEEM detection range for all residues studied. However, ²H atoms on the Leu sidechain were found to have the shortest distances; thus, suggesting that Leu would be a strong candidate to significantly improve this ESEEM approach.

Figure 1 shows three-pulse ESEEM data for *i*+3 ²H-labeled d₁₀ Leu10 M26 peptide incorporated into DMPC/DHPC lipid bicelles. Also, a non-deuterated Leu (control) sample is shown for comparison. Both low frequency ²H modulation and high frequency proton modulation appear in the time domain data (Figure 1(A)) for the ²H-labeled d₁₀ Leu10 *i*+3 sample (red). However, only proton modulation appears in the control sample prepared in the absence of ²H-labeled Leu (black). FT frequency domain data for the ²H-labeled d₁₀ Leu10 M26 *i*+3 sample reveal a peak centered at the ²H Larmor frequency of 2.3 MHz. No such peak was observed for the control sample. The ESEEM spectra clearly indicate that the dipolar interaction between the SL and ²H-labeled Leu can be detected for the *i*+3 Leu10 M26 sample.

The three-pulse ESEEM data for the ²H-labeled d₁₀ Leu10 (*i*+1 through *i*+4) M26 peptide are shown in Figure 2, respectively. For ²H-labeled d₁₀ Leu10 M26 peptides, the ²H modulation is observed in the time domain for the *i*+1, *i*+3 and *i*+4 samples. Also, a ²H peak

is clearly observed at the ^2H Larmor frequency in the frequency domain. However, there is no ^2H modulation for the ^2H -labeled d_{10} Leu10 $i+2$ M28 sample. To support these data, molecular modeling and molecular dynamics studies were conducted to estimate distances between the ^2H nuclei on the Leu10 sidechain and the N-O bond on the SL at the $i+1$, $i+2$, $i+3$, and $i+4$ positions¹. ^2H -SL distances for the $i+2$ (9–15 Å) were found to be outside the ESEEM detection range of 8 Å. However, ^2H -SL distances for the $i+1$ (6–11 Å), $i+3$ (5–11 Å) and $i+4$ (3–10 Å) positions were found to be within the ESEEM detection range with different probabilities.

Figure 3(A) shows the comparison of the ESEEM frequency domain data between ^2H -labeled d_{10} Leu10 and ^2H -labeled d_8 Val9 peptides at the $i+4$ position. These frequency domain data reveal a dramatic signal enhancement when ^2H -labeled d_{10} Leu is used instead of ^2H -labeled d_8 Val. Distances and conformations provided by molecular dynamic simulation supported this result. Figure 3(B) displays the minimal energy conformations of the AChR M28 peptides with ^2H -labeled d_{10} Leu10 (left) and ^2H -labeled d_8 Val9 (right) with MTSL at the $i+4$ positions. The figure indicates that the distance from the N-O bond on the nitroxide to the nearest deuteron is much smaller in AChR M28 peptides with ^2H -labeled d_{10} Leu10 (4.4 Å) than that with ^2H -labeled d_8 Val9 (7.6 Å)¹.

DISCUSSION

In previous ^2H -labeled d_8 Val9 experiments, three-pulse ESEEM data showed ^2H peaks for both $i+3$ and $i+4$, but not for $i+1$ and $i+2$ for the AChR M28 peptide¹. This unique pattern is indicative of an α -helical secondary structure. MD simulations suggest that ^2H -labeled Leu d_{10} atoms on the side chain will be closer in distance to the SL for $i+3$ and $i+4$ positions when compared to Val, Ala, Phe. ^2H -labeled Leu d_{10} is a strong candidate for this new methodology method due to its longer chain and larger number of available deuterium atoms within the ESEEM ~ 8 Å detection limit.

