Published in final edited form as:

Metab Brain Dis. 2018 December ; 33(6): 1975–1984. doi:10.1007/s11011-018-0305-4.

Changes in Biophysical Characteristics of PFN1 Due to Mutation Causing Amyotrophic Lateral Sclerosis

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Abstract

Single amino acid mutations in profilin 1 (PFN1) have been found to cause amyotrophic lateral sclerosis (ALS). Recently, we developed a mouse model for ALS using a PFN1 mutation (glycine 118 to valine, G118V), and we are now interested in understanding how PFN1 becomes toxically lethal with only one amino acid substitution. Therefore, we studied mutation-related changes in the PFN1 protein and hypothesized that such changes significantly disturb its structure. Initially, we expressed and studied the purified PFN1^{WT} and PFN1^{G118V} proteins from bacterial culture. We found that the PFN1^{G118V} protein has a different mean residue ellipticity, as measured by far-UV circular dichroism, accompanied by a spectral shift. The intrinsic fluorescence of PFN1^{G118V} showed a small fluctuation in maximum fluorescence absorption and intensity. Moreover, we examined the time course of PFN1 aggregation using SDS-PAGE, western blotting, and MALDI-TOF/TOF and found that compared with PFN1^{WT}, PFN1^{G118V} had an increased tendency to form aggregates. Dynamic light scattering data confirmed this, showing a larger size distribution for PFN1^{G118V}. Our data explain why PFN1^{G118V} tends to aggregate, a phenotype that may be the basis for its neurotoxicity.

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Author Contributions

M.K. introduced the idea, designed the study, analyzed the data, and wrote the manuscript. A.G. designed the study, analyzed the data, and wrote the manuscript. M.N. designed and performed experiments, analyzed data, and wrote the manuscript. P.G. assisted with MALDI analysis and helped prepare the manuscript. A.A. assisted with the expression of recombinant PFN1, and S.A. assisted with sub-cloning experiments.

Competing interests: Dr. Kiaei and UAMS have a financial interest in the technology discussed in this publication. These financial interests have been reviewed and approved in accordance with the UAMS conflict of interest policies. Other authors have no conflict to disclose.

Keywords

PFN1^{G118V}; PFN1^{WT}; ALS; Aggregation; Actin binding domain

Introduction

Amyotrophic lateral sclerosis (ALS) is a fatal neurodegenerative disease caused by the progressive degeneration of motor neurons in the brain and spinal cord leading to the loss of voluntary movement and muscle control. It is known as Lou Gehrig's disease in the United States and as motor neuron disease in Europe. Two different forms of ALS exist: familial (fALS) and sporadic (sALS) (Zarei et al. 2015). Although the etiology of ALS is largely unknown, inheritable genetic mutations have been found to cause fALS in humans (Hardiman et al. 2017; van Es et al. 2017). One such mutation was found in the gene *PFN1*, which encodes the profilin 1 (PFN1) protein (Wu et al. 2012). Thus far, eight PFN1 mutations (A20T, C71G, T109M, M114T, E117G, G118V, R136W, and Q139L) have been linked to fALS and sALS (Chen et al. 2013; Ingre et al. 2013; Smith et al. 2014; Wu et al. 2012).

Evolutionarily, PFN1 is a highly conserved small protein, with 139 amino acid residues in humans and 140 amino acid residues in rodents. Among the four known profilin isoforms (PFN 1–4), PFN1 is ubiquitously expressed and abundant in most cell and tissue types (Alkam et al. 2017; Witke 2004). The tertiary structure of PFN1 (Fig. 1) consists of seven anti-parallel β -sheets and five α -helices, two of which are located at the termini (Krishnan et al. 2009; Metzler et al. 1995; Schutt et al. 1993). PFN1 is best known for its actin-binding function and is an essential regulator in converting G-actin to F-actin, a critical step in actin polymerization. Furthermore, multiple other functions were discovered by virtue of its interaction with ligands and cellular signaling partners via its poly-L-proline (PLP)-binding domain (Witke 2004). PFN1 has more than 50 ligands and binding partners, at least two of which are disease-causing proteins: survival of motor neuron protein (SMN), which is implicated in spinal muscular atrophy, and valosin-containing protein (VCP), which is implicated in ALS (Fan and Simard 2002; Hensel and Claus 2017; Johnson et al. 2010). The links between PFN1 and SMN and VCP are of great interest, and further studies are needed to confirm whether their interactions regulate the actin cytoskeleton (Hensel and Claus 2017; Nolle et al. 2011).

