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Shortening heparan sulfate chains prolongs survival and reduces parenchymal plaques in prion disease caused by mobile, ADAM10-cleaved prions

Patricia Aguilar-Calvo[1] · Alejandro M. Sevillano[1] · Jaidev Bapat[1] · Katrin Soldau[1] · Daniel R. Sandoval[2] ·
Hermann C. Altmeppen[3] · Luise Linsenmeier[3] · Donald P. Pizzo[1] · Michael D. Geschwind[4] · Henry Sanchez[4] ·
Brian S. Appleby[5,6] · Mark L. Cohen[5,6] · Jiri G. Safar[5,6] · Steven D. Edland[7,8] · Markus Glatzel[3] · K. Peter R. Nilsson[9] ·
Jeffrey D. Esko[2] · Christina J. Sigurdson[1,10,11]

Abstract

Cofactors are essential for driving recombinant prion protein into pathogenic conformers. Polyanions promote prion aggregation in vitro, yet the cofactors that modulate prion assembly in vivo remain largely unknown. Here we report that the endogenous glycosaminoglycan, heparan sulfate (HS), impacts prion propagation kinetics and deposition sites in the brain. Exostosin-1 haploinsufficient ($Ext1^{+/-}$) mice, which produce short HS chains, show a prolonged survival and a redistribution of plaques from the parenchyma to vessels when infected with fibrillar prions, and a modest delay when infected with subfibrillar prions. Notably, the fibrillar, plaque-forming prions are composed of ADAM10-cleaved prion protein lacking a glycosylphosphatidylinositol anchor, indicating that these prions are mobile and assemble extracellularly. By analyzing the prion-bound HS using liquid chromatography–mass spectrometry (LC–MS), we identified the disaccharide signature of HS differentially bound to fibrillar compared to subfibrillar prions, and found approximately 20-fold more HS bound to the fibrils. Finally, LC–MS of prion-bound HS from human patients with familial and sporadic prion disease also showed distinct HS signatures and higher HS levels associated with fibrillar prions. This study provides the first in vivo evidence of an endogenous cofactor that accelerates prion disease progression and enhances parenchymal deposition of ADAM10-cleaved, mobile prions.

Keywords Amyloid · Neurodegeneration · Glycosaminoglycans · ADAM10 cleavage

✉ Christina J. Sigurdson
  csigurdson@ucsd.edu

[1] Department of Pathology, University of California, San Diego (UCSD), 9500 Gilman Dr., La Jolla, CA 92093, USA

[2] Department of Cellular and Molecular Medicine, University of California, San Diego (UCSD), La Jolla, CA, USA

[3] Institute of Neuropathology, University Medical Center Hamburg-Eppendorf (UKE), Hamburg, Germany

[4] Department of Neurology, Memory and Aging Center, University of California, San Francisco (UCSF), San Francisco, CA, USA

[5] Departments of Pathology and Neurology, Case Western Reserve University, Cleveland, OH, USA

[6] National Prion Disease Pathology Surveillance Center, Case Western Reserve University, Cleveland, OH, USA

[7] Department of Family Medicine and Public Health, University of California, San Diego (UCSD), La Jolla, CA, USA

[8] Department of Neurosciences, University of California, San Diego (UCSD), La Jolla, CA, USA

[9] Department of Physics, Chemistry, and Biology, Linköping University, Linköping, Sweden

[10] Department of Medicine, University of California, San Diego (UCSD), La Jolla, CA, USA

[11] Department of Pathology, Immunology, and Microbiology, University of California, Davis (UCD), Davis, CA, USA

Introduction

More than 35 proteins aggregate into pathogenic fibrils in systemic and neurodegenerative disorders, including diabetes and Alzheimer's disease [22]. The fibril assembly is accelerated in vitro by diverse anionic polymers, such as RNA, heparin, and heparan sulfate (HS) [14, 16, 23], raising the question of whether endogenous polymers affect disease progression in vivo. Heparin and HS are glycosaminoglycans (GAGs), long polysaccharide chains containing repeating disaccharide units assembled on extracellular and membrane-bound proteins. HS is structurally similar to the anticoagulant heparin, and has been found associated with nearly all amyloids [5, 61, 66, 69, 97, 98, 100]. The specificity of HS–protein binding is profoundly affected by HS chain length as well as by the density and pattern of sulfation [62, 105, 114], which varies among cell types [93].

Prion diseases are rapidly progressive neurodegenerative disorders [86] caused by the misfolding of the cell surface-expressed prion protein monomer, PrP^C, into a β-sheet-rich multimer, PrP^{Sc} [18, 20, 76, 87]. PrP^{Sc} can be arranged as distinct conformers, or strains, having different biochemical properties, cellular targets, and varying abilities to spread into the central nervous system (CNS) [8–10, 13]. For example, fibrillar prions spread poorly from extraneural entry sites into the CNS, whereas subfibrillar prions, which do not form plaques or ultrastructurally visible fibrils in situ, spread efficiently into the CNS [10–12, 52]. Yet the molecular mechanisms that govern prion assembly into fibrils or subfibrils and that determine how structure is linked to the capacity to spread are unknown.

Similar to other amyloidogenic proteins, prions bind GAGs, and in vitro studies have revealed that GAGs promote prion conversion and fibril assembly [6, 46, 113] and enhance prion uptake [40–42], but can also PrP^C [109]. Shortening or de-sulfating HS chains abrogated the acceleration in fibril assembly and enhanced uptake of prions [46], supporting a role for HS in prion conversion. Additionally, HS co-localizes with prion plaques in brain sections from patients and prion-infected mice [66, 96], which, together with the in vitro findings, suggest HS may impact prion aggregation, cell targeting, and spread in vivo.

Paradoxically, GAG mimetics delay prion disease in animals [1, 26, 28, 29, 56, 57], potentially due to stabilization of PrP^C [109] or competition with endogenous GAGs [17, 19]. These encouraging initial reports led to the treatment of five variant Creutzfeldt–Jakob disease (vCJD) patients with intraventricular pentosan polysulfate; four of five patients showed an extended survival as compared to the mean for vCJD. While the results were promising, a treatment effect due to differences in patient care could not be excluded [68, 80, 107].

To determine how HS modifications impact prion spread through the brain, here we altered HS chain length and tested the impact on prion assembly and replication in vivo. Using $Ext1^{+/-}$ mice that express shorter HS chains due to a reduction in exostosin-1 (Ext1) [60], the co-polymerase that assembles the HS chains [72], we assessed the spread of three conformationally diverse prion strains. $Ext1$ haploinsufficiency did not alter the conformation of any prion strain. Instead, we found evidence that HS binds ADAM10-cleaved, fibrillar prions, facilitates parenchymal plaque formation, and accelerates disease, yet only minimally binds to GPI-anchored, subfibrillar prions. Using mass spectrometry and immunohistochemistry, we found high levels of HS bound to fibrillar prions in mouse and human brain, and localized HS to the plaque core. Collectively, our results elucidate how HS impacts parenchymal prion plaque distribution in a strain-specific manner, and indicate a rationale for selectively targeting the plaque-forming prions for treatment trials designed to disrupt the HS–prion interface.

Methods

Prion transmission experiments in transgenic mice

$Ext1^{+/-}$ mice [60] were bred to wild-type mice (C57BL/6) or tga20 mice, which overexpress mouse PrP^C [31]. Mice were maintained under specific pathogen-free conditions on a 12:12 light/dark cycle. Groups of 5–16 male and female $Ext1^{+/-}$ and $Ext1^{+/+}$ littermate control mice (6–8 weeks old) were anesthetized with ketamine and xylazine and inoculated into the left parietal cortex with 30 μl of 1% prion-infected brain homogenate prepared from terminally ill mice.

Prion-inoculated mice were monitored three times weekly for the development of terminal prion disease, including ataxia, kyphosis, stiff tail, hind leg clasp, and hind leg paresis, and were then euthanized. The brain was halved, and one hemisphere was immediately fixed in formalin. Fixed brains were treated for 1 h in 96% formic acid, post-fixed in formalin, cut into 2 mm transverse sections, and paraffin embedded for histological analysis. A 2–3 mm transverse section was removed from the remaining hemisphere at the level of the hippocampus/thalamus, embedded in optimal cutting temperature (OCT) compound, and immediately frozen on dry ice. The remaining brain tissue was frozen for biochemical studies. Survival time was calculated from the day of inoculation to the day of terminal clinical disease.

Histopathology and immunohistochemical stains

Four micron sections were cut onto positively charged silanized glass slides and stained with hematoxylin and eosin (HE), or immunostained using antibodies for PrP [SAF84,

epitope in the globular domain, amino acids 160–170 of mouse PrP (Cayman Chemical); 12F10, epitope in the globular domain, amino acids 144–152 of human PrP (Cayman Chemical); sPrP^{G228}, epitope in the proteolytically cleaved C-terminus, including amino acid 228G of mouse PrP (generated in the laboratory of co-author MG) [63])], as well as astrocytes (glial fibrillary acidic protein, GFAP), microglia (Iba1), and endothelial cells (CD31). For all mouse and most human PrP^{Sc} immunolabeling, sections were deparaffinized and incubated for 5 min in 96% formic acid, then washed in water for 5 min, treated with 5 µg/ml of proteinase-K (PK) for 7 min to digest PrP^C, and washed in water for 7 min. Sections were then placed in citrate buffer (pH 6), heated in a pressure cooker for 20 min, cooled for 5 min, and washed in distilled water. Sections were blocked and incubated with anti-PrP antibodies SAF84, sPrP^{G228} (mouse brain sections), or 12F10 (human brain sections, sCJD and GSS-F198S cases) for 45 min, followed by anti-mouse or anti-rabbit biotin (Jackson Immunolabs) for 30 min, and then streptavidin–HRP (Jackson Immunoresearch) for 45 min. Slides were then incubated with DAB reagent (Thermo Scientific) for 15 min and counterstained with hematoxylin. For the GSS-P102L case, brain sections were placed in 1.5 mM HCl in a microwave for 15 min, blocked and incubated with 3F4 (epitope in amino terminus of PrP, amino acids 109–112 of human PrP, [49]) for 1 h, incubated with enzyme-conjugated polymer DAKO Envision for 30 min, and then DAKO DAB reagent for 1 min prior to counterstaining with hematoxylin. The immunostaining was performed on a Lab Vision 480 Autostainer.

GFAP immunohistochemistry for astrocytes (1:6000; DAKO) was performed on an automated tissue immunostainer (Ventana Discovery Ultra, Ventana Medical Systems, Inc) with a protease antigen retrieval (P2, Ventana) for 16 min.

For the PrP^{Sc} and CD31 (endothelial cells) dual immunolabeling, tissue sections were stained sequentially using anti-PrP SAF84 (1:250) and CD31 antibodies (1:150; Dianova) using the tyramide signal amplification system (TSA; Thermo Fisher). Slides were stained on a Ventana Discovery Ultra (Ventana Medical Systems, Tucson, AZ, USA). Antigen retrieval was performed using a slightly basic treatment solution (CC1; pH 8.5, Ventana) for 92 min at 95 °C. Sections then were incubated in anti-PrP antibody for 32 min at 37 °C, followed by anti-mouse-HRP (UltraMap Detection Kit, Ventana) and TSA-Alexa 594. The antibodies were denatured by treatment in a citric acid-based solution, pH 6 (CC2, Ventana) for 24 min at 95 °C. Subsequently, the slides were incubated with anti-CD31 antibody (rat) for 32 min at 37 °C followed by rabbit anti-rat (1:500; Jackson ImmunoResearch) and detected using the OmniMap system (Ventana) to fluorescently label the CD31-expressing cells with TSA-Alexa 488.

