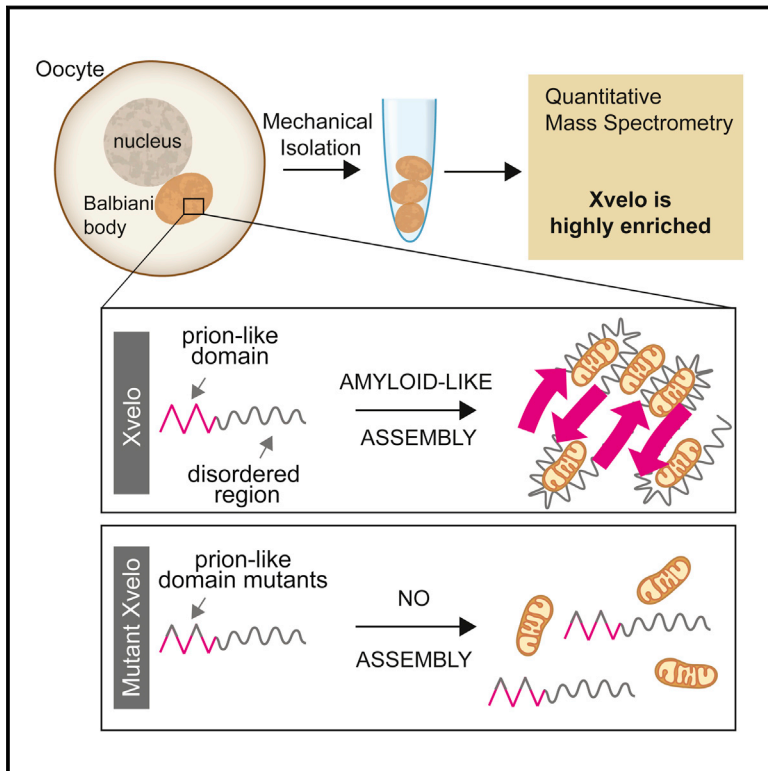


Amyloid-like Self-Assembly of a Cellular Compartment

Graphical Abstract



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In Brief

Amyloid-like self-assembly of a specific protein drives formation of a cellular compartment in oocytes.

Highlights

- The organelle content of the Balbiani body is held together by an Xvelo matrix
- Xvelo forms amyloid-like networks in vitro, which can recruit RNA and mitochondria
- Prion-like domain of Xvelo dictates specificity in amyloid assembly
- Amyloid-like polymerization is conserved among vertebrate Balbiani body organizers



Amyloid-like Self-Assembly of a Cellular Compartment

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SUMMARY

Most vertebrate oocytes contain a Balbiani body, a large, non-membrane-bound compartment packed with RNA, mitochondria, and other organelles. Little is known about this compartment, though it specifies germline identity in many non-mammalian vertebrates. We show Xvelo, a disordered protein with an N-terminal prion-like domain, is an abundant constituent of *Xenopus* Balbiani bodies. Disruption of the prion-like domain of Xvelo, or substitution with a prion-like domain from an unrelated protein, interferes with its incorporation into Balbiani bodies *in vivo*. Recombinant Xvelo forms amyloid-like networks *in vitro*. Amyloid-like assemblies of Xvelo recruit both RNA and mitochondria in binding assays. We propose that *Xenopus* Balbiani bodies form by amyloid-like assembly of Xvelo, accompanied by co-recruitment of mitochondria and RNA. Prion-like domains are found in germ plasm organizing proteins in other species, suggesting that Balbiani body formation by amyloid-like assembly could be a conserved mechanism that helps oocytes function as long-lived germ cells.

INTRODUCTION

Female germ cells, oocytes, are highly specialized cells. In many species, oocytes are long-lived and lie dormant for months or years before they are activated prior to fertilization (Li and Albertini, 2013). They ensure the continuity of the species by providing the female genome and mitochondria, along with most of the housekeeping machinery and nutrients the early embryo will need after fertilization. Oocytes have a unique subcellular organization, with a large nucleus, called the germinal vesicle, and a large cytoplasm. In many species, the cytoplasm of the early oocytes contains a highly specialized compartment called the Balbiani body. It is non-membrane bound and densely packed with mitochondria, RNA, ER, and Golgi. The Balbiani body assembles early in oocyte formation (Lei and Spradling, 2016), disappears in late-stage oocytes in mammals (Pepling et al., 2007), and forms dispersed isles at the vegetal pole in late-stage oo-

cytes of *Xenopus* and zebrafish (Bontems et al., 2009; Kloc et al., 2004).

How, or why, the Balbiani body forms is largely mysterious (Gupta et al., 2010; Heim et al., 2014; Kloc and Etkin, 1995; Kloc et al., 1993; Marlow and Mullins, 2008). In some species, including frogs and fish, but not newts and mammals, one function of the Balbiani body is to serve as an embryonic determinant that specifies germline identity by forming germ plasm (Lesch and Page, 2012; Richardson and Lehmann, 2010). Germ plasm, a special part of oocyte cytoplasm, protects specific maternal RNAs from degradation and is believed to host healthy mitochondria during development to pass on to future generations (Houston and King, 2000; Kogo et al., 2011). Germline specification is the only known function of the Balbiani body, but it is notable that oocytes from species that use inductive processes to specify their germline, including mammals, still contain a Balbiani body, whose ultrastructure is similar to those of organisms that use germ plasm (Hertig, 1968). We do not know what functions the Balbiani body serves in all vertebrates. We speculate their most conserved function is to protect the quality of mitochondria and other organelles during long periods of oocyte dormancy, which can extend for decades in humans.