The recent reports of the work by several groups describe their investigation into the mechanism of mutant PFN1 neurotoxicity in ALS (Boopathy, et al., 2015, Freischmidt et al., 2015, Del Poggetto et al., 2015, Del Poggetto et al., 2016). The mutations in PFN1 are located in the actin binding domain and in the PLP binding domain. The hypothesis that the neurotoxicity of PFN1 due to loss of actin binding and by loss of interaction with PLP containing domain may not be true. Therefore, there may be different mechanism(s) at play, such as gain-of-toxic function. There is strong evidence that specific mutations thermodynamically destabilize PFN1 and also cause structural changes in its native state that affect its propensity for aggregation, as characterized *in vitro* (Boopathy et al. 2015; Del Poggetto et al. 2015a; Del Poggetto et al. 2016; Wu et al. 2012). A more recent study

showed that a C71G mutation causes well-folded PFN1 to co-exist with unfolded PFN1. Unfolded PFN1^{C71G} has accessible amphiphilic and/or hydrophobic regions that might induce a non-native helical conformation (Lim et al. 2017), initiating ALS as a result of interacting abnormally with membranes. On the other hand, the accumulation of TAR DNA-binding protein 43 (TDP-43) aggregates was found to be associated with PFN1 toxicity (Matsukawa et al. 2016; Tanaka and Hasegawa 2016), and this may be an initiating factor in the neuron toxicity and cytoskeletal dysfunction found in ALS.

Of the fALS PFN1 mutations identified, the G118V mutation is located most proximal to the actin-binding site, according to a model generated with PyMol software and published bovine PFN1 X-ray crystallography data (Alkam et al. 2017; Fil et al. 2017). A recent study showed that in a hPFN1^{C71G} mouse model, motor neuron degeneration preceded mutant protein aggregation and caused ALS phenotypes by a gain of toxicity (Yang et al. 2016). However, the mechanism by which PFN1^{G118V} initiates ALS remains unknown. A recently developed hPFN1^{G118V} mouse model showed pathogenic and behavioral phenotypes of ALS disease, including mutant PFN1 aggregation and abnormally low F/G-actin ratios in lumbar spinal cord sections from fully symptomatic and end-stage animals (Fil et al. 2017). Although some mutations can alter actin binding, the G118V mutation causes PFN1 to lose its ability to bind actin (Freischmidt et al. 2015; Wu et al. 2012). It is possible that this amino acid substitution uniquely alters the biophysical properties of PFN1 to produce a conformational change in the binding site, allowing the valine residue to adapt more energetically favorable dihedral angles (Boopathy et al. 2015).

The misfolding and aggregation of PFN1, due to a single amino acid substitution, could be the underlying cause of neurotoxicity and neurodegeneration in human ALS and in the PFN1 transgenic mouse model (Fil et al. 2017). However, the molecular details behind the misfolding and aggregation of PFN1 are not known; therefore, we sought to investigate the changes that lead to this anomaly. Here, we have investigated the biophysical characteristics of PFN1 containing a G118V mutation. We expressed and purified wild type and mutant PFN1 from *E. coli* and analyzed the proteins with common biochemical, biophysical, and analytical techniques (e.g., western blotting, MALDI-TOF/TOF, dynamic light scattering [DLS], far-UV circular dichroism [CD], and fluorescence spectroscopy). Our study identified structural changes in PFN1^{G118V} and provides valuable insight into the mechanism of PFN1 toxicity.