For the PrP and HS dual immunolabeling, tissue sections were deparaffinized and epitopes were exposed by sequential treatment with formic acid, PK, and heated citrate buffer (pH 6) as described above for PrP staining. Sections were blocked and incubated with anti-PrP [SAF-84 (1:250) or 12F10 (1:400)] and anti-HS [10E4 (1:200; AMS Bioscience)] antibodies for 45 min followed by anti-mouse IgM biotin (1:500; Jackson Immunolabs) for 30 min and streptavidin–HRP (1:2000; Jackson Immunoresearch) for 45 min, and then incubated with tyramide-Alexa488 (Invitrogen) for 10 min. Finally, sections were incubated with anti-mouse IgG–CY3 (1:200; Jackson Immunolabs), nuclei were labeled with DAPI, and slides were mounted with fluorescent mounting medium (Dako). As controls for the HS stain, a subset of duplicate slides were treated with 8 milli-units of heparin lyases I, II, and III for 1 h prior to immunostaining. Isotype immunoglobulin controls, single sections immunostained for PrP or HS, and prion-negative cases were also included.

For quantitative analysis of microglial inflammation in tga20$^{+/-}Ext1^{+/-}$ and tga20$^{+/-}Ext1^{+/+}$ mice, brain regions containing cerebral cortex, hippocampus, thalamus, hypothalamus, and cerebellum were imaged (approximately 6 fields per mouse) using the Olympus EX41 microscope with DP Controller. Images were converted to grayscale and FIJI (an ImageJ-based image processing software) was used to measure the total brain area and quantify microglia using the "Measure" function. Activated microglia were demarcated using the "Find the edges" function and particle analysis was used to measure the area occupied by microglia. The total area covered by microglia was divided by the total area in each brain region.

For the Alcian blue stain, brain sections were deparaffinized and incubated for 3 min in 3% acetic acid, then stained in Alcian blue solution in the microwave for 30 s, cooled for 40 min, washed in distilled water, and counterstained in nuclear fast red solution for 5 min. For the Congo red staining, slides were deparaffinized, fixed in 70% ethanol for 10 min, immersed in an alkaline solution, and then stained with Congo red solution for 20 min.

Histopathologic lesion profiles

Two investigators blinded to animal identification performed the histological analyses. Brain lesions from $Ext1^{+/-}$ and $Ext1^{+/+}$ prion-infected mice were scored for the level of spongiosis, gliosis, and PrP immunological reactivity on a scale of 0–3 (0 = not detectable, 1 = mild, 2 = moderate, 3 = severe) in nine regions including gray and white matter: (1) dorsal medulla, (2) cerebellum, (3) hypothalamus, (4) medial thalamus, (5) hippocampus, (6) septum, (7) medial cerebral cortex dorsal to hippocampus, (8) cerebral peduncle, and (9) cerebellar peduncle. A sum of the three scores

resulted in the value obtained for the lesion profile for the individual animal and was depicted in the 'radar plots'.

Western blot and glycoprofile analyses

PrP^{Sc} was concentrated from 10% brain homogenate in phosphate-buffered saline (PBS) (*w/v*) by performing sodium phosphotungstic acid precipitation prior to western blotting [110]. Briefly, 20 µl of 10% brain homogenate in an equal volume of 4% sarkosyl in PBS was digested with benzonase™ (Sigma-Aldrich) followed by treatment with 20 µg/ml PK at 37 °C for 30 min. After addition of 4% sodium phosphotungstic acid in 170 mM MgCl$_2$ and protease inhibitors (Complete TM, Roche), extracts were incubated at 37 °C for 30 min and centrifuged at 18,000×*g* for 30 min at 25 °C. Pellets were resuspended in 2% sarkosyl prior to electrophoresis and immunoblotting. Samples were electrophoresed through a 10% NuPage Bis–Tris gel (Thermo Fisher Scientific) and transferred to nitrocellulose by wet blotting. Membranes were incubated with monoclonal antibody POM19 [discontinuous epitope at C-terminal domain, amino acids 201–225 of the mouse PrP [85]] or polyclonal antibody sPrP^{G228} followed by incubation with an HRP-conjugated IgG secondary antibody. The blots were developed using a chemiluminescent substrate (Supersignal West Dura ECL, Thermo Fisher Scientific) and visualized on a Fuji LAS 4000 imager. Quantification of PrP^{Sc} glycoforms was performed using Multigauge V3 software (Fujifilm). For mCWD, 100 µl of 10% brain homogenate was digested with 100 µg/ml PK prior to electrophoresis through a 10% NuPage Bis–Tris gel.

For quantifying the level of PrP^C expression in uninfected mice and PrP^{Sc} in prion-infected mice, the total protein concentration was measured in brain homogenates by bicinchoninic acid assay (Pierce) and equivalent protein levels were immunoblotted as described. Membranes were also probed for actin (Genetex) (PrP^C analysis only).

Conformation stability assay

Prion strain stability in guanidine hydrochloride (GdnHCl) was measured as previously described [82]. In brief, 10% brain homogenates in PBS (w/v) were denatured for 1 h in increasing concentrations of GdnHCl from 0 to 6 M. Samples were then diluted in a Tris-based lysis buffer (10 mM Tris–HCl, 150 mM NaCl, 10 mM EDTA, 2% sarkosyl, pH 7.5) to 0.15 M GdnHCl and digested with PK at a ratio of 1:500 (1 µg PK: 500 µg total protein) for 1 h at 37 °C. The digestion was stopped with 2 mM phenylmethylsulfonyl fluoride (PMSF) and protease inhibitors (Complete TM, Roche), followed by centrifugation at 18,000×*g* for 1 h. The pellets were washed in 0.1 M NaHCO$_3$ (pH 9.8) and centrifuged at 18,000×*g* for 20 min. The pellets were then denatured in 6 M guanidine isothiocyanate (GdnSCN), diluted with 0.1 M NaHCO$_3$, and coated passively onto an ELISA plate. PrP was detected with biotinylated-POM1 antibody (epitope in the globular domain, amino acids 121–231 of the mouse PrP [85]), a streptavidin–HRP-conjugated secondary antibody, and a chemiluminescent substrate. The PrP stability was measured in a minimum of three independent experiments for each strain, always comparing *Ext1*$^{+/-}$ with *Ext1*$^{+/+}$ (3–4 mice per strain).

h-FTAA staining and fluorescence lifetime imaging (FLIM)

Sections (10 µm) of OCT-embedded brain samples were cut onto positively charged silanized glass slides, dried for 1 h, and fixed in 100% then 70% ethanol for 10 min each. After washing with deionized water, sections were equilibrated in PBS, pH 7.4, for 10 min. Heptamer-formyl thiophene acetic acid (h-FTAA) was diluted in PBS to a final concentration of 1.5 µM and added to the sections. The sections were incubated with h-FTAA for 30 min at room temperature, washed with PBS, and mounted using Dako fluorescence mounting medium. The fluorescence decay of h-FTAA bound to PrP aggregates was collected using an inverted Zeiss (Axio Observer.Z1) LSM 780 microscope (Carl Zeiss MicroImaging GmbH) equipped with a modular FLIM system from Becker and Hickl. In this setup, the emitted photons were routed through the direct coupling confocal port of the Zeiss LSM 780 scanning unit and detected by a Becker and Hickl HPM-100-40 hybrid detector. Data were recorded by a Becker and Hickl Simple-Tau 152 system (SPC-150 TCSPC FLIM module) with the instrument recording software SPCM version 9.42 in the FIFO image mode, 256×256 pixels, using 256 time channels (Becker and Hickl GmbH). For all acquisitions, a T80R20 main beam splitter was used and the pinhole was set to 20.2 µm. Scanning area was set to 235.7 µm×235.7 µm, with a scanning resolution of 512×512 pixels. A Plan-Apochromat 40×/1.3 Oil DIC objective lens was used and a 510 nm longpass filter was positioned in front of the hybrid detector. Excitation utilized the 490 nm laser line from the pulsed tunable In Tune laser (Carl Zeiss MicroImaging GmbH) with a repetition rate of 40 MHz. Data was subsequently analyzed in SPCImage version 3.9.4 (Becker and Hickl GmbH), fitting each of the acquired decay curves to a tri-exponential function and color-coded images, as well as distribution histograms, showing the intensity-weighted mean lifetimes generated with the same software. The procedure of staining and FLIM imaging protein aggregates with h-FTAA is described in detail in Ref. [71].

Purification of PrP^Sc to quantify and characterize the bound heparan sulfate

Samples were purified from mouse (whole brain) and human brain (cerebellum: GSS-F198S, sCJD MM1 and sCJD MV1, frontal cortex: GSS-P102L and sCJD MM2, and occipital cortex: sCJD MM1) as previously described [89] with minor modifications. One ml of 10% brain homogenate in PBS (w/v) was mixed with an equal volume of TEN(D) buffer (5% sarkosyl in 50 mM Tris–HCl, 5 mM EDTA, 665 mM NaCl, 0.2 mM dithiothreitol, pH 8.0), containing complete TM protease inhibitors (Roche). Samples were incubated on ice for 1 h and centrifuged at 18,000×g for 30 min at 4 °C. All but 100 µl of supernatant was removed, and the pellet was resuspended in 100 µl of residual supernatant and diluted to 1 ml with 10% sarkosyl TEN(D). Each supernatant and pellet was incubated for 30 min on ice and then centrifuged at 18,000×g for 30 min at 4 °C. Supernatants were recovered while pellets were held on ice. Supernatants were added separately into ultracentrifuge tubes with 10% sarkosyl TEN(D) buffer containing protease inhibitors and centrifuged at 150,000×g for 2.5 h at 4 °C. Supernatants were discarded, while pellets were rinsed with 100 µl of 10% NaCl in TEN(D) buffer with 1% sulfobetaine (SB 3–14) and protease inhibitors and then combined and centrifuged at 225,000×g for 2 h at 20 °C. The supernatant was discarded and the pellet was washed and then resuspended in ice cold TMS buffer containing protease inhibitors (10 mM Tris–HCl, 5 mM MgCl$_2$, 100 mM NaCl, pH 7.0). Samples were incubated on ice overnight at 4 °C. Samples were then incubated with 25 units/ml benzonase™ (Sigma-Aldrich) and 50 mM MgCl$_2$ for 30 min at 37 °C, followed by a digestion with 10 µg/ml PK for 1 h at 37 °C. PK digestion was stopped by incubating samples with 2 mM PMSF on ice for 15 min. Samples were incubated with 20 mM EDTA for 15 min at 37 °C. An equal volume of 20% NaCl was added to all tubes followed by an equal volume of 2% SB 3–14 buffer. For the sucrose gradient, a layer of 0.5 M sucrose, 100 mM NaCl, 10 mM Tris, and 0.5% SB 3–14, pH 7.4, was added to ultracentrifuge tubes. Samples were then carefully transferred and the tubes topped with TMS buffer. Samples were centrifuged at 200,000×g for 2 h at 20 °C. The pellet was rinsed with 0.5% SB 3–14 in PBS. Pellets were resuspended in 50 µl of 0.5% SB 3–14 in PBS and stored at − 80 °C. Gel electrophoresis and silver staining were performed to confirm the purity of the PrP^Sc. To quantify PrP levels, samples were compared against a dilution series of recombinant PrP by immunoblotting and probing with POM19 and 3F4 anti-PrP antibodies.