An elegant maternal effect screen in zebrafish identified only one gene, *bucky ball*, required for Balbiani body formation (Dosch et al., 2004). In *buc* mutants, the Balbiani body did not form, and oocyte polarity was disrupted. Overexpression of Bucky ball protein stimulated numerous Balbiani bodies and ectopic germ cell formation (Bontems et al., 2009; Heim et al., 2014; Marlow and Mullins, 2008). Bucky ball is proposed to be a structural organizer of zebrafish Balbiani bodies, but the biochemical basis of this proposed function has been unclear.

Xenopus laevis provides complementary advantages for analysis of Balbiani bodies because oocytes are abundant and accessible, easy to manipulate due to their large size, and amenable to live imaging. *Xenopus* oocytes are classified in six different stages according to Dumont, 1972. Stage I oocytes contain a clearly visible Balbiani body (Figure 1A). The Balbiani body gradually disappears during oocyte development to give rise to dispersed germ plasm islands at the vegetal pole of the stage VI oocytes during oocyte maturation (Kloc et al., 2004).

Here, we report that the Balbiani body forms via amyloid-like self-assembly of Xvelo, the *Xenopus* homolog of Bucky ball. Amyloids and amyloid-like proteins have largely been studied in the context of neurodegenerative diseases, such as Alzheimer's

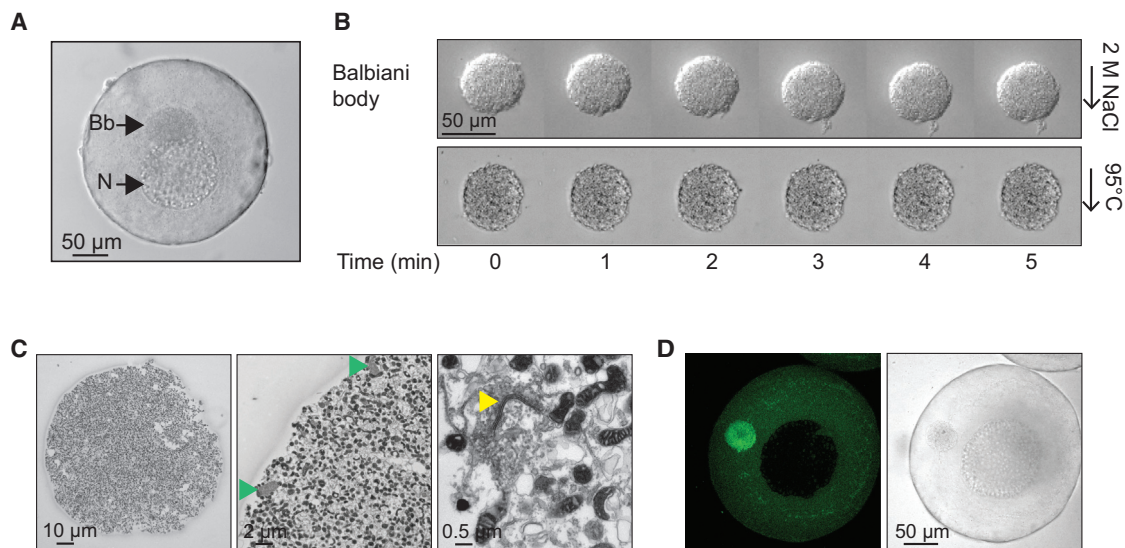


Figure 1. A Balbiani Body Is a Non-Membrane-Bound Compartment Packed with Membranous Organelles

(A) Phase contrast image of a stage I *Xenopus laevis* oocyte. Bb, Balbiani body; N, nucleus or germinal vesicle.

(B) Balbiani body immobilized in perfusion chambers. 2 M NaCl (first panel) or 95°C 50 mM HEPES, 100 mM KCl (pH 7.6) buffer (second panel) was perfused into the chambers.

(C) Thin-section electron microscope (EM) images of isolated Balbiani bodies from stage I *Xenopus* oocytes. Mitochondria (dark spots), RNP particles (green arrow head), and Golgi stacks (yellow arrow head) are clearly visible.

(D) Stage I oocytes were incubated in 10 μM Thioflavin T in 1× MMR for 10 min and washed twice with 1× MMR.

See also [Figure S1](#), [Table S1](#), and [Movie S1](#).

disease, amyotrophic lateral sclerosis, or prion diseases (Haass and Selkoe, 2007; Koo et al., 1999; Polyimenidou and Cleveland, 2011). Several examples of amyloid-like aggregation mechanisms with normal biological functions have recently emerged (Berchowitz et al., 2015; Fowler et al., 2006; Hou et al., 2011; Li et al., 2012). Both pathological and physiological amyloid-forming proteins contain low-complexity regions, which are intrinsically disordered but can undergo conformational conversions into an amyloid-like fibrillar state (Kato et al., 2012). These regions are also found in yeast prion proteins, a class of infectious proteins that generate heritable phenotypic diversity in clonal population of yeast cells (Alberti et al., 2009; Halfmann et al., 2012; Wickner et al., 2013). Protein regions with sequence compositions similar to yeast prions are called prion-like domains (PLDs) (Alberti et al., 2009; Si et al., 2003).

RESULTS

A Balbiani Body Has a Rigid Structure that Stains with Thioflavin T

We began our studies on Balbiani bodies by manually dissecting them from *Xenopus laevis* stage I oocytes, which are transparent and 50–300 μm in diameter (Figure 1A). Stage I oocytes were placed in a glass-bottom dish together with a cytoskeleton stabilizing buffer and mechanically disrupted using tweezers, leaving the Balbiani body intact. The Balbiani body preserved its circular shape and did not disintegrate under the considerable shear forces applied during isolation (Movie S1). Balbiani bodies also retained their structure under harsh conditions, including

high salt (2 M NaCl) (Figure 1B, first panel) and high temperature (up to 95°C) (Figure 1B, second panel). Confirming previous work on intact oocytes, thin-section electron microscopy of the isolated Balbiani bodies revealed densely packed mitochondria, ER, and Golgi stacks as well as compact fibrillar elements that others have shown to be made of ribonucleoprotein (RNP) (Figure 1C) (al-Mukhtar and Webb, 1971; Balinsky and Devis, 1963; Billett and Adam, 1976).