Materials and Methods

Cloning and expression of recombinant PFN1

Plasmid DNA constructs for human PFN1, both wild type (PFN1^{WT}) and the G118V mutation (PFN1^{G118V}), were obtained from NorClone (Canada). PFN1 cDNA was subcloned into pET21a (Novagen). *E. coli* BL21 cells transformed with PFN1 constructs were grown in LB medium (Liofilchem, Italy) containing 100 µg/mL ampicillin at 37°C to an optical density at 600 nm of 0.6 (Fedorov et al. 1994). Subsequently, 1 mM IPTG (SinaClon, Iran) was added to induce PFN1 expression. To allow sufficient protein production, the cells were incubated for either 3 hours at 37°C (wild type) or overnight at 18°C (G118V) before harvesting by centrifugation.

Purification of PFN1 by ion exchange

Bacterial cells were collected and sonicated in 50 mM Tris-HCl and 100 mM NaCl buffer, pH 7.4, containing protease inhibitor (Roche, Germany), and bacterial cell debris was removed by centrifugation (12,000g). The supernatant was subjected to a Diethylaminoethyl (DEAE)-Sephacel Fast Flow anion exchange column (GE Healthcare Bio-Sciences AB, Sweden) pre-equilibrated with 50 mM Tris-HCl pH 7.4 (buffer A) with a KNAUER Azura FPLC system coupled to a UV detector (Germany). The protein sample was loaded onto the column at a flow rate of 0.5 ml/min. The column was washed with buffer A (3 column volumes) at a flow rate of 1 ml/min. Then, a linear gradient was established with buffer A and buffer B (1 M NaCl), and the rest of the bound proteins were eluted by increasing the percentage of buffer B from 0% to 100% during 100 minutes. The UV wavelength was set at 280 nm.

Purification of PFN1 by affinity column

Cyanogen bromide (CNBr)-activated sepharose was prepared based on Kohn's method (Kohn and Wilchek 1984). Poly-L-proline, with molecular weight 1 –10 kDa (Sigma-Aldrich, USA), was dissolved in the coupling buffer (0.1 M NaHCO₃ and 0.5 M NaCl, pH 8.5), and added to CNBr-activated resin. The resin was stirred for 20 hours at 4°C then washed with the coupling buffer and blocked with 0.1 M Tris-HCl (pH 8.0) plus 0.5 M NaCl for 2 hours. Then, the lysate was washed 3 times with 0.1 M Tris buffer (pH 8.0) and 0.1 M acetate buffer (pH 4.0) plus 0.5 M NaCl. The lysate was diluted with wash buffer (10 mM Tris, 100 mM NaCl, 1 mM EDTA, and 2 mM DTT, pH 7.4) and incubated with PLP-sepharose resin, pre-equilibrated with wash buffer, in an ice bath for 2 hours with gentle shaking. PFN1 was eluted from the resin with wash buffer plus 8 M urea and then dialyzed against wash buffer without urea for 48 hours.

The purified proteins were concentrated using Vivaspin® centrifugal concentrators with a 5 kDa molecular-weight cutoff. The total protein concentration was determined with a Pierce BCA Protein Assay Kit (Thermo-Fisher Scientific, USA), according to the manufacturer's protocol.

SDS-PAGE analysis and western blotting

Samples were mixed with SDS-PAGE sample buffer (0.5 M Tris-HCl pH 6.8, 10% SDS, 3 M glycerol, 0.05% bromophenol blue) and separated with 15% tris-glycine SDS-PAGE according to the Laemmli procedure (Laemmli 1970). SDS-PAGE gels were run with or without reducing agent. If added reducing agent (10X, ThermoFisher, Waltham, MA), it was diluted to 0.25% and added to the buffer in the inner chamber of gel tank to maintain the protein samples in a reduced state during electrophoresis. Reducing agent (1%) added to the protein samples prior to loading. The gels were stained with Coomassie Brilliant Blue G-250 or silver nitrate. Separated proteins on the SDS gel were transferred to a PVDF membrane using the Invitrogen iBlot device. The membranes were blocked with 5% skim milk in PBS-T (0.1% Tween 20) buffer for 1 hour, incubated overnight at 4°C with anti-PFN1 primary antibody (Sigma-Aldrich #P7749) diluted in PBS-T 5% milk (1:1000), and rinsed 3 times with PBS-T buffer. Then, proteins were visualized with an HRP-conjugated secondary

antibody (Amersham Corp., USA) diluted in PBS-T 5% milk (1:5000) and detected with the ECL system (Amersham).