For the ^2H -labeled d_{10} Leu10 ESEEM experiments conducted, similar data were obtained with significantly enhanced ^2H modulation as shown in Figure 3(B). MD simulations reveal a significant population of ^2H -SL distances below the 8 Å ESEEM detection limit for ^2H -labeled d_{10} Leu10 when compared to ^2H -labeled d_8 Val9. In the case of Leu, there are two torsion angle rotations about χ_1 and χ_2 and two free rotation modes about the $\text{C}\gamma$ and $\text{C}\delta$ bonds, which correspond to two (CD_3) methyl groups^{9,21,22}. As shown in Figure 3(B), the additional C-C bonds in the Leu sidechain bring the deuterons closer to the N-O nitroxide bond when compared to the Val sidechain. Therefore, the MTSL has a higher probability of being able to detect the ^2H nuclei at a closer distance and resulting in a significant increase in ^2H modulation depth for Leu at this position. However, different conformations of Val and Leu might be favored due to unique sidechain or tertiary interactions. For MTSL, there are three torsional angle rotations about χ_1 , χ_2 and χ_3 and two additional free torsion angle rotations about χ_4 and χ_5 ²³⁻²⁵. Thus, various orientations could be favored due to the interaction of MTSL and peptide backbone or amino acid side chains. All of these factors play a role in the ^2H modulation depth and can alter the corresponding FT intensity. Thus, the observed ^2H modulation depth for ^2H -labeled Leu or Val can vary depending upon the biological system studied.

The ^2H -labeled d_{10} Leu10 ESEEM spectra reveal a ^2H peak for the $i+1$ sample, indicating that ^2H nuclei are weakly coupled to the MTSL. However, no coupling was detected for the $i+2$ sample. Since α -helices have 3.6 amino acids per turn, the Leu residue in the $i+2$ sample lies on the opposite side of the helix as the MTSL. It is expected that distances between ^2H nuclei on the Leu10 sidechain and the $i+1$ SL are closer than that of $i+2$. Also, MD simulations indicated ^2H -SL distances ranged from 9–15 Å for the $i+2$ and 6–11 Å for $i+1$.

sample which matches well with the ESEEM data of the $i+1$ and $i+2$ samples. MD simulations show similar ^2H -SL distance ranges for the $i+1$ and $i+3$ samples. However, the population of distance distributions within the 8 Å limit for $i+3$ is significantly greater than $i+1$ (data not shown), which is consistent with the $i+3$ sample having larger ^2H modulation and a bigger FT ^2H peak in experimental ESEEM data (Figure 2).

Theoretically, the distance between a SL and a ^2H nucleus can be determined by simulation of the ESEEM spectra. For this three-pulse ESEEM ($\pi/2$ - τ - $\pi/2$ -T- $\pi/2$), the modulation depth produced by the dipolar-coupled nucleus is proportional to $1/r^6$. However, due to the variation of conformations and dynamics of the leucine side chain and the MTSL, it is difficult to get quantitative distance information from this method so far^{21,26-28}. Since the three-pulse ESEEM measurements were conducted at 80K, the leucine side chain and MTSL adopt a variety of conformations with respect to each other, yielding a range of distances between the MTSL and multiple ^2H nuclei. In order to improve this approach to get more quantitative distance information, different ^2H -labeled amino acids need to be investigated. Amino acids with only one ^2H atom or magnetically equivalent ^2H nuclei, such as ^2H -labeled d_3 Ala can be employed with this approach to provide quantitative distance measurements via ESEEM spectral simulations.

This ESEEM structural approach is comparable to rotational echo double resonance (REDOR) solid-state NMR spectroscopy. Solid State NMR spectroscopy is a very powerful technique for studying the structure of membrane proteins¹². REDOR NMR can be used to probe the alpha helical component when coupled with spectral simulation by measuring dipolar couplings between NMR active nuclei, such as ^{13}C and ^{15}N ^{29,30}. However, this technique suffers from low signal sensitivity. As a result, REDOR experiments usually require milligram quantities of isotopically labeled protein or peptide and abundant instrumental time. By using ESEEM and SDSL, secondary structure determination for membrane peptides and proteins can be performed on the μmol scale and with minimal data acquisition time (5 to 10 minutes).