To probe the molecular properties of the Balbiani body, we introduced a number of small molecules into the oocytes. Thioflavin T (ThT), a dye that stains the β sheet-rich structures of amyloid (Alberti et al., 2010; Nilsson, 2004), strongly accumulated in the Balbiani body (Figure 1D), suggesting it is rich in β sheet structures, which is a hallmark of amyloids.

Xvelo Is Highly Concentrated in Balbiani Bodies

To analyze the composition of Balbiani bodies, we used quantitative mass spectrometry (McAlister et al., 2014; Wühr et al., 2014). This revealed that the most enriched Balbiani body proteins that are not part of organelles are Xvelo (commonly known as Xvelo) and fetal hemoglobin subunits, Hba1 and Hbg1 (Figures S1A–S1C). We focused on Xvelo because it is homologous to zebrafish Bucky ball, which plays a crucial role in Balbiani body organization (Bontems et al., 2009). Computational analysis of Xvelo sequence predicted an intrinsically disordered protein with no known domains, apart from a C-terminal positively charged region that could bind RNA.

Quantitative western blotting using a peptide antibody against the C terminus of Xvelo (Figure S2A) provided an estimate of

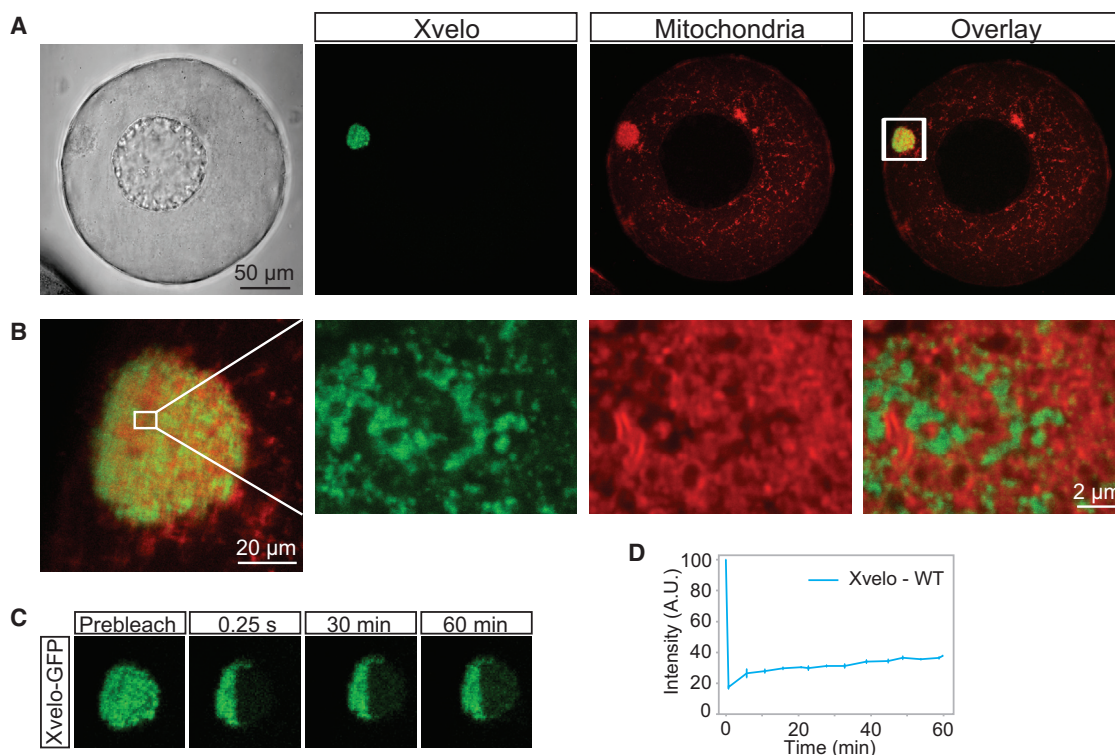


Figure 2. Xvelo Forms a Stable Matrix

(A) mRNA encoding for Xvelo-GFP was microinjected into stage I oocytes. MitoTracker Deep Red was used to label mitochondria. Oocytes were imaged live with a laser scanning confocal microscope with a 40 \times water-immersion objective.

(B) Magnification of the Balbiani body in (A).

(C) Internal rearrangement of fluorescent Xvelo-GFP particles after half bleach over time.

(D) The fluorescent recovery of the half-bleached Xvelo-GFP in the Balbiani body in (C) and two other biological replicates is shown by quantification of fluorescence in bleached region over time. Fluorescent intensity changes in the bleached region per pixel over time were plotted after it was normalized for photobleaching by using an unbleached neighboring area and background subtraction.

See also [Figure S2](#).

Xvelo concentration in oocytes of $\sim 8 \mu\text{M}$ ([Figures S2B](#) and [S2C](#)). If all the Xvelo was concentrated in the Balbiani body, and assuming an average late stage I oocyte diameter is 250 μm and Balbiani body diameter is 60 μm , this corresponds to $\sim 560 \mu\text{M}$ in the Balbiani body. Compared with the published concentrations of proteins in a frog egg, where the most concentrated proteins were alpha-1-antitrypsin (Serpina1) and actin (Actg1) at 17.6 and 14.3 μM , respectively ([Wühr et al., 2014](#)), one can appreciate the exceptionally high concentration of Xvelo in Balbiani bodies. This high local concentration of Xvelo, together with its homology to the zebrafish protein Bucky ball, makes it a likely candidate for a structural organizer of the Balbiani body.