Protein identification by mass spectrometry

Intact PFN1 proteins and in-gel digested PFN1 peptides underwent peptide mass fingerprinting with MALDI mass spectrometry (Applied Biosystems 4800 MALDI-TOF/TOF). In-gel trypsin digestion was performed on the protein bands of interest excised from the Coomassie blue-stained gels, as described previously (Shevchenko et al. 2007). The tryptic peptide mixtures were mixed with an equal volume of α -cyano-4-hydroxycinnamic acid matrix solution in 50% acetonitrile containing 0.1% trifluoroacetic acid, spotted (0.7 μ L) onto a MALDI plate, air dried, and analyzed with a MALDI-TOF/TOF operated in reflector positive mode. The monoisotopic peptide mass values of digested PFN1 were searched in the Swiss-Prot and NCBI databases using the MASCOT peptide mass fingerprinting search program (www.matrixscience.com). For intact mass analysis, sinapinic acid was used as a matrix, and equine cytochrome c (average mass 12,362 Da) and apomyoglobin (average mass 16,952 Da) were used as internal and external standards for calibration. All were analyzed with MALDI-TOF/TOF in linear positive mode. Unless otherwise stated, all materials in this experimental section were purchased from Sigma-Aldrich.

Far-UV CD

Far-UV CD (190–260 nm) measurements were carried out on a JASCO J-715 spectropolarimeter (Tokyo, Japan) at a protein concentration of 0.1 mg/ml in 50 mM Tris-HCl (PH 7.4) at 25°C. The blank-subtracted data were then converted to mean residue ellipticity $[\Theta]$. A quartz cuvette with a 1-mm path length was used for analysis.

Intrinsic fluorescence

Intrinsic fluorescence spectra were recorded at a protein concentration of 0.4 mg/ml in 50 mM Tris-HCl (PH 7.4) using the Cary Eclipse spectrofluorometer (Varian Ltd, Spain) and a 10-mm quartz cuvette with excitation at 280 nm and emission in the 280–500 nm range. The slit widths were set at 5 nm.

Dynamic light scattering

A DLS device (Nanophox Sympatech GmbH, Germany) was used to assess the size distribution of proteins in 0.4 mg/ml protein samples in 50 mM Tris-HCl (PH 7.4) at 25°C (refraction index of 1.33 and a viscosity of 0.88 cP in a 10-mm plastic cell).

All samples used in the far-UV CD, intrinsic fluorescence, and DLS experiments were purified by anion-exchange chromatography, and all analyses were performed immediately after centrifugation (15,700g, 10 minutes) and filtration (0.22 μ m).

Aggregation analysis

Purified recombinant PFN1^{WT} and PFN1^{G118V} samples (0.7 mg/ml, purified with anion-exchange chromatography and stored at –20°C) were thawed, centrifuged at 15,700g for 10 minutes, and then filtered (0.22 μ m filters). Samples in aggregation buffer (100 mM Tris-

HCl and 11.2 mM NaCl, pH 7.4) were incubated at 37°C with shaking on an orbital shaker at 200 rpm. Aliquots were removed from each sample over a time course of 96 hours at 0, 4, 24, 48, 72, and 96 hours, and the samples were analyzed by MALDI-TOF/TOF and SDS-PAGE.