Heparan sulfate purification and analysis by mass spectrometry

The purified PrP^Sc samples were digested with 0.5 M NaOH (final concentration) on ice for 16 h at 4 °C, neutralized with 0.5 M acetic acid (final concentration), and digested with pronase for 25 h at 37 °C. HS was next extracted by anion exchange chromatography using diethyl-aminoethyl (DEAE) Sepharose columns (Healthcare Life Sciences). For depolymerization, HS was extensively digested with 1 milliunit each of heparinases I, II, and III. The disaccharides resulting from enzymatic depolymerization were tagged by reductive amination with [^{12}C$_6$]aniline [58, 59]. The [^{12}C$_6$] aniline-tagged disaccharides were mixed with [^{13}C$_6$]aniline-tagged disaccharide standards. Samples were analyzed by liquid chromatography–mass spectrometry (LC–MS) using an LTQ Orbitrap Discovery electrospray ionization mass spectrometer (Thermo Fisher Scientific). Internal disaccharides were identified based on their unique mass and quantified relative to the wet weight of tissue [58, 59]. The level of measured HS relative to the PrP^Sc level (measured by western blotting with a recombinant PrP dilution series) was calculated and plotted.

Human patients with sporadic or familial prion disease

Brain samples from patients with sporadic Creutzfeldt–Jakob disease (sCJD) or Gerstmann–Sträussler–Scheinker disease (GSS) (Table 1) were used to investigate how the level of HS molecules bound to PrP^Sc correlated with the histopathologic lesions. The sCJD brain samples originated from patients referred to the UC San Francisco (UCSF) Memory and Aging Center for rapidly progressive neurologic disease. All patients had extensive clinical testing, including brain MRI as well as CSF analysis for 14-3-3, neuron-specific enolase (NSE), and total tau. All were classified as probable sCJD by UCSF clinical and radiological diagnostic criteria [33, 102]. The mean age of sCJD patients at disease onset was 59 ± 5 years (mean ± SD) and the duration of clinical neurologic signs ranged from 1.5 to 24 months (10 ± 9 months).

The GSS brain samples were received and characterized at the National Prion Disease Pathology Surveillance Center (NPDPSC). The mean age of GSS-F198S patients at disease onset was 57 ± 8 years. The duration of clinical neurologic signs markedly varied from 63 to 120 months (86 ± 25 months). Of the familial prion disease cases, the GSS-P102L patient showed the earliest age of disease onset (22 years) and the shortest disease duration (23 months).

For all patients, the *PRNP* open reading frame was sequenced to define the genotype at polymorphic codon 129 (methionine or valine) and to test for mutations in the PrP sequence (performed at NPDPSC). Genomic DNA was extracted from frozen brain tissue samples using Qiagen QIAamp DNA minikit (Qiagen, Gaithersburg, MD) according to the manufacturer's protocol, and a 760-bp fragment corresponding to the human PrP gene (residues 5–258) was amplified by PCR using primers HRM-F (5′-TATGTG
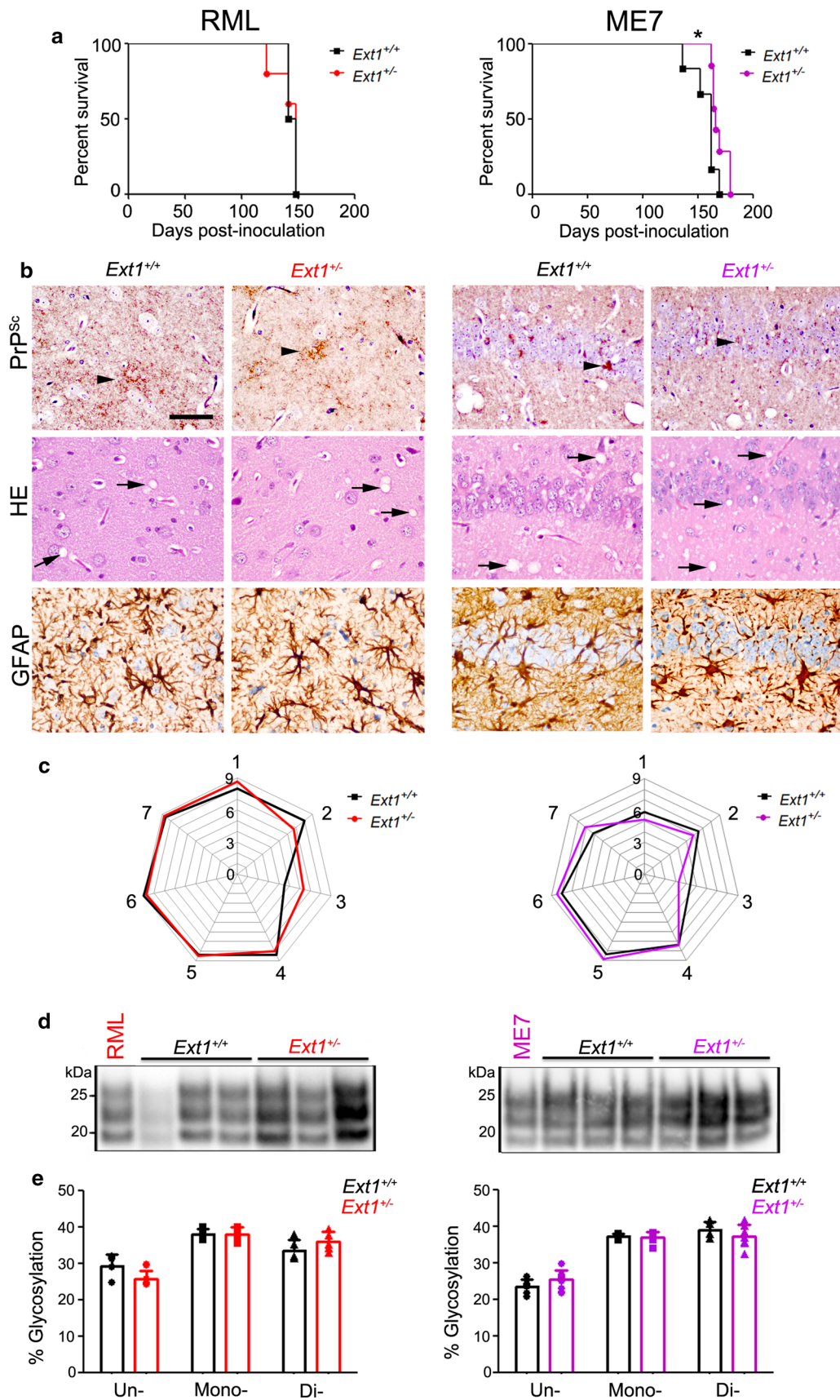
◀**Fig. 1** *Ext1*$^{+/+}$ and *Ext1*$^{+/−}$ mice infected with subfibrillar prions show similar survival times, brain lesions, and biochemical properties of PrP^{Sc}. **a** A comparison of survival times revealed no differences for RML-infected *Ext1*$^{+/−}$ mice and a modest delay in ME7-infected *Ext1*$^{+/−}$ mice as compared to the *Ext1*$^{+/+}$ mice. **b** Brain sections immunolabeled for PrP and GFAP, or stained with hematoxylin and eosin (HE), show indistinguishable prion aggregate distribution and morphology (arrowheads), spongiform degeneration (arrows), and astrogliosis in *Ext1*$^{+/+}$ and *Ext1*$^{+/−}$ brains. **c** Lesion profiles of RML- and ME7-infected *Ext1*$^{+/+}$ and *Ext1*$^{+/−}$ mice (1-dorsal medulla, 2-cerebellum, 3-hypothalamus, 4-medial thalamus, 5-hippocampus, 6-septum, and 7-cerebral cortex) are almost superimposable. **d** Electrophoretic mobility and **e** glycoprofiles of RML and ME7 strains in *Ext1*$^{+/+}$ and *Ext1*$^{+/−}$ mice. The RML and ME7 inocula were loaded for comparison ([first lane 9 (d)]). *$P < 0.05$, Log-rank (Mantel–Cox) test (**a**). Cerebral cortex (RML) and hippocampus (ME7) shown in (**b**). Scale bar = 50 μm (**b**). RML: $n = 6$–7 mice/group; ME7: $n = 4$–5 mice/group

GACTGATGTCGGCCTCTGCAAGAAGCGC-3′) and HRM-R (5′-CCACCTCAATTGAAAGGGCTGCAGGTG GATAC-3′) with defined cycling conditions [39, 53]. The Met/Val polymorphism at codon 129 and mutation of the *PRNP* coding region were determined by deep (63) or direct Sanger sequencing as previously described [53, 79]. Nucleotide sequences from both deep and Sanger sequencing were analyzed using DNAStar Lasergene Software Suite v.7.1.0 (Madison, WI).

The sCJD patients included 129 MM (2), MV (3), and VV (1). The four GSS-F198S patients had a missense mutation at *PRNP* codon 198 (F198S mutation) and were 129 MV (3) or VV (1). The GSS-P102L case had a missense mutation at *PRNP* codon 102 (P102L) and was 129 MV (Table 1).

A diagnosis of prion disease was confirmed for all cases by detection of PK-resistant PrP on immunoblots of brain samples. Characterization of sCJD prion subtype was based on PrP^{Sc} electrophoretic mobility and *PRNP* genotype. All four GSS-F198S patient brain samples showed 8 kDa and higher molecular weight PrP bands by immunoblot, consistent with the published N- and C-terminally truncated PrP fragment ($_{74}$GQPHGGGWGQPHGGGWGQGGGTH-SQWNKP$_{102}$) [83] and multicentric plaques histologically, whereas the GSS-P102L case showed a ≈ 21 kDa (type 1) PrP band and no lower molecular weight bands, as well as diffuse, synaptic PrP deposits in the cerebral cortex, and multicentric plaques in the cerebellum, similar to a subset of GSS-P102L cases previously published [78].