Xvelo Forms a Stable Matrix in Balbiani Bodies

To investigate Xvelo function in *Xenopus*, we began by injecting mRNAs encoding for Xvelo tagged with GFP (Xvelo-GFP) into the oocyte cytoplasm. Xvelo-GFP mainly localized to the Balbiani body with little soluble protein present in the cytoplasm ([Figure 2A](#)). More detailed analysis at higher magnification showed that Xvelo-GFP filled the gaps between the mitochondria in the Balbiani body ([Figure 2B](#)). These results suggest that Xvelo

may form a matrix in which mitochondria and other organelles are embedded.

To test whether Xvelo was part of a stable matrix, we used FRAP (fluorescence recovery after photobleaching) to probe its dynamics. We again injected mRNA encoding for Xvelo-GFP, and after overnight incubation, we imaged the oocytes. After photobleaching Xvelo-GFP, maximum recovery of $\sim 20\%$ was seen after 1 hr ([Figures 2C](#) and [2D](#)). We conclude that Xvelo matrix is highly stable, with slow turnover, consistent with a role as a structural matrix.

Xvelo's Association with the Balbiani Body Requires Its Prion-like Domain

We next investigated how the Xvelo matrix is formed and held together. Despite the lack of any conventional domains, Xvelo has a PLD at its N terminus, which is detected by several prion detection algorithms ([Figure 3A](#); [Figure S3A](#)) ([Alberti et al., 2009](#); [Lancaster et al., 2014](#); [Toombs et al., 2012](#)). It also has a lysine/ arginine rich region at its C terminus that might act as an RNA binding domain ([Figure 3A](#)). To assess the behavior of these different parts of Xvelo in vivo, we performed a structure-function analysis by dividing the protein into four fragments

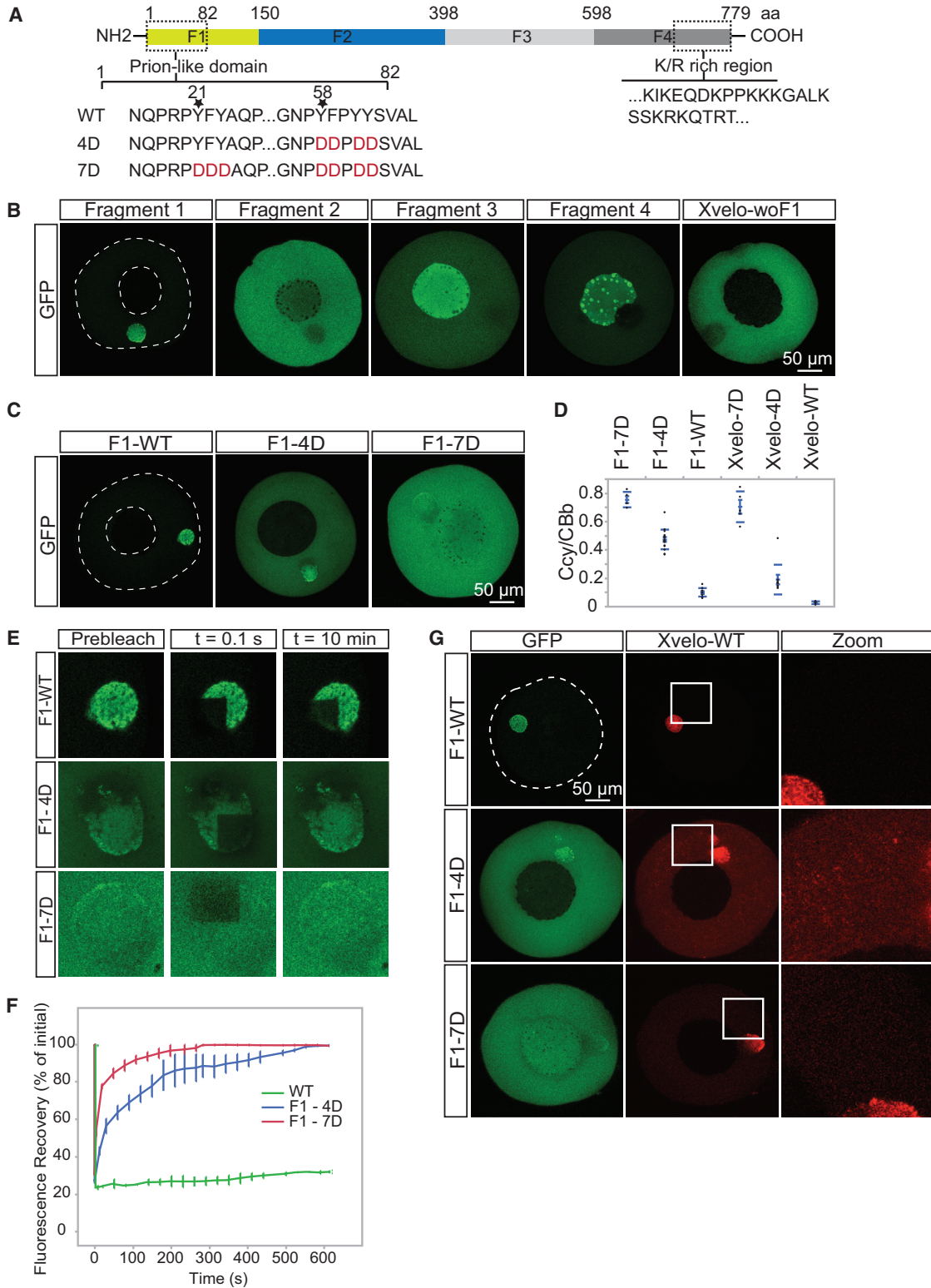


Figure 3. Xvelo Self-Assembly Is Dependent on Its Prion-like Domain

(A) Diagram of the known structural elements of Xvelo. Prion-like domain, mutants (4D and 7D) and the fragments of Xvelo (F1–F4) are marked in the figure. (B) mRNAs encoding for Xvelo fragments shown in (A) and Xvelo without fragment 1 (Xvelo-woF1) are in vitro synthesized and micro-injected into stage I oocytes. Oocytes were imaged after overnight incubation in oocyte culture medium (OCM).