Results

Expression, purification, and validation of recombinant PFN1^{WT} and PFN1^{G118V}

Using *E. coli* as a simple and robust system to overexpress recombinant protein, we obtained high quality and reasonable yields of both wild type (PFN1^{WT}) and mutant (PFN1^{G118V}) PFN1 protein. Prominent bands at a molecular weight corresponding to PFN1 (14–15 kDa) were visible after SDS-PAGE of the recombinant protein homogenates suggesting that PFN1 proteins were expressed (Fig. S11). Moreover, the proteins from the excised bands were analyzed with MALDI-TOF/TOF to validate the gel-based identification of PFN1 (Fig. S11). The peptide sequences were matched with the Swiss-Prot database and rated by a scoring system. MALDI-TOF/TOF analysis identified 21 peptides from the complete digestion of PFN1^{WT} (81% sequence coverage) and 11 matched peptides from the partial digestion of PFN1^{G118V} (47% sequence coverage) (Table 1). This bottom-up proteomic analysis resulted in 100% confidence with human PFN1 for both expressed recombinant proteins (Tables S11 and S12).

To purify recombinant PFN1^{WT} and PFN1^{G118V}, we performed PLP-sepharose affinity chromatography with bacterial cell lysates and obtained high quality protein, as indicated by western blots and mass spectrometry peaks. We found that samples of PFN1^{G118V} contained multiple bands (higher molecular weights) by western blotting, suggesting that PFN1^{G118V} oligomerized (Fig. 2a). As shown by western blotting, the presence of higher molecular-weight bands recognized by anti-PFN1 (Fig. 2b, arrows) confirmed that the bands seen with silver staining (Fig. 2a, arrows) were SDS-resistant PFN1 oligomers. The arrows in figure 2b point to two bands that are dimers and trimers. These bands are much more prominent in the PFN1^{G118V} sample; however, the 30–35 kDa band is less prominent in the PFN1^{WT} sample, and the 45 kDa trimer is absent. Furthermore, PFN1^{WT} and PFN1^{G118V} were purified with DEAE-Sepharose Fast Flow anion-exchange resin to avoid the use of urea. As shown in Figure S12, an aliquot of each fraction (labeled 1–4) was subjected to SDS-PAGE, and PFN1 bands (arrows) were processed for sequencing with MALDI-TOF/TOF. Fraction 1 contained most of the PFN1^{WT} or PFN1^{G118V} proteins, while fractions 2–4 contained very little of either protein.

Biophysical characterization of PFN1^{WT} and PFN1^{G118V}

The conformational properties of the purified recombinant PFN1^{WT} and PFN1^{G118V} were assessed with far-UV CD, intrinsic fluorescence, and DLS techniques.

Far-UV CD—The far-UV CD spectrum of PFN1^{WT} revealed a typical α + β structure (Fig. 3a) that is consistent with observations reported by others (Boopathy et al. 2015; Del Poggetto et al. 2015b; Lambrechts et al. 2002). PFN1^{WT} exhibited negative ellipticity over the range of approximately 204–240 nm and positive ellipticity over the range of

approximately 192–204 nm, with a maximum at 195 nm. In contrast, the CD plot of PFN1^{G118V} showed lower values of positive and negative mean residue ellipticity $[\Theta]$ (deg cm² dmol⁻¹). Additionally, this spectrum showed a maximal positive CD signal at 193 nm and a small shift from approximately 204 nm to a lower wavelength (200 nm) corresponding to a “blue shift” compared to PFN1^{WT} (Fig. 3a).

Intrinsic fluorescence—The intrinsic tryptophan fluorescence of PFN1, due to tryptophan residues 3 and 31 in the PLP-binding site located on a solvent-exposed hydrophobic surface, was used to determine structural fluctuations in the protein by comparing the fluorescence intensities and maximum wavelength (McLachlan et al. 2007). As shown in Figure 3b, the intrinsic fluorescence spectrum of PFN1^{WT} showed a peak from 325–350 nm. Although this peak, with a maximum absorption of 335 nm, represented the compact folded structure of PFN1^{WT} in its native form, small changes in maximum fluorescence absorption to approximately 340 nm (a “red shift”) and in fluorescence intensity were apparent in PFN1^{G118V}, indicating that the change from glycine to valine caused this shift (Fig. 3b). This suggests that a structural change within a more flexible hydrophobic surface, although the exposure of the tryptophan indole groups to solvent is similar to the native protein.