CONCLUSION

^2H -labeled d_{10} Leu was shown to be a very powerful secondary structure probe with enhanced ^2H modulation by a factor of 4 when compared to Val. This modulation enhancement leads to a significant increase in sensitivity and corresponding decrease in data acquisition time. Thus, the amount of time required to achieve the same signal-to-noise using ^2H -labeled d_{10} leucine as a probe instead of ^2H -labeled d_8 valine decreases by a factor of ~16. The Leu ESEEM data further validates this structural biology approach and provide researchers with another amino acid to probe secondary structure for proteins and peptides. This ESEEM method uses SDSL and selective deuterium labels, both of which can be incorporated into standard expression systems using routine molecular biology techniques for applications to larger protein systems (not just peptides)^{31,32}. ^2H and ^{15}N residue specific isotopic labeling schemes are used in a variety of NMR structural biology experiments^{33,34}.

This new ESEEM secondary structure approach can be applied to a wide variety of different protein systems that are not amiable to x-ray crystallography or solution NMR including membrane proteins. Several other biophysical techniques such as CD, NMR, CW-EPR, FRET, and 2D IR can provide protein secondary structural information. This ESEEM approach is advantageous because it has no size limitations, is straightforward, uses small spectroscopic labels, minimal sample requirements, and provides selective secondary structural information on specific protein segments. For SDSL EPR researchers, this ESEEM approach will allow researchers to probe not only the dynamics of a specific spin-

labeled site with CW-EPR lineshape analysis, but also the secondary structure. Traditional EPR experiments that probe secondary structure typically require multiple SDSLs in a row to detect differences in lineshape or relaxation that follow a pattern that depends upon the environment and local secondary structure of the protein. Sometimes the EPR data can be ambiguous or challenging to interpret. Certain biophysical techniques such as CD provide global secondary structural information, whereas this ESEEM approach can probe the secondary structure of specific segments of a protein like the solid-state NMR REDOR technique. Direct identification of site-specific secondary structure can be obtained using this ESEEM approach with small amounts of sample and minimal instrumentation time when compared to NMR.

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ABBREVIATIONS

ESEEM	Electron Spine Echo Envelope Modulation
AChR	Acetylcholine Receptor
MTSL	(<i>S</i> -(2,2,5,5-tetramethyl-2,5-dihydro-1H-pyrrol-3-yl)methyl methanesulfonylthioate)
SDSL	Site-Directed Spin-Labeling
DMSO	Dimethyl sulfoxide
MALDI	Matrix-Assisted Laser Desorption/Ionization
TOF	Time of Flight
DMPC	1,2-Dimyristoyl-sn-glycero-3-phosphorylcholine
DHPC	1,2- dihexanoyl-sn-glycero-3- phosphorylcholine
CW	Continuous Wave
EPR	Electron Paramagnetic Resonance
MD	Molecular Dynamic
REDOR	Rotational Echo Double Resonance

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Figure 1.

(A) Time domain three-pulse ESEEM spectra of Leu10 *i*+3 M2δ in bicelle. The top (red) is ²H-labeled d₁₀ Leu10 *i*+3 and the bottom (black) is the non-deuterated Leu10 *i*+3 control sample. (B) Frequency domain spectra where red illustrates the ²H-labeled d₁₀ Leu10 *i*+3 M2δ sample and black illustrates the control.

Figure 2. Three-pulse ESEEM experimental data with a $\tau=200$ ns of $i+1$ to $i+4$ ^2H -labeled d_{10} Leu10 in lipid bicelle samples. (A) Time domain spectra, (B) Frequency domain spectra.

Figure 3.

Comparison between ^2H -labeled d_8 Val9 and ^2H -labeled d_{10} Leu10 M28 bicelle samples. (A) Normalized FT frequency domain modulation data for ^2H -labeled d_{10} Leu10 $i+4$ (blue) and ^2H -labeled d_8 Val9 $i+4$ (red). (B) Likely conformation of M28 with and ^2H -labeled d_{10} Leu10 (left) and M28 with ^2H -labeled d_8 Val9 (right) both with MTSL at $i+4$ from MD simulation