(legend continued on next page)

(see [Supplemental Experimental Procedures](#) for details; [Data S1A](#)).

We in vitro synthesized mRNAs encoding for the four Xvelo fragments tagged with GFP and injected them into oocytes at equal concentrations. Each fragment of Xvelo localized differently ([Figure 3B](#)). The C-terminal F4 fragment, which carries the positively charged region, localized to nucleoli. Nucleoli are RNA-rich, thus this localization may reflect predicted RNA binding activity of the F4 fragment ([Figure 3B](#)). The N-terminal F1 fragment, which carries the PLD, localized to the Balbiani body. Its localization pattern was indistinguishable from the full-length protein ([Figures 2A and 3B](#)). An additional fragment lacking the F1 fragment but retaining the rest of the Xvelo protein (Xvelo-woF1) did not localize to the Balbiani body ([Figure 3B](#)). Thus, we conclude that the N terminus, which contains the PLD, is the key region that targets Xvelo to the Balbiani body.

We next designed mutants that would disrupt the propensity of Xvelo for amyloid-like self-assembly. Charged amino acids are strongly disfavored in PLDs, as they interfere with the formation of a hydrophobic nucleus ([Alberti et al., 2009](#); [López de la Paz and Serrano, 2004](#); [Serio et al., 2000](#)). Thus, to create defective PLD mutants of Xvelo, we mutated either four or seven aromatic residues in its PLD to a negatively charged amino acid (aspartate) ([Figure 3A](#)). We call these 4D and 7D mutants, respectively ([Figure 3A](#)). The mutants no longer scored positive in prion detection algorithms ([Figures S3B and S3C](#)). We injected mRNAs encoding for wild-type or mutant versions of Xvelo tagged with GFP into the oocytes and imaged after overnight incubation. We used mRNAs encoding for both fragment 1 (the PLD) and full-length Xvelo to observe the effects of the mutants in the oocytes in case the full-length protein interferes with the assembly pattern of the mutants. Fragment 1 proteins carrying the 4D and 7D mutations still exhibited a partial localization to the Balbiani body, but they also showed a diffuse signal in the cytoplasm, which was not observed for the wild-type protein ([Figures 3C and 3D](#)). The effect was much stronger for the 7D mutant, which was barely enriched in the Balbiani body ([Figures 3C and 3D](#)). Full-length proteins behaved similar to their F1 counterparts ([Figures 3D and S3D](#)). Thus, we conclude that the PLD of Xvelo is essential and sufficient for Balbiani body localization, and the two conserved motifs enriched for aromatic residues provide an essential structural role in Xvelo targeting.

To investigate whether the mutations in the Xvelo PLD change the association dynamics of the GFP-tagged constructs with the endogenous Xvelo matrix, we performed FRAP on wild-type and mutants. Fragment 1 had a slow turnover rate, similar to the full-length protein ([Figures 2C, 2D, 3E, 3F, S3E, and S3F](#)). However,

the mutants recovered from photobleaching significantly faster than wild-type; the 100% recovery time was 10 min for 4D and 3 min for 7D mutants after photobleaching ([Figures 3E and 3F](#)). (This should be compared to the wild-type, which recovers to 20% after 1 hr in [Figures 2C and 2D](#).) The recovery times of the full-length mutants were similar to the F1 fragments ([Figure S3F](#)). These results suggest that the PLD drives the association of Xvelo with the pre-assembled Xvelo in the matrix.

If the mutants change the association dynamics of Xvelo with the matrix, they might impose a dominant-negative effect (i.e., inhibit the assembly of wild-type Xvelo into the matrix). To test this, we co-injected mRNAs encoding for full-length Xvelo-WT-mCherry and GFP-tagged F1 wild-type and mutants at equal concentrations and imaged by live confocal microscopy the next day. Wild-type Xvelo co-localized with the F1 fragment in the Balbiani body, and there was little signal in the cytoplasm ([Figure 3G](#)). However, in the presence of the 4D mutant, the wild-type Xvelo also formed discrete small aggregates in the cytoplasm ([Figure 3G](#)). The 7D mutant also increased the amount of diffusely localized wild-type Xvelo, but it did not cause the punctate localization pattern seen with the 4D mutant ([Figure 3G](#)). The difference between the mutants may be due to the fact that the 4D mutant binds more strongly to wild-type Xvelo because of its remaining hydrophobic motif. The same patterns were observed when the experiments were repeated with full-length mutants ([Figure S3G](#)). Thus, we conclude that mutant PLDs can partially inhibit the self-assembly of wild-type Xvelo, most likely by binding to it and blocking growth into larger assemblies.

Recombinant Xvelo Forms Micron-Scale Networks In Vitro

To test whether Xvelo can form amyloid-like fibers on its own, we analyzed recombinant Xvelo-GFP in vitro. Xvelo-GFP did not express at all in bacteria (data not shown) but expressed well in baculovirus-infected insect Sf9 cells. Purification of Xvelo-GFP from insect cells was challenging due to its strong tendency to aggregate, but we found that addition of 300 mM arginine to Xvelo-GFP solubilized it in an otherwise physiological buffer ([Figure S4A](#)). The guanidino fragment of arginine may function similar to guanidinium ion as a solubilizing agent ([England and Haran, 2011](#); [Tsumoto et al., 2004](#)).

We purified soluble Xvelo-GFP in 300 mM arginine and then diluted into 30 mM arginine, which induced self-assembly ([Figure 4A](#), first panel). Xvelo-GFP self-assembled first into small, then large micron-scale networks over time ([Figure 4A](#), first panel). To test if network assembly required the PLD of Xvelo,

(C) mRNAs encoding for wild-type and PLD mutants of fragment 1-GFP were microinjected into oocytes. Oocytes were incubated overnight and imaged.