Statistics

Log-rank (Mantel–Cox) tests were performed to assess survival differences between groups. A Student's *t* test (two-tailed, unpaired) was used to determine the statistical significance between the *Ext1*$^{+/−}$ and *Ext1*$^{+/+}$ mouse groups for the PrP^C expression level, lesion profiles, activated microglia,

PrP^{Sc} glycoprofiles, PrP^{Sc} conformation stability, and PrP^{Sc} fibril structure (FLIM). A non-parametric Fisher's exact test was used to compare the proportion of *Ext1*$^{+/−}$ versus *Ext1*$^{+/+}$ mice having plaques by brain region. A two-way ANOVA with Bonferroni's post-test was used to compare the number of parenchymal, vascular, and periventricular plaques in *Ext1*$^{+/−}$ versus *Ext1*$^{+/+}$ mice, as well as the HS composition associated with different prion strains. A non-parametric Spearman correlation test was used to test the correlation of plaque numbers in the corpus callosum versus velum interpositum within a mouse group (*Ext1*$^{+/−}$ or *Ext1*$^{+/+}$). The statistical significance in the level of HS bound to PrP and the level of ADAM10-cleaved PrP among prion strains was determined by performing a non-parametric, two-tailed Wilcoxon rank sum test (when 2 strains were compared) or a one-way ANOVA with Tukey's post test (when three or more strains were compared). For all analyses, $p < 0.05$ was considered significant.

The Wilcoxon test is an exact test, meaning the minimum observable *p* value is constrained when the sample size is small. Notably, there were two comparisons (Figs. 4f and 5d) where there was a complete separation of values, but nonetheless the exact test *p* value did not reach statistical significance. In this circumstance, the lack of formal statistical significance should not be interpreted as a null finding.

Study approval

All animal studies were performed following procedures to minimize suffering and were approved by the Institutional Animal Care and Use Committee at UC San Diego. Protocols were performed in strict accordance with good animal practices, as described in the Guide for the Use and Care of Laboratory Animals published by the National Institutes of Health.

This study was approved by the Human Research Protection Committees at UC San Francisco and Case Western Reserve University (IRB study numbers 10-04905 and 03-14-28, respectively). All brain tissues utilized were deidentified samples collected at autopsy (NIH Office of Human Subjects Research Protections, exemption 4).

Results

HS proteoglycans (HSPG) promote prion internalization and replication in vitro [46, 113]. To determine how HS modulates prion infection in vivo, we challenged *Ext1*$^{+/−}$ [60] and *Ext1*$^{+/+}$ (WT) littermate control mice with prions. Groups of mice were inoculated intracerebrally with three mouse-adapted prion strains, RML, ME7, and mCWD, or uninfected control brain (mock). mCWD prion infection is prolonged in WT mice (approximately 550 days);

therefore, *Ext1*^{+/−} mice were bred to *tga*20 mice, which express four–six-fold higher levels of mouse PrP[C] [31], and the F1 generation (*tga*20[+/−];*Ext1*^{+/−} and *tga*20[+/−];*Ext1*^{+/+} mice) was inoculated with mCWD. The PrP[C] expression in the brain was similar for *Ext1*^{+/−} and *Ext1*^{+/+} mice as well as for *tga*20[+/−];*Ext1*^{+/−} and *tga*20[+/−];*Ext1*^{+/+} mice (Online Resource 1).

Prion disease phenotype triggered by subfibrillar strains is minimally altered in mice expressing short HS chains

Ext1^{+/−} and *Ext1*^{+/+} mice exposed to RML subfibrillar prions showed no difference in the survival time [141 ± 11 versus 145 ± 4 days post-inoculation (dpi), respectively] (Fig. 1a); however, *Ext1*^{+/−} mice exposed to ME7 prions showed a modest but significantly prolonged survival [169 ± 7 dpi versus 157 ± 12 dpi (*Ext1*^{+/−} and *Ext1*^{+/+} mice, respectively)] (Fig. 1a). In comparing the two subfibrillar strains histologically, we found that RML prions induce fine to punctate 1–5 μm aggregates and ME7 prions induce punctate and small, plaque-like deposits, both accompanied by spongiform change and astrocytic gliosis (Fig. 1b). The prion-infected *Ext1*^{+/−} and *Ext1*^{+/+} mice showed similar aggregate morphologies and affected brain regions within each strain, resulting in nearly overlapping lesion profiles (spongiform change, gliosis, and prion deposition) and indicating no major differences in the histopathology due to the shortened HS chains (Fig. 1c).

To determine whether the biochemical properties of RML and ME7 were altered in mice expressing short HS chains, we compared the electrophoretic mobility, glycoform profile, and PrP[Sc] level in brain, and found no differences (Fig. 1d, e and Online Resource 2). Thus the shorter HS chains in *Ext1*^{+/−} mice do not markedly affect the brain lesions, PrP[Sc] levels, or the biochemical properties of subfibrillar prions, but led to a modest prolongation in survival following infection with one subfibrillar strain, ME7.

Short HS chains prolong survival and alter the distribution of fibrillar prions

Unlike RML and ME7, mCWD prions form long fibrils visible in situ by electron microscopy [95] and accumulate as extracellular plaques containing acidic polysaccharides (Online Resource 3). In contrast to the mice infected with subfibrillar prions, *tga*20[+/−]*Ext1*^{+/−} mice infected with mCWD prions showed a markedly prolonged survival that was more than 30 days longer as compared to the *tga*20[+/−]*Ext1*^{+/+} mice (275 ± 25 versus 237 ± 37 dpi, respectively) (Fig. 2a), yet with no differences in the total PrP[Sc] levels at terminal disease (Online Resource 2). However, the histopathology in the *tga*20[+/−]*Ext1*^{+/−} mice was profoundly

altered. Most of the *tga*20[+/−]*Ext1*^{+/+} mice infected with mCWD (80%, 8/10 mice) developed large (50–100 μm) extracellular plaques in the corpus callosum (Fig. 2b), as is typical for this prion strain (100% of mCWD-infected *tga*20 mice, $n = 11$) [12]. In contrast, only approximately 30% (5/16) of *tga*20[+/−]*Ext1*^{+/−} mice accumulated plaques in the corpus callosum (Fig. 2c), and the plaque number in each mouse was significantly reduced (Fig. 2d). Instead, mCWD prions accumulated within and around blood vessels (amyloid angiopathy) more broadly distributed throughout the brain, including unusual locations rarely observed in the *tga*20[+/−]*Ext1*^{+/+} mice or in any of the previous mCWD-infected *tga*20 mice examined, such as the thalamus and hypothalamus (Fig. 2c and Online Resource 4a).

Interestingly, *tga*20[+/−]*Ext1*^{+/−} brains showed significantly more perivascular plaques in the cerebellum and velum interpositum, a continuum of the subarachnoid space [30], as compared to *tga*20[+/−]*Ext1*^{+/+} brains (Fig. 2d). In the *tga*20[+/−]*Ext1*^{+/+} brains, the plaque numbers in the corpus callosum and velum interpositum were inversely correlated ($p < 0.01$, Spearman correlation test), suggesting that with PrP[Sc] forming fewer parenchymal plaques, deposition occurred instead perivascularly.

We next compared the plaque distribution with survival time in the *tga*20[+/−]*Ext1*^{+/−} mice and found that the mice that survived the longest (more than 260 dpi) had the most widely distributed plaques, with plaques in the hippocampus, cortex, thalamus, and basal ganglia (Online Resource 4b), despite having total plaque numbers that did not significantly vary from the mice with shorter survival times.

The spongiform change and gliosis in the infected *tga20*^{+/−}*Ext1*^{+/−} brains closely resembled the *tga20*^{+/−}*Ext1*^{+/+} brains (Fig. 2e). The plaques remained congophilic (Fig. 2f) and showed a similar plaque morphology as seen in the *tga20*^{+/−}*Ext1*^{+/+} brains (Fig. 2b). Dual immunostaining of brain sections for PrP and endothelial cells revealed that all the plaques in the corpus callosum were parenchymal, whereas nearly all plaques in non-callosal regions were perivascular (*tga20*^{+/−}*Ext1*^{+/+}: 88% and *tga20*^{+/−}*Ext1*^{+/−}: 94%) (Fig. 2g). Notably, some plaques were also observed subjacent to the ventricle (*tga20*^{+/−}*Ext1*^{+/+}: 8% and *tga20*^{+/−}*Ext1*^{+/−}: 3%).

To determine how the expanded mCWD plaque distribution impacted the neuroinflammatory response, we next measured the level of activated microglia in the brains of *tga*20[+/−]*Ext1*^{+/−} and *tga*20[+/−]*Ext1*^{+/+} mice by immunolabeling with ionizing calcium-binding adaptor molecule 1 (Iba1) antibody [47]. The level of microglial activation and clustering around mCWD plaques was indistinguishable between mice at terminal prion disease (Fig. 2h), suggesting there were no differences in the inflammatory response.

Short HS chains could alter the mCWD conformation, which may change the plaque distribution and prolong prion

disease. Thus, we next studied whether the mCWD fibril structure differed in tga20$^{+/-}$$Ext1$$^{+/-}$ mice using fluorescence lifetime imaging (FLIM) of h-FTAA stained deposits. h-FTAA is a heptameric oligothiophene that has previously been used to distinguish prion aggregates associated with distinct prion strains [2, 65]. FLIM experiments were conducted with excitation of h-FTAA at 490 nm and the acquired decay curves were fitted with a bi-exponential decay function and two components of the fit. We found that the intensity-weighted mean lifetime (ti) displayed by h-FTAA showed similar distributions for mCWD deposits in tga20$^{+/-}$$Ext1$$^{+/-}$ and tga20$^{+/-}$$Ext1$$^{+/+}$ mice (Fig. 3a). Thus, h-FTAA binds in an analogous fashion to these prion deposits, suggesting that mCWD is not structurally altered in the tga20$^{+/-}$$Ext1$$^{+/-}$ mice. Consistent with this result, the biochemical properties of the mCWD PrP^{Sc} in tga20$^{+/-}$$Ext1$$^{+/+}$ and tga20$^{+/-}$$Ext1$$^{+/-}$ mice, including electrophoretic mobility, glycoprofile, and aggregate stability in GdnHCl, were indistinguishable (Fig. 3b, c), indicating that the altered prion distribution observed with $Ext1$ haploinsufficiency was not due to a change in the prion conformation.