Dynamic light scattering—The hydrodynamic diameter of protein samples were measured by DLS, where the intensity of light scattering is proportional to the sixth power of the particle diameter, causing the scattered light from small species to be very small and undetectable (Lorber et al. 2012). Our DLS analysis of PFN1^{WT} showed an average peak for particle sizes of 4–5 nm, which was consistent with the size of a native PFN1 monomer (Fig. 4a). On the other hand, DLS analysis of PFN1^{G118V} showed an average peak for size distribution in the range of 230–320 nm. This rather large hydrodynamic diameter for mutant PFN1^{G118V} (40 to 60 times larger than for PFN1^{WT}) suggested that aggregated proteins were present (Fig. 4b).

Aggregation analysis of PFN1^{WT} and PFN1^{G118V} *in vitro*

To gain further insight into the characteristics and mechanism of PFN1^{G118V} aggregation, we subjected PFN1^{WT} and PFN1^{G118V} to a time-course condition that might influence the aggregation process. As shown in figure 5, MALDI-TOF/TOF and SDS-PAGE assays showed a decrease in soluble PFN1 during the time-course study. At time zero, intact mass analysis of PFN1^{WT} and PFN1^{G118V} showed the most abundant peak to be the PFN1 monomer in both samples (Fig. 5a and 5c). Moreover, we observed a weak signal consistent with the mass-to-charge (m/z) ratio of a PFN1 dimer in these MALDI mass spectra (Fig. 5a and 5c, arrows). When PFN1 protein samples were incubated at 37°C with constant shaking and analyzed with MALDI, we observed a peak for monomeric PFN1^{WT}, consistent with a corresponding band on the gel, suggesting that PFN1^{WT} did not aggregate into larger protein complexes during the 96 hours of incubation as shown in (Fig. 5b). On the other hand, the band corresponding to monomeric PFN1^{G118V} began to diminish after four hours and was undetectable after two days of incubation (Fig. 5d). Likewise, MALDI analysis of samples incubated for more than 48 hours no longer showed the monomeric spectra of PFN1^{G118V} that was present in the baseline samples (data not shown). We also confirmed that

PFN1^{G118V} was present in the sample by re-solubilizing the insoluble fraction (that formed during incubation) in loading buffer with 2-mercaptoethanol (to reduce the samples) and performing SDS-PAGE. Coomassie-blue staining indicated a band corresponding to monomeric PFN1 (~15 kDa). We also noticed that some aggregated PFN1^{G118V} (high molecular-weight protein aggregates) was not solubilized in the presence of reducing agent and did not migrate on the SDS gel, but remained in the stacking gel. As shown in Fig. 5d, a Coomassie-stained band that is likely PFN1^{G118V} was noticeably reduced, and there was no detectable Coomassie-stained band in the 72-hour lane (Fig. 5d).

Discussion and Conclusion

PFN1 has attracted attention because of its significant role in many cellular activities and its contribution to various diseases. Mutations in its actin- and PLP-binding domains (Fig. 1) potentially interfere with its *de novo* activities, resulting in the toxicity associated with disorders like ALS. Here, we expressed, purified, validated, and investigated the physical properties and differences between wild type and mutant recombinant human PFN1. These types of analyses require the appropriate methodology, a sufficient amount of quality protein, and tools like MALDI-TOF/TOF and SDS-PAGE for manipulation and analysis. First, we ensured that we had isolated pure (90–95%) wild type and mutant human PFN1, as confirmed by MALDI-TOF/TOF (Tables 1, SI1, and SI2).