(D) Ratio of GFP concentration in the oocyte cytoplasm (Ccy) to the Balbiani body (Cbb) in oocytes injected with mRNAs encoding for indicated proteins. Relative concentrations were calculated by using oocyte or the Balbiani body volume from z stacks and the fluorescent intensity of GFP. Mean values and SEs of 10 oocytes are plotted.

(E) Internal rearrangement of fluorescent wild-type or mutant F1-GFP particles after photobleaching over time.

(F) The fluorescent recovery of photobleached wild-type or mutant F1-GFP in Balbiani bodies in (E) and two other biological replicates for each are shown by quantification of fluorescence in bleached region over time normalized by an unbleached neighboring region.

(G) mRNAs encoding for full-length Xvelo-mCherry wild-type and GFP-tagged fragments (F1-WT-GFP, F1-4D-GFP, and F1-7D-GFP) were in vitro synthesized. F1-WT-GFP or mutants were mixed with equal amounts of full-length Xvelo-mCherry mRNA and microinjected into the oocytes. After overnight incubation, the oocytes were imaged by scanning confocal microscopy.

See also [Figure S3](#).

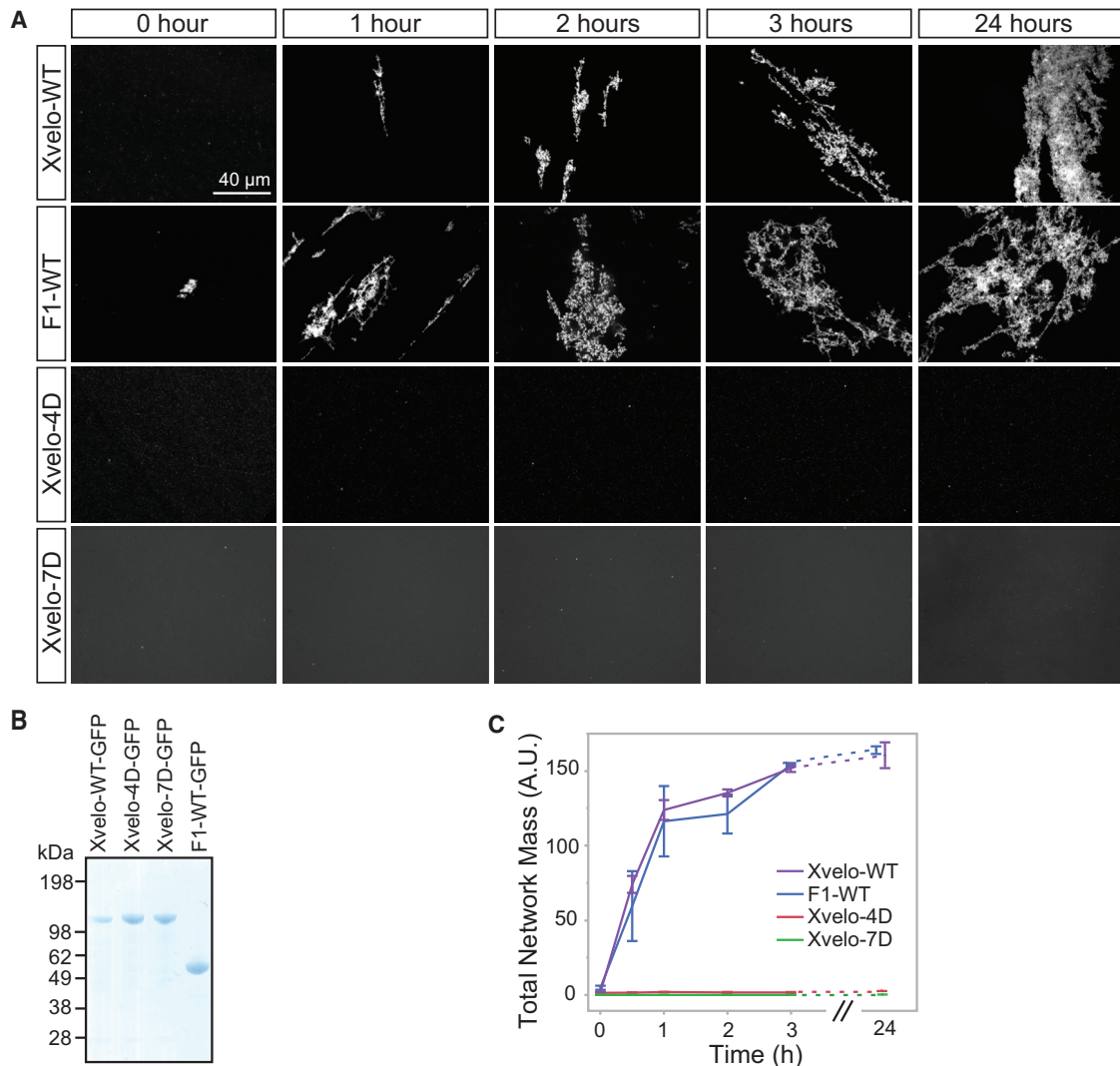


Figure 4. Xvelo Forms Micron-Scale Networks In Vitro

(A) 15 μ M of recombinant Xvelo-GFP, F1-WT-GFP, and full-length mutants (Xvelo-4D-GFP and Xvelo-7D-GFP) were diluted into a low-arginine buffer (30 mM) to promote their self-assembly. The reaction mixtures were incubated at 25°C for the indicated time intervals and squashed under a coverslip to be imaged by spinning-disc confocal microscopy.

(B) Coomassie-stained gels depicting recombinant Xvelo-GFP, Xvelo-4D-GFP, Xvelo-7D-GFP, and F1-WT-GFP.