mCWD prions are GPI anchorless

PrP^C is tethered to the outer leaflet of the plasma membrane by a GPI anchor; however, 10–15% of PrP^C is constitutively cleaved between Gly228 and Arg229 (murine PrP) by the metalloproteinase ADAM10, releasing nearly full-length PrP^C from the cell surface [4, 106] (Fig. 4a). Prion aggregates reportedly contain the GPI anchor [103], yet conformers are diverse and it is possible that not all prions are GPI anchored. To determine whether mCWD prions harbor a GPI anchor, we employed a newly developed antibody that recognizes ADAM10-cleaved PrP (sPrP^{G228}) [63] and an antibody that recognizes cleaved and uncleaved (total) PrP (POM19) [85]. We found that abundant mCWD PrP^{Sc}, yet minimal RML and ME7 PrP^{Sc}, were recognized by the sPrP^{G228} antibody, indicating that PK-resistant mCWD is largely composed of ADAM10-cleaved PrP lacking the GPI anchor and three C-terminal amino acid residues (Fig. 4b). The level of ADAM10-cleaved PrP^{Sc} correlated with the plaque morphology among the three strains, as mCWD had the highest ratio of GPI-anchorless PrP^{Sc} to total PrP^{Sc} signal (630) and showed 100% extracellular plaques. In contrast, RML and ME7 had lower ratios (0.36 and 3.25, respectively) (Fig. 4c) and no or few extracellular plaques, respectively, consistent with being GPI anchored, as previously reported for PrP^{Sc} [67, 104]. To localize ADAM10-cleaved PrP^{Sc} in mCWD-infected brain, we performed immunostaining on brain sections using the sPrP^{G228} antibody. These experiments revealed strong immunolabeling of mCWD plaques, indicating GPI-anc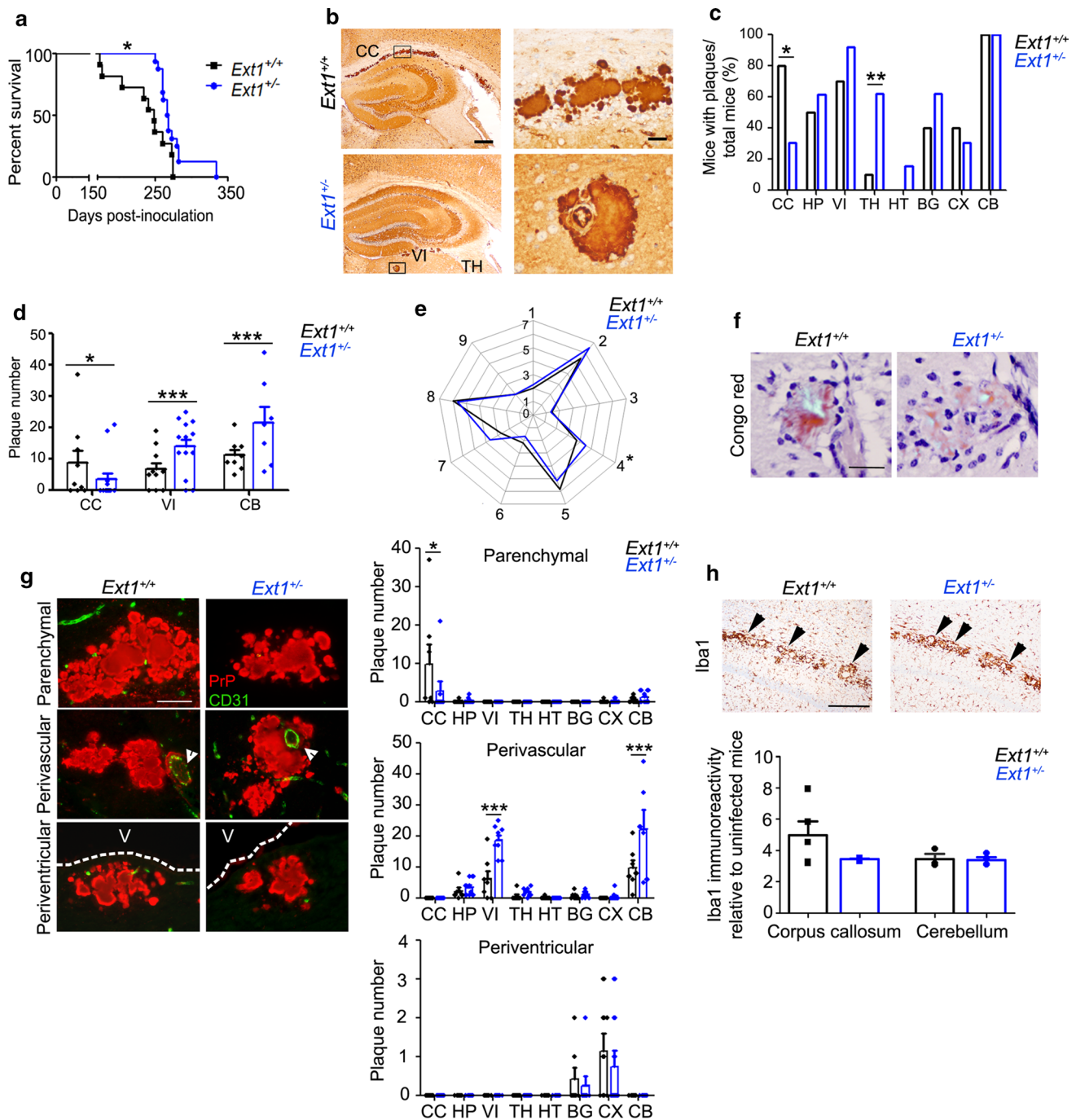horless PrP forms plaques of near full-length PrP. Conversely, there was very weak immunolabeling of RML aggregates (Fig. 4d).

GPI-anchorless prions contain high levels of HS

We reasoned that the GPI-anchorless state may enable shed PrP^C and small prion aggregates to better interact with HS in the extracellular matrix. In this case, GPI-anchorless prions would be predicted to have more HS bound. To compare the levels of HS bound to GPI-anchored and -anchorless prions, we utilized two GPI-anchored (sPrP^{G228} negative), subfibrillar prion strains, RML and (22L shown in Online Resource 3 and 5), that had been serially passaged in transgenic mice expressing GPI-anchorless PrP [tg(GPI-PrP)] [21], generating GPI-anchorless RML and 22L (GPI-anchorless RML shown in Online Resource 3). In the GPI-anchorless state, RML and 22L prions are fibrillar and vasculotropic [3], as originally reported by Chesebro and colleagues for RML [21]. We purified the GPI-anchored and anchorless prions from brain, assessed the PrP^{Sc} purity by silver stain (Online Resource 6), quantified the PrP^{Sc} levels, measured the HS levels by liquid chromatography–mass spectrometry (LC–MS), and calculated the HS: PrP^{Sc} ratio. Remarkably, we found approximately 20-fold more HS bound to GPI-anchorless RML and 22L prions as compared to GPI-anchored RML and 22L (Fig. 4e, f). Furthermore, upon testing mCWD prions we also found relatively high levels of HS bound (Fig. 4f), consistent with the other GPI-anchorless prions.

Although the LC–MS was performed on highly purified prion preparations, it is possible that HS associates with PrP during the isolation process. To further investigate the association of HS and PrP, we localized PrP and HS in brain sections from mCWD-infected tga20$^{+/-}$$Ext1$$^{+/+}$ and tga20$^{+/-}$$Ext1$$^{+/-}$ by dual immunolabeling in situ. Supporting the in vivo binding of PrP and HS, we found that HS intensely co-labeled with prion plaques, whereas the heparinase-digested (Fig. 4g) and isotype control samples were HS negative. HS and PrP single stains confirmed antibody specificity. We found that parenchymal plaques consistently showed strong HS immunolabeling (100%, $n = 35$ plaques).

The GPI-anchorless prions had exceedingly more HS bound than their GPI-anchored counterparts, suggesting that GPI-anchored prions rarely or only transiently interact with HS. We reasoned that adding a GPI anchor onto mCWD should create a prion that binds less HS. Therefore, we next tested a recently developed *GPI-anchored* mCWD strain obtained by passaging GPI-anchorless mCWD into tga20 mice (new mCWD) [3] (Fig. 5a–c). The new GPI-anchored mCWD formed diffuse deposits in the corpus callosum and hippocampus, and no longer formed plaques, bound Congo red, or stained strongly with Alcian blue (Fig. 5a). We performed LC–MS on prion-bound HS isolated from the

brains of mice infected with GPI-anchorless mCWD prions (tg(GPI-PrP) mice) and the new GPI-anchored mCWD prions (*tg*a20 mice) to compare with the original ADAM10-cleaved mCWD. Strikingly, while the ADAM10-cleaved mCWD and GPI-anchorless mCWD prions had abundant HS bound, the new GPI-anchored mCWD prions resembled other GPI-anchored prions in consistently showing very low levels of HS binding (Fig. 5d, e), supporting earlier findings that GPI-anchored prions harbor less HS. Interestingly, the HS disaccharide composition was somewhat similar between the ADAM10-cleaved mCWD and the new GPI-anchored mCWD, yet differed from the GPI-anchorless mCWD, as HS was generally more sulfated in the GPI-anchorless mCWD, specifically in the level of N-sulfated and 2-O-sulfated HS (Fig. 5f and Online resource Table 1). In HS-immunolabeled brain sections, we also found that HS co-localized with ADAM10-cleaved and GPI-anchorless mCWD, but not with the new GPI-anchored mCWD prions, indicating that HS mainly accumulates in extracellular, fibrillar prion plaques (Fig. 5g).

◀**Fig. 2** mCWD-infected *tg*a20$^{+/-}$*Ext1*$^{+/-}$ mice show prolonged survival times and altered plaque distribution. **a** mCWD-infected *tg*a20$^{+/-}$*Ext1*$^{+/-}$ ("*Ext1*$^{+/-}$") mice show a significant delay in survival time. **b** PrP immunolabeled brain sections show mCWD prion plaques in the corpus callosum (CC) of *Ext1*$^{+/+}$ mice, whereas in *Ext1*$^{+/-}$ mice, plaques are present in other brain regions including thalamus (TH), and velum interpositum (VI). The plaque morphology was unchanged. **c** The distribution of mCWD plaques varied between the *Ext1*$^{+/+}$ and *Ext1*$^{+/-}$ mice, as fewer *Ext1*$^{+/-}$ mice developed plaques in the corpus callosum (CC), whereas more *Ext1*$^{+/-}$ mice developed plaques in the basal ganglia (BG) and thalamus (TH) (*HP* hippocampus, *HT* hypothalamus, *CX* cerebral cortex, and *CB* cerebellum). **d** *Ext1*$^{+/-}$ mice show fewer plaques in the corpus callosum and more plaques in the velum interpositum and cerebellum. **e** Lesion profiles comparing spongiform degeneration, astrogliosis, and PrP^{Sc} deposition are similar in *Ext1*$^{+/+}$ and *Ext1*$^{+/-}$ mice (1—medulla, 2—cerebellum, 3—hypothalamus, 4—medial thalamus, 5—hippocampus, 6—septum, 7—cerebral cortex, 8—cerebral peduncle and 9—cerebellar peduncle). **f** mCWD plaques are congophilic in *Ext1*$^{+/+}$ and *Ext1*$^{+/-}$ mice [shown is cerebellum (*Ext1*$^{+/+}$) and hippocampus (*Ext1*$^{+/-}$)]. **g** Dual immunostaining for PrP and endothelial cells (CD31) shows typical non-vascular plaques in the corpus callosum (upper panels), perivascular plaques in the basal ganglia, and periventricular plaques adjacent to the lateral ventricle (middle panels, white arrowheads show blood vessel; bottom panels, *V* ventricle). *Ext1*$^{+/-}$ mice show fewer parenchymal plaques in the corpus callosum and more vascular plaques in the velum interpositum and cerebellum. **h** Iba1 immunolabeling of activated microglia shows similar clustering of activated microglia around mCWD plaques in *Ext1*$^{+/+}$ and *Ext1*$^{+/-}$ brain sections (arrowheads). Quantification of activated microglia shown in the upper panel (for corpus callosum *Ext1*$^{+/+}$: $n = 5$ and *Ext1*$^{+/-}$: $n = 3$; for cerebellum *Ext1*$^{+/+}$: $n = 3$ and *Ext1*$^{+/-}$: $n = 4$). Scale bars = 500 μm (left) and 50 μm (right) (**b**), 50 μm (**f**, **g**), and 1 mm (panel **h**). *$P < 0.05$, Log-rank (Mantel–Cox) test (**a**). *$P < 0.05$, **$P < 0.01$, ***$P < 0.001$, Fisher's exact test (**c**), two-way ANOVA with Bonferroni's post test (**d**, **g**), unpaired, 2-tailed Student's *t* test (**e**). *Ext1*$^{+/+}$: $n = 11$ mice; *Ext1*$^{+/-}$: $n = 16$ mice

Human plaque-forming prions also bind abundant HS

In certain human familial prion diseases, particularly Gerstmann–Sträussler–Scheinker (GSS) disease, N- and C-terminally cleaved, GPI-anchorless PrP forms parenchymal plaques or perivascular amyloid [35–38, 48, 83, 116], whereas in other diseases, near full-length to full-length PrP forms fine granular synaptic deposits and plaque-like deposits [50]. To test whether human plaque-forming prions also bind significantly more HS than the non-plaque-forming prions, we used LC–MS to measure the prion-bound HS levels and composition in post-mortem brain samples from 11 patients diagnosed with familial or sporadic prion disease. The familial cases consisted of five patients having missense mutations in codon 198 (F198S) or codon 102 (P102L) of *PRNP*. The six sCJD cases were previously characterized by subtype [74], and consisted of subtypes MM1 (2), MV1 (1), MV2K (2), and VV2 (1) (Table 1).