We used appropriate techniques such as DLS, far-UV CD, and intrinsic fluorescence that were able to reveal the molecular dynamics and characteristics of PFN1^{WT} and PFN1^{G118V} proteins, from size distribution profiles to secondary and tertiary structures. Our data on PFN1^{WT} (Fig. 3 and 4a) were consistent with data on the native, folded PFN1 protein published by others (Boopathy et al. 2015; Del Poggetto et al. 2015b). When we analyzed our DLS data, the average hydrodynamic diameter for PFN1^{G118V} was 265 nm. However, there was no evidence of aggregation by PFN1^{WT}, as the average hydrodynamic diameter was less than 5 nm (Fig. 4a and 4b). Furthermore, differences in the far-UV CD spectrum for PFN1^{G118V} (Fig. 3a), along with a red shift in fluorescence and a slight decrease in fluorescence intensity (Fig. 3b), indicated changes in the secondary and tertiary structure of PFN1^{G118V} compared to wild type. These results are in agreement with those previously published by Del Poggetto et al. (2015b), but they differ from those published by Boopathy et al. (2015), who reported that PFN1^{G118V} has a native-like CD spectrum. Given the phi and psi angles for Gly118 according to the Ramachandran plot, the substitution of glycine with valine induces conformational changes that form more energetically favorable dihedral angles. The far-UV CD values for PFN1^{G118V} suggest that the glycine to valine mutation may produce a significant intermolecular force, resulting in an abnormally different secondary structure and affecting the fifth alpha helix. This is consistent with the results of our own *in silico* study (Kiaei et al., 2018, manuscript under review). The small blue shift at approximately 204 nm to 200 nm, where the molar ellipticity is 0, and lower positive and negative ellipticity values for PFN1^{G118V} in the CD spectrum relative to PFN1^{WT} (Fig. 3a) can also be attributed to high molecular-weight species. Likewise, with western blot of purified PFN1^{G118V}, we observed bands corresponding to the molecular weights of PFN1 dimers and trimers (approximately 30 kDa and 45 kDa) (Fig. 2).

In addition, the intrinsic tryptophan fluorescence assay indicated a folded structure for both PFN1^{WT} and PFN1^{G118V}, as the maximum absorption at 350–357 nm indicating solvent-exposed tryptophan residues in the unfolded protein was not observed here (Del Poggetto et al. 2015b). Given the minor red shift in maximum fluorescence presented here and considering that the G118V mutation has a modest effect on the interaction between PFN1 and PLP (Boopathy et al. 2015), we propose that the environment of tryptophan residues is very similar in the mutant and native state. One may speculate that, altogether, the structural changes that occur when PFN1^{G118V} is folded may make the hydrophobic core somewhat solvent accessible.

This type of structural change following a single, critical amino acid substitution may favor the oligomerization and aggregation of PFN1^{G118V}. Consistent with our observations, *in vitro* and *in vivo* studies by others support our evidence for the tendency of PFN1^{G118V} to aggregate (Fil et al. 2017; Smith et al. 2014; Tanaka and Hasegawa 2016; Tanaka et al. 2016; Yang et al. 2016). There are also reports of a correlation between changes in how mutant PFN1 variants fold and their tendency for aggregation (Del Poggetto et al. 2015a; Del Poggetto et al. 2016). Indeed, our evidence that PFN1^{G118V} has a larger hydrodynamic diameter than PFN1^{WT} further supports the conclusion that PFN1^{G118V} forms aggregates (Fig. 4b). Looking back at the challenges we encountered when purifying human PFN1^{G118V} from our bacterial system, the fact that a majority of the mutant protein was recovered from the pellet instead of the supernatant suggests that the protein forms aggregates (data not shown). Additionally, in our time-course study with SDS-PAGE and MALDI-TOF/TOF, we found a significant decrease in the band and peak of soluble PFN1^{G118V} monomer after 24 hours of incubation at 37°C, and we observed the formation of larger insoluble protein complexes that were stable under denaturing conditions (Fig. 5d). However, we observed bands of SDS-soluble, monomeric PFN1^{WT} on the gel after 96 hours of incubation, indicating that PFN1^{WT} is less prone to aggregating (Fig. 5b). The work of other colleagues supports the idea that SDS-resistant oligomers of PFN1^{G118V} (see Fig. 2b) may seed the formation of inclusions and accelerate the recruitment of monomeric PFN1^{G118V} into aggregates (Tanaka et al. 2016). Previous studies demonstrated that the G118V mutation, which is proximal to the actin-binding site (Fig. 1), causes PFN1 to lose its ability to bind actin, sequester G-actin, and regulate the F/G-actin ratio, thus affecting the dynamics of actin polymerization and de-polymerization (Fil et al. 2017; Freischmidt et al. 2015). Our data suggest that the G118V mutation causes structural alterations in PFN1 that may impair its interaction with actin as a consequence of abnormal binding-site conformation and/or PFN1 aggregation.