(C) Quantification of networks in (A). 20 images were taken and stitched together, a threshold was applied, and the network intensities were measured. The integrated intensity of networks per sample at each time point (total network mass) is plotted. Means and SEs of three biological replicates are shown. See also Figure S4.

we expressed and purified the F1 fragment containing the PLD. F1-WT-GFP also formed large networks over time, with striking similarity to networks formed by full-length Xvelo, showing that network assembly is driven by the PLD. We confirmed the critical role of the PLD by showing that two full-length PLD mutants, Xvelo-4D-GFP and Xvelo-7D-GFP, did not form networks (Figure 4A, lower panels) (Figure 4B).

To quantify Xvelo self-assembly in vitro, we tested Xvelo concentrations similar to its physiological concentration (Figure S2C). To balance between the aggregation propensity of Xvelo at high concentrations and the measured physiological concentration of 8 μ M in oocytes if it was uniformly distributed,

we chose a stock concentration of 15 μ M for each recombinant protein so that after 10-fold dilution, the Xvelo concentration in the mixture was 1.5 μ M. We quantified total network mass by summing the fluorescence intensity of networks in an area corresponding to a total of 20 images (Figures 4B and S4B). This analysis further confirmed that the F1 and full-length Xvelo-GFP have indistinguishable kinetics of network formation while the PLD mutants could not form any networks (Figure 4C). Xvelo-4D-GFP formed small precipitates after dilution out of high arginine, but these precipitates remained in solution without forming networks after overnight incubation (Figure 4A, third panel). Xvelo-7D-GFP was completely soluble with no precipitates upon

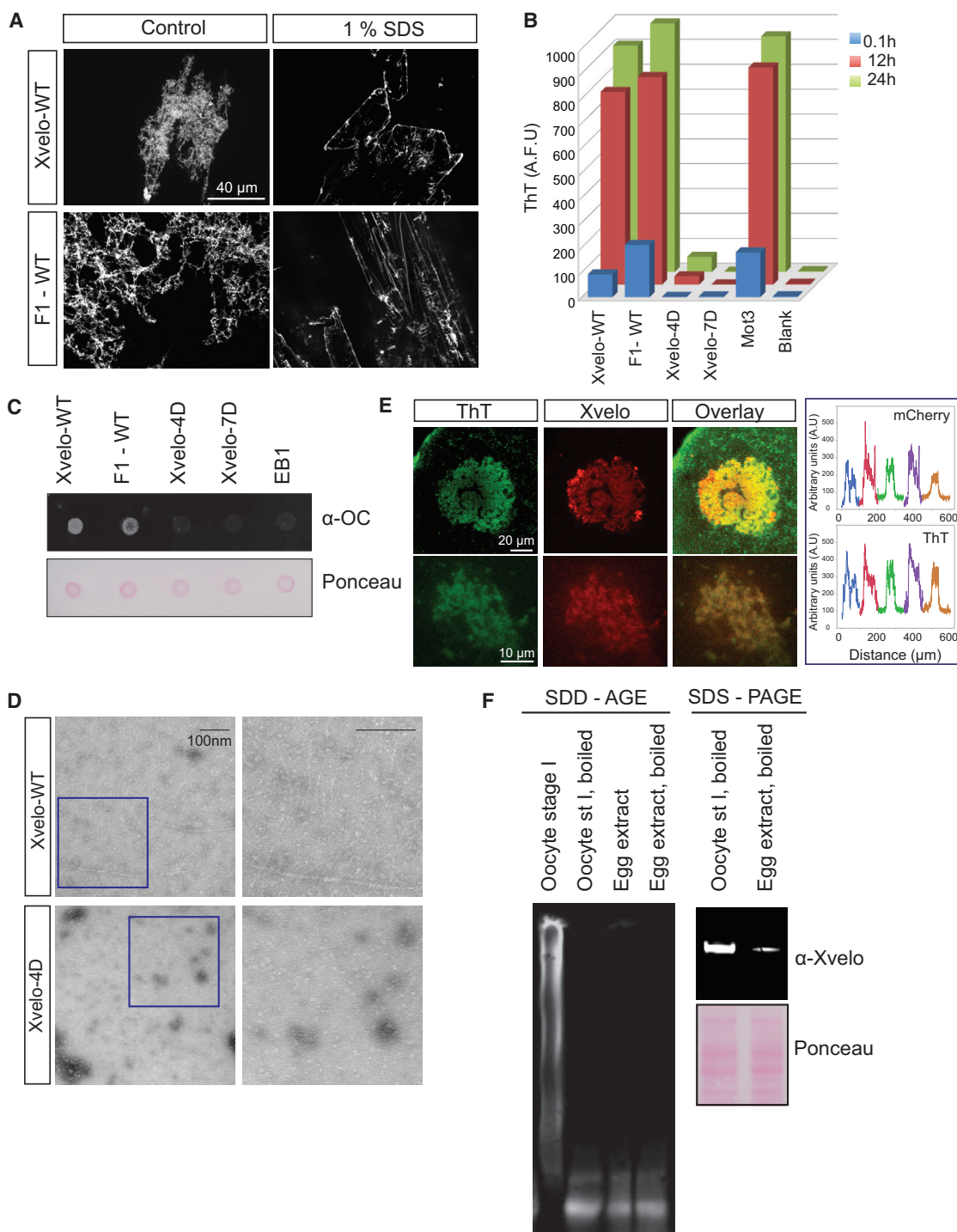


Figure 5. Xvelo Shows Amyloid-like Features In Vivo and In Vitro

(A) SDS was added to Xvelo and F1-WT-GFP networks to a final 1% concentration, and the reactions were incubated at room temperature for 15 min. The resulting mixtures were squashed under a coverslip and imaged by a spinning-disc confocal microscope.

(B) A final concentration of 5 μM of Thioflavin T was added to the wild-type, F1, and mutant network reactions at the indicated time points. Yeast prion Mot3 was used as a positive control, whereas blank was only buffer and ThT. ThT fluorescence was measured (a.u.) by a fluorescence plate reader.