Four of the patients with familial prion disease (GSS-F198S) had developed compact multicentric PrP plaques involving deep layers of the neocortex and subpial regions

with no spongiform degeneration (Online resource 7). Confluent plaques were identified throughout the gray matter structures within the basal ganglia, thalamus, midbrain, cerebellar molecular layer, and dentate nuclei. PK-resistant PrP fragments of 8 kDa and higher were detected by immunoblotting (Fig. 6a), consistent with published reports [24, 25, 37, 38, 44, 84]. The fifth familial case (GSS-P102L) developed moderate, full-thickness spongiform degeneration with synaptic deposits (no plaques) throughout the cortex (Online resource 7) and scattered pleomorphic unicentric and multicentric plaques and plaque-like deposits in the cerebellum and thalamus. Immunoblotting of the frontal cortex revealed PK-resistant PrP composed of three bands from ~21 to 30 kDa, similar to PrP from the sCJD MM1 cases although with a different glycoform ratio (Fig. 6a), consistent with a previously published study [78].

The three sCJD cases with rapidly progressive disease (1.6–6 months) (MM1 and MV1) histologically showed a moderate degree of spongiform change with small, sometimes confluent vacuoles and fine granular and synaptic PrP aggregates (Online Resource 7). PK-resistant PrP having an electrophoretic mobility of 21 kDa (unglycosylated PrP) was detected by immunoblot (Fig. 6a). The MV2 and VV2 cases were characterized histologically by mild spongiform change (MV2) or focal areas of spongiform change (VV2) and diffuse aggregates as well as small plaques and plaque-like deposits (Online resource 7 and Table 1). Upon immunoblotting, PK-resistant bands were present at 19 and 20 kDa or 19 kDa, respectively (Fig. 6a).

Consistent with the results in mice, mass spectrometry of HS bound to PrP purified from the human brain samples revealed abundant HS bound to the plaque-forming prions (GSS-F198S cases), with approximately 10- to 100-fold higher HS levels than HS present in the seven cases having plaque-like deposits (sCJD) or diffuse deposits [sCJD and GSS-P102L (cortex)], respectively (Fig. 6b, c). Remarkably, the lowest HS levels (~100-fold lower than GSS-F198S) were associated with the diffuse prion aggregates [sCJD MM1 and MV1 and GSS-P102L (cortex)], while intermediate levels of HS (~10-fold lower) were associated with the sCJD cases having small plaques and plaque-like deposits (VV2 and MV2 K) (Fig. 6c). Thus, the HS levels once again correlated with the presence of plaques or plaque-like deposits in the brain.

Notably, the composition of the HS bound to the multicentric plaque-forming GSS prions was similar to that bound to sCJD prions. The primary differences were in the relatively low levels of unsulfated N-acetylated HS and 6-O-sulfated HS in the GSS-F198S cases as compared to the sCJD cases (Fig. 6d).

To localize the abundant HS detected in the GSS-F198S cases in tissue sections, we next immunolabeled brain
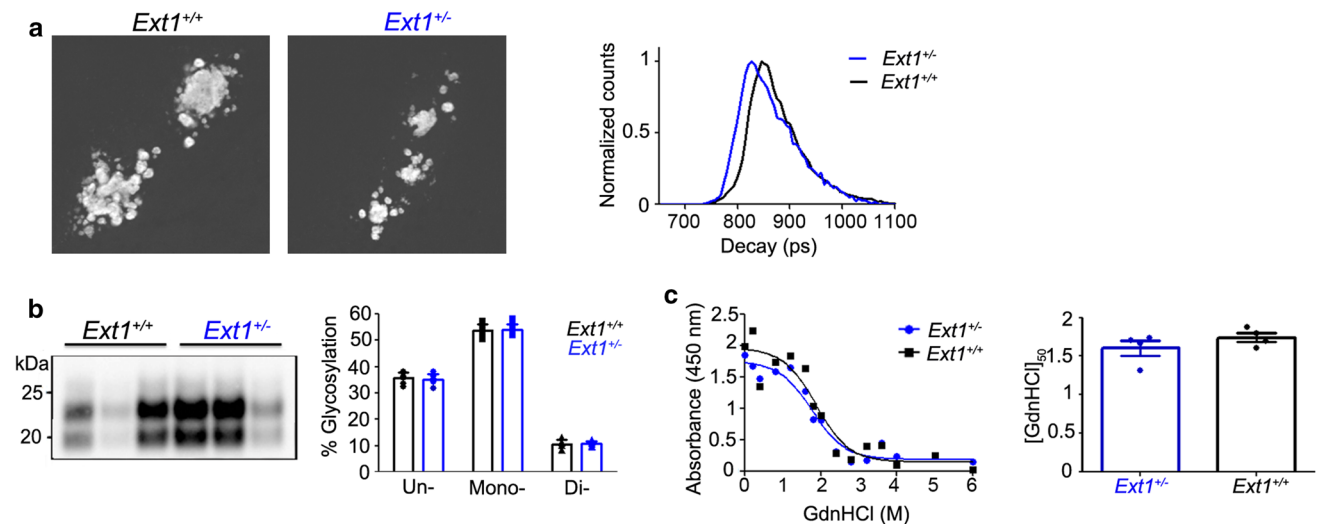
Fig. 3 mCWD prion conformation is similar in mice having long or short HS chains. **a** h-FTAA fluorescence lifetime decay of mCWD prion plaques in *tg*a20$^{+/−}$*Ext1$^{+/+}$* ("*Ext1$^{+/+}$*") and *tg*a20$^{+/−}$*Ext1$^{+/−}$* ("*Ext1$^{+/−}$*") brain sections are similar. **b** mCWD electrophoretic mobility and glycoprofile are also similar in *Ext1$^{+/+}$* and *Ext1$^{+/−}$* mice. **c** Representative example of PrP^{Sc} aggregate stability as measured by GdnHCl denaturation in mCWD-infected *Ext1$^{+/+}$* and *Ext1$^{+/−}$* mice. [GdnHCl]$_{1/2}$ values are shown for mCWD in *Ext1$^{+/+}$* and *Ext1$^{+/−}$* brain ($n = 4$ mice/genotype; each run in triplicate)

sections and found intense HS labeling concentrated in the plaque core (Fig. 6e), reported to contain the N- and C-terminally truncated PrP [38]. This finding indicates that HS binds prion plaques in human brain parenchyma, and less so to synaptic and punctate deposits, consistent with the findings in the prion-infected mice.

Discussion

Here, we provide the first in vivo evidence of an endogenous cofactor, HS, that facilitates parenchymal prion plaque deposition in the brain. mCWD prion-infected mice with *Ext1* haploinsufficiency showed fewer parenchymal plaques, an increase in vascular plaques, and a prolonged survival time. In addition, we found that the mCWD plaques were composed of ADAM10-cleaved, extracellular PrP and exceedingly high levels of HS, particularly 6-O-sulfated HS. Although mCWD-infected *tg*a20$^{+/−}$; *Ext1$^{+/−}$* mice showed a longer survival time, there was no evidence for a change in the prion conformation, biochemically or histologically, implicating a role for HS in the altered plaque distribution. Taken together, our studies suggest that HS binds shed, GPI-anchorless prion protein and enhances parenchymal plaque formation, potentially through scaffolding prion fibril assembly and reducing prion transit toward blood vessels.

Surprisingly, *Ext1* haploinsufficiency led to either a modest or no change in the disease progression induced by GPI-anchored, subfibrillar prions. This remarkable and unexpected finding suggests little or only a transient interaction

between HS and GPI-anchored prions, in stark contrast to the GPI-anchorless prions in mice and humans typically form parenchymal plaques or cerebral amyloid angiopathy (CAA), depending on the specific mutation [21364890]. We found a strong correlation between the GPI-anchored state of the prion and the level of HS bound. Two GPI-anchorless prion strains (GPI$^−$ RML and GPI$^−$ 22L) bound abundant HS, whereas their GPI-anchored counterparts bound very little, suggesting that GPI-anchorless prions interact more frequently with HS or with higher affinity. Additionally, while ADAM10-cleaved, (GPI-anchorless) mCWD prions bound abundant HS, the new GPI-anchored mCWD prions bound very little HS. Collectively, our results suggest that GPI anchoring of prions to the cell membrane topologically constrains prions, limiting access and binding to extracellular HS. Thus, we propose two key factors tightly linked to the formation of parenchymal plaques in prion disease that ultimately impact the disease phenotype: (1) mobile, GPI-anchorless PrP and (2) extracellular HS.