In conclusion, we investigated the substitution of glycine 118 in PFN1, a solvent-exposed residue within a flexible loop, with valine, a residue with a larger side chain. Our findings support our hypothesis that this mutation significantly alters the secondary and tertiary structures of PFN1. Therefore, the toxicity of mutant PFN1 could result from the placement of two valines side by side, thus changing the structure and leading to misfolding, oligomerization, and aggregation. Our results present a possible mechanism by which mutant PFN1 aggregates, with important implications for the development of drugs that may block this process.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements:

We gratefully acknowledge Dr. Seyed Omid Ranaei Siadat from the Protein Research Center (Shahid Beheshti University) for allowing us to access his laboratory, which focuses on protein expression. Financial support by the Shahid Beheshti University Research Council is gratefully acknowledged. This manuscript was edited by the Science Communication Group at the University of Arkansas for Medical Sciences (UAMS); we also thank Caroline Barham (UAMS) for proofreading the manuscript. The authors acknowledge funding from the UAMS startup fund, UAMS Center for Translational Neurosciences, NIGMS IDeA Program Award P30 GM110702, P20GM109005, and NINDS NS088653 and NS101334.

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Figure 1:

Structure of human PFN1^{WT} generated with PyMol (PDB: 1PFL).

The G118 residue is depicted in pink, and the remaining ALS-linked residues are shown in yellow and labeled in white. Residues involved in actin binding are shown in blue, and PLP-binding domains are in green. Tryptophan residues (W3 and W31) involved in PLP binding are shown in green and labeled in red. The structure on the left shows the actin-binding site, where the G118V mutation located. The structure on the right is tilted to show the PLP-binding area and other features of PFN1.

Figure 2:

Silver-stained SDS-PAGE and western blot of purified PFN1^{WT} and PFN1^{G118V}.

a) Samples purified with PLP-sepharose were subjected to SDS-PAGE and stained with silver nitrate. These gels were run without any reducing agent. The G118V sample contains higher molecular weight bands (arrows), evidence for PFN1 oligomerization. **b)** Western blots of purified wild type (WT) and mutant samples were probed with anti-PFN1, and oligomers are confirmed to be PFN1 with specific antibody that only recognize PFN1. M, pre-stained protein ladder; molecular weights are shown in kDa.

Figure 3:
Structural characterization of PFN1^{WT} and PFN1^{G118V} with different probes.
a) Far-UV circular dichroism spectra and **b)** intrinsic fluorescence spectra of PFN1^{WT} and PFN1^{G118V}.

Figure 4:
Size distribution profile of PFN1^{WT} and PFN1^{G118V}.
Dynamic light scattering graph showing the average size distribution of **a)** PFN1^{WT} and **b)** PFN1^{G118V}.

Figure 5:

Intact mass and time-course aggregation analysis of PFN1^{WT} and PFN1^{G118V}.

MALDI-TOF/TOF spectra of **a**) PFN1^{WT} and **c**) PFN1^{G118V} proteins at time zero before aggregation analysis. SDS-PAGE of aliquots from **b**) PFN1^{WT} or **d**) PFN1^{G118V} samples incubated at 37°C under constant shaking for 0, 24, 48, or 72 hours. Insoluble (Ins) fraction from PFN1^{G118V} sample solubilized in sample buffer containing 2-mercapthoethanol; arrowhead pointing to the Coomassie-stained band in the top of the Ins lane (d) indicates PFN1 aggregates. Other fractions were solubilized in sample buffer without reducing agent.

Table 1

Peptide size, number, and sequence coverage.

Protein	PFN1 ^{wt}	PFN1 ^{G118V}
No. of peptides	21 (4–20 amino acid residues)	11 (4–20 amino acid residues)
Sequence coverage	81%	47%