The modest effect of *Ext1* haploinsufficiency on GPI-anchored prion disease seemingly contradicts previous studies that HS or heparin binds and promotes PrP internalization and conversion in vitro [7, 42, 46, 77, 94]. In vitro, exogenously applied sulfated polyanions were shown to reduce prion replication in cells, potentially by inhibiting prion binding to endogenous HS [19]. Additionally, heparanase overexpression in RML-infected mice infected led to a prolonged survival time [54]; however, the impact on prion aggregate conformation and brain distribution was not described. Here, HS chains were shortened but
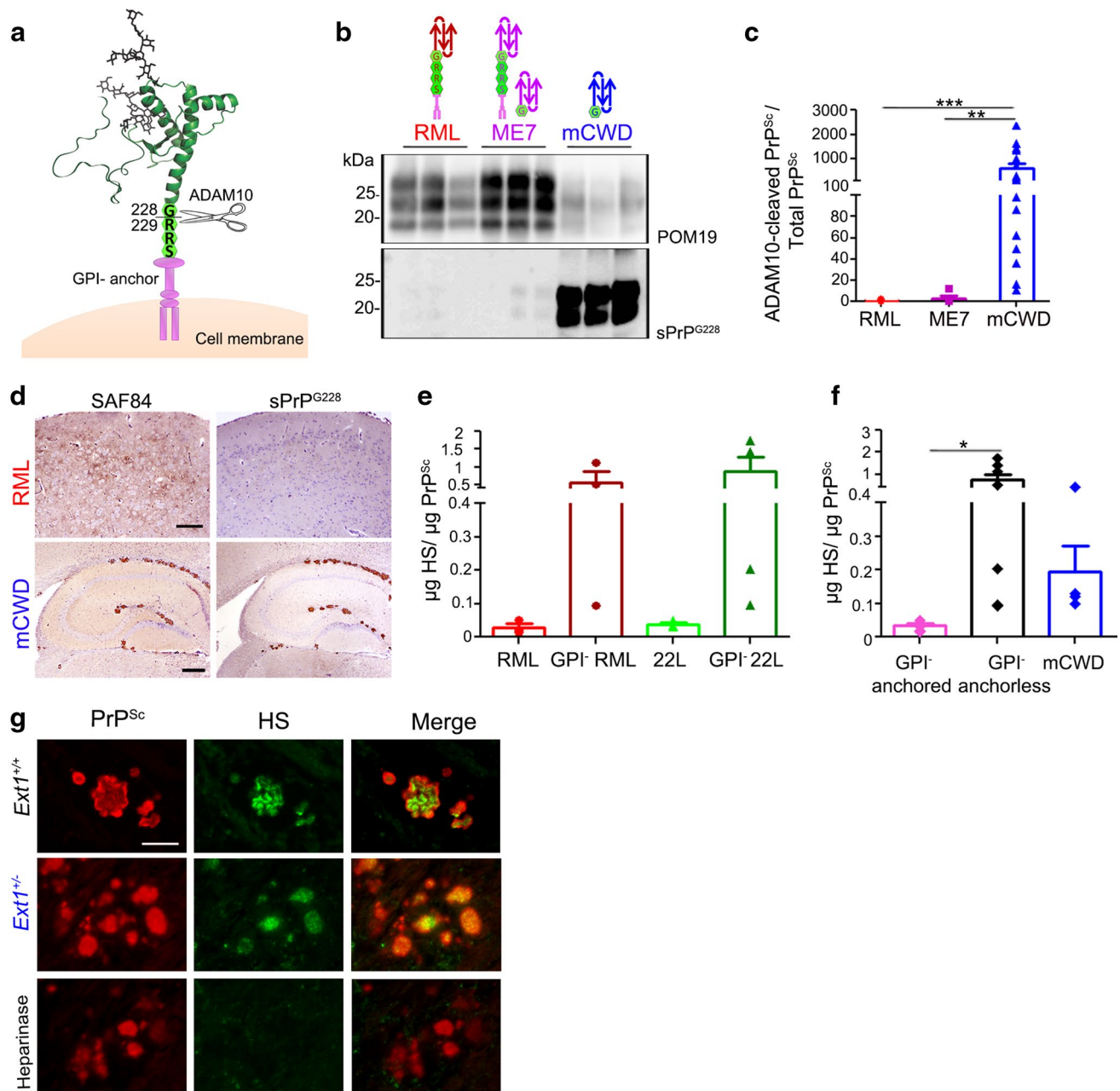
Fig. 4 ADAM10-cleaved and full-length GPI-anchorless prions bind HS. **a** Schematic representation of ADAM10 cleavage at mouse PrP residue 228 shows the release of shed PrP lacking the GPI anchor and three C-terminal amino acid residues (RRS). **b** Immunoblots of brain homogenate from prion-infected $Ext1^{+/+}$ mice using POM19 antibody (PrP) and sPrP^{G228} antibody (ADAM10-cleaved PrP). **c** Ratios of ADAM10-cleaved PrP^{Sc} relative to total PrP^{Sc} reveal significantly higher levels of ADAM10-cleaved PrP in mCWD as compared to the RML and ME7 strains. **d** Brain immunolabeled for PrP with SAF84 (amino acids 163–169 of mouse PrP) and sPrP^{G228} antibodies reveals all mCWD plaques, but few diffuse RML aggregates, are labeled by sPrP^{G228}. **e** Quantification of HS bound to GPI-anchored and –anchorless RML and 22L prions by LC–MS, and **f** a grouped comparison of HS bound to GPI-anchored prions (RML and 22L) versus GPI-anchorless prions (GPI$^-$ RML, GPI$^-$ 22L) and ADAM10-cleaved mCWD shows that full-length, GPI-anchorless prions and ADAM10-cleaved prions (mCWD) bind more HS than their GPI-anchored counterparts (for RML and 22L). **g** Dual immunostaining of mCWD-infected brain sections for PrP and HS shows parenchymal prion plaques in the corpus callosum label strongly for HS. Pre-treating brain sections with heparinases abolished HS labeling of plaques. Scale bars = 200 μm and 500 μm for upper and lower panel (**d**) and 25 μm (**g**). *$P < 0.05$, **$P < 0.01$ and ***$P < 0.001$, Wilcoxon rank sum test (**c**) and one-way ANOVA with Tukey's post test (**f**)
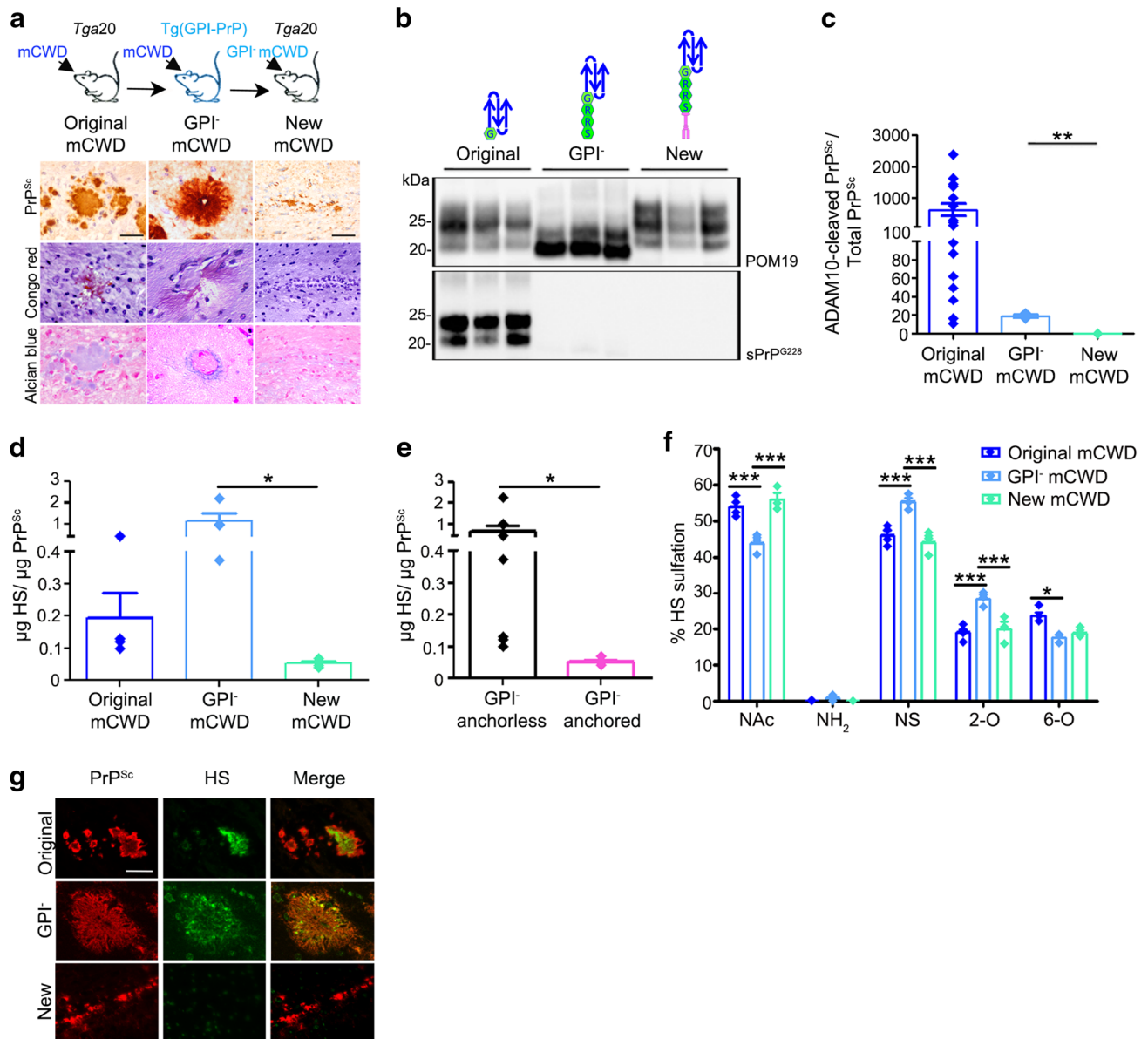
Fig. 5 GPI-anchored mCWD prions do not form plaques and bind low levels of HS. **a** Schematic representation of mCWD inoculated into *tg*a20 and GPI-anchorless PrP^C expressing mice [tg(GPI-PrP)] with corresponding prion plaque morphology, Congo red, and Alcian blue staining of brain sections (corpus callosum). Note that the new mCWD prions do not bind Congo red or Alcian blue. **b** Western blots of PK-treated PrP labeled with anti-PrP POM19 (total PrP) or sPrP^{G228} (ADAM10-cleaved PrP) antibodies reveal ADAM10-cleaved PrP in the original mCWD, but not the GPI-anchorless mCWD or the new mCWD–infected brain. **c** Ratios of ADAM10-cleaved: total PrP^{Sc} are higher in the original mCWD than in the GPI-anchorless mCWD or in the new mCWD-infected brains (original mCWD results are also shown in Fig. 4c). **d** The new mCWD binds less HS than the original mCWD or GPI-anchorless mCWD (original

mCWD results are also shown in Fig. 4f). **e** A grouped comparison shows HS bound to GPI-anchorless prions (mCWD and GPI⁻ mCWD, black bar) versus the new GPI-anchored mCWD (pink bar). **f** The HS bound to GPI-anchorless mCWD is more sulfated than the HS associated with ADAM10-cleaved mCWD and the new GPI-anchored mCWD, as it contains less unsulfated N-acetylated (NAc) HS and higher levels of N-sulfated (NS) and 2-O-sulfated (2-O) HS. **g** HS does not co-localize to the new mCWD aggregates (bottom panel). Scale bars = 100 μm for original and GPI⁻ mCWD and 200 μm for new mCWD (**a**) and 25 μm (**g**). *$P < 0.05$, **$P < 0.01$ and ***$P < 0.001$, Wilcoxon rank sum test (**c**, **e**), one-way ANOVA with Tukey's post test (**d**), and two-way ANOVA with Bonferroni's post-test (**f**)

not eliminated, therefore we cannot exclude a role for HS chains in advancing GPI-anchored prion disease. Further in vivo work using additional enzyme mutants involved in

Table 1 Human cases used for heparan sulfate analysis

Code	Age of onset	Disease duration (months)	Gender	*PRNP* genotype (mutation-129 codon)	PrP^Sc subtype	Symptoms at disease onset	Family history of prion disease
1	49	120	M	F198S-129MV	GSS, 8 kDa	Cerebellar	Sibling and niece
2	51	72	M	F198S-129VV	GSS, 8 kDa	Cerebellar	Sibling and daughter
3	60	89	M	F198S-129MV	GSS, 8 kDa	Not reported	Parent
4	67	63	M	F198S-129MV	GSS, 8 kDa	Not reported	Not reported
5	22	23	M	P102L-129MV	fCJD, Type 1	Seizures	Cousin
6	60	20	M	129MV	sCJD, Type intermediate + 2	Visuospatial	Not reported
7	55	6	F	129VV	sCJD, Type 2	Apraxia	Not reported
8	55	24	F	129MV	sCJD, Type intermediate + 2	Behavior	Not reported
9	69	6	F	129MV	sCJD, Type 1	Behavior/Memory	Not reported
10	60	1.5	M	129MM	sCJD, Type 1	Cognitive/Visual	Not reported
11	57	4	F	129MM	sCJD, Type 1	Motor	Not reported

HS synthesis would be required to better understand how cell surface HS impacts GPI-anchored prion disease.

The mCWD plaque redistribution to perivascular sites and the delay in disease observed in the *Ext1*^+/− mice are reminiscent of findings from an Alzheimer's disease mouse model (APP/PS1) deficient in neuronal HS chains, which showed a marked reduction in amyloid-β parenchymal plaques and an increased amyloid angiopathy compared to age-matched APP/PS1 mice [64]. Interestingly, clearance of the amyloid-β monomer was increased in mice lacking neuronal HS [64], suggesting that HS may trap amyloid-β in the parenchyma and prevent clearance by bulk fluid flow. In human brain, HS proteoglycans co-localize with amyloid-β plaques [97, 108], similar to what we observed in the familial prion disease cases with plaques (F198S). Thus, HS may be a fundamental component in parenchymal fibril assembly common to a subset of prion diseases and Alzheimer's disease. Future mass spectrometry studies may inform on the HS composition in the amyloid-β plaques, enabling further comparison with the prion-bound HS.

The mass spectrometry studies reported here are the first to quantify the levels and composition of HS tightly bound to any amyloidogenic protein in the brain. The findings show striking differences among the prion strains. For example, in comparing the GPI-anchored and -anchorless mCWD conformers, we noted significantly higher levels of N-acetylated and 6-O-sulfated HS obtained from the anchorless prions. It has been previously shown that HS sulfation plays a role in the internalization and assembly of prions, tau, amyloid-β, and α-synuclein [45, 46, 88, 105]. Additionally, sulfated GAGs stimulate prion conversion in vitro [46]. Our detection of major variation in the prion-associated HS among strains suggests either difference in the affinity of a prion fold for a particular HS structure or in the HS molecules

synthesized near the prion deposition site. These findings underscore the need to define the HS structures in different brain regions as well as the molecular determinants for HS binding to protein aggregates.

Collectively, these studies suggest that HS binds shed, GPI-anchorless prions in the parenchyma, the question remained, does HS scaffold prions, accelerating fibril assembly and plaque formation? HS chains are abundant, long flexible polymers that are highly anionic and typically act as a scaffold, with electrostatic interactions being major contributors to protein binding [15, 73, 93]. HS chains normally tether and locally concentrate proteins [27, 114], foster protein homo-oligomerization [81, 115], and stabilize extracellular proteins [92], all of which would promote prion assembly. In gelsolin fibril assembly, HS is not required for the nucleation phase, but instead promotes the extension phase, possibly by binding and scaffolding oligomers for conversion into fibrils [99]. In prion assembly, Ma and colleagues found that cofactors were required for recombinant prion protein to form a fibrillar conformation that was both infectious and pathogenic [111]. We suggest that prions co-opt HS as an endogenous cofactor post-nucleation, which facilitates fibril assembly, as here ADAM10-cleaved and GPI-anchorless prions consistently bound high levels of HS and formed fibrils. Nevertheless, we cannot exclude the possibility that HS passively binds prions post-plaque assembly, although this would not explain the plaque redistribution in the context of the shortened HS chains.

The present findings may help explain the extraordinarily rapid progression of most prion diseases, as the majority are subfibrillar. Diffuse, subfibrillar prion deposits (noncongophilic) correlate with rapidly progressive disease, whereas plaque-forming, fibrillar prions (congophilic) tend to correlate with longer, more slowly
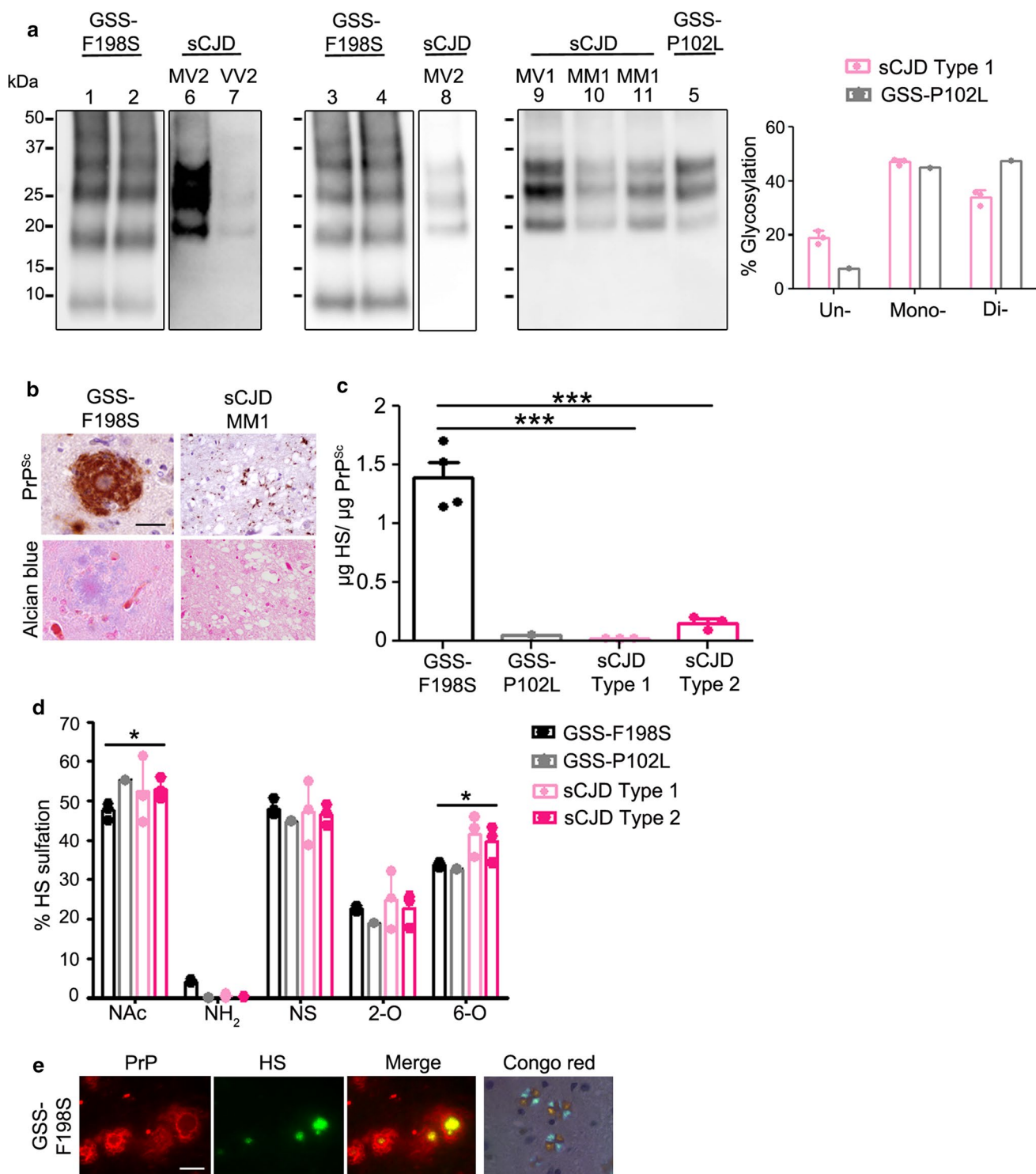
Fig. 6 Abundant HS is associated with prion plaques in human brain. **a** Immunoblots of PK-digested PrP purified from GSS and sCJD brain samples show differences in electrophoretic mobility and gly-coprofile. Note that PrP from the GSS-P102L brain shows a PK core size of 21 kD and a different glycoprofile than PrP from the sCJD MM1 and MV1 brain samples. **b** Fibrillar prion plaques from a GSS F198S patient and an sCJD patient immunolabeled for PrP. Plaques bind Alcian blue. **c** Mass spectrometry of HS from purified prion preparations derived from the GSS and sCJD patient brain samples reveal significantly higher HS levels associated with the GSS-F198S purified prions. **d** The HS bound to GSS-F198S prions shows lower levels of N-acetylated (NAc) and 6-O (6-O) sulfated disaccharides than the HS bound to sCJD. **e** Cerebellar GSS-F198S plaques show intense HS immunolabeling, primarily in the plaque core. Plaque cores in GSS-F198S affected brains are Congo red positive (shown is a representative example). Frontal cortex (GSS-F198S) and thalamus (sCJD MM1) are shown in (**b**). Scale bars = 50 μm (sCJD) and 100 μm GSS-F198S (**b**), 25 μm (dual IF, **e**) and 250 μm (Congo red, **e**). ***$P < 0.001$, One-way ANOVA with Tukey's post test (**c**). *$P < 0.05$, Two-way ANOVA with Bonferroni's post test (**d**)

progressive disease, with some exceptions. Patients with sporadic CJD (subtype MM1 or MV1) or familial CJD (fCJD), from missense mutations in codon 200 (E200K) [101] or 208 (R208H) [91] in *PRNP*, typically develop subfibrillar aggregates (synaptic, granular, and plaque-like deposits) and show a short survival time of less than 1 year [32, 51], whereas patients with PrP cerebral amyloidosis, for example, from missense or nonsense mutations in codon 102 (P102L) [43, 55, 112], 145 (Y145X), 226 (Y226X), or 227 (Q227X) in *PRNP* [36, 48], develop parenchymal or vascular amyloid and show a longer survival time, with a mean of 5 years [34, 51]. How HS impacts the kinetics of fibril assembly and ultimately survival time would be worth further study. The rapidly progressive disease in sCJD or fCJD could be explained, in part, by the limited access of GPI-anchored prions to extracellular HS scaffolds, reducing fibril assembly and resulting in primarily subfibrillar aggregates, although whether these aggregates are GPI anchored in human prion disease is not clear. GPI-anchorless PrP fragments have been detected in the brain of sCJD patients [70]. Our findings also have direct implications for therapy, as interfering with HS binding or modifying endogenous HS structure would be predicted to have the most significant impact on the progression of fibrillar, plaque-forming prion diseases, including GSS. As most systemic amyloids form extracellularly and bind HS, including islet amyloid polypeptide in type 2 diabetes [75], these results also suggest that interfering with HS binding to amyloidogenic proteins may be considered more broadly as a therapeutic strategy for amyloidoses.

Compliance with ethical standards

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