(C) 1 μg RFP-tagged wild-type, F1, and mutant recombinant proteins were dot-blotted on a nitrocellulose membrane and assayed for reactivity with α-amyloid fibril OC. EB1-RFP was used as a negative control.

(D) Negative stain electron microscopy images of the untagged Xvelo-WT and Xvelo-4D self-assembly reactions (scale bars, 100 nm).

(legend continued on next page)

dilution, indicating its complete inability to self-assemble. We also found that the assembly kinetics of Xvelo networks were dependent on Xvelo concentration (Figure S4C). We conclude that Xvelo can form networks on its own in vitro and that the Xvelo PLD is essential and sufficient for the formation of these networks.

Xvelo Networks Exhibit Amyloid-like Properties In Vitro and In Vivo

We next tested whether the Xvelo networks we see in vitro have amyloid-like properties. For this purpose we tested a number of different criteria. First we showed that both Xvelo-WT and F1-GFP networks were resistant to SDS treatment (Figure 5A). We noticed a partial solubilization of the Xvelo network by SDS (Figures 5A and S5A), suggesting the presence of a more stable fiber backbone, which is decorated with more loosely associated Xvelo that can be released by treatment with SDS. Next we monitored acquisition of Thioflavin T (ThT) fluorescence, using RFP-tagged Xvelo, fragment 1 and mutants Xvelo-4D and Xvelo-7D, and the yeast Mot3 prion as a rapidly aggregating positive control (Alberti et al., 2009). F1 and full-length Xvelo networks quickly acquired ThT fluorescence in a PLD-dependent manner (Figure 5B). We supported these findings by showing that full-length and F1 networks were recognized by an anti-amyloid fibril antibody, whereas the mutants or an unrelated protein (EB1-RFP) were not (Figure 5C). Finally, negative-stain transmission electron microscopy (TEM) revealed that Xvelo-WT forms fibers reminiscent of amyloids in vitro in a PLD-dependent manner (Figure 5D).

To confirm that the Xvelo matrix behaves like an amyloid in vivo, we examined whether Xvelo staining coincides with the strong staining of the Balbiani body with ThT. For this purpose, we injected oocytes with mRNA encoding for Xvelo-mCherry, incubated the oocytes with ThT, and imaged them by live confocal microscopy. Xvelo and ThT signals overlapped strongly (Figure 5E). Considering Xvelo is the only enriched protein in the Balbiani body with a PLD and that the endogenous Xvelo concentration exceeds 500 μ M in the Balbiani body, the majority of the ThT signal likely comes from endogenous Xvelo.

Next, we used semi-denaturing detergent agarose gel electrophoresis (SDD-AGE) to check the solubility of Xvelo. SDD-AGE allows the resolution of a wide size range of SDS-resistant aggregates (Alberti et al., 2010; Bagriantsev et al., 2006). We collected stage I oocytes and used egg extracts as a comparison. Indeed, Xvelo formed SDS-resistant aggregates in vivo detected by the SDD-AGE gel. Moreover, it did not form any detectable SDS-resistant aggregates in the mature egg (Figure 5F).

Therefore, because Xvelo forms SDS-resistant, filamentous assemblies that bind ThT in a manner that depends on the presence of its prion-like domain, we conclude that Xvelo forms am-

yloid-like networks in stage I oocytes and in vitro. Although Xvelo is present in the eggs at a detectable, albeit a much lower, concentration of \sim 80 nM (Figures 5F and S2D), it does not form SDS-resistant assemblies in the eggs (Figure 5F). Thus, the amyloid-like characteristics of Xvelo are transient and regulated during development.

Prion-like Domain Specificity for Targeting to the Balbiani Body

To examine whether other proteins with prion-like domains might also be involved in organizing the Balbiani body, we first looked in our mass spectrometry list for other proteins with a PLD. However, apart from the common contaminant yolk protein, vitellogenin, none of the other enriched proteins in the Balbiani body were predicted to contain a prion-like domain.

To investigate whether Xvelo is unique in its ability to form a stable matrix in the Balbiani body, we selected five RNA binding proteins with prion-like domains, namely CPEB3, Dazap1, FUS, hnRNPA1, and Tia1, as well as the aggregation-prone mutant of FUS, FUS-156E, and injected mRNAs encoding for GFP-tagged versions of these proteins into the oocytes. Apart from CPEB3, all these proteins are present naturally in *Xenopus* eggs and oocytes (Wühr et al., 2014). Among these proteins, hnRNPA1, CPEB3, FUS, and FUS156E did not localize to the Balbiani body (Figure 6A). Tia1 and Dazap1 localized to the Balbiani body (as well as the cytoplasm and nucleus, respectively). However, fast turnover rates of both Tia1 and Dazap1 after photobleaching strongly suggest that they do not incorporate into a stable matrix (Figures S6A and S6B). Both Tia1 and Dazap1 are implicated in translational repression by binding to 3' UTRs of mRNAs (Dixon et al., 2003; Steger, 2001), and thus, their localization pattern can be explained by their binding to the repressed mRNAs in the Balbiani bodies upon overexpression.

Our experiments suggest that the PLD of Xvelo has specific features to target to and form a stable matrix in the Balbiani body. Part of the evidence that Xvelo structurally organizes Balbiani bodies is its sequence homology to zebrafish Bucky ball, whose genetics pointed to such a function (Bontems et al., 2009). To test if the two proteins exhibit similar sub-cellular dynamics, we expressed Bucky ball in *Xenopus* oocytes as a GFP fusion and looked at the characteristics of incorporation of Bucky ball into *Xenopus* Balbiani bodies. Bucky ball targeted to the Balbiani body, co-localizing with Xvelo (Figure 6B). Bucky ball turnover time after photobleaching was still a little faster than Xvelo, but of a similar order of magnitude (Figures 6C and 6D). Thus, we conclude that sequence features conserved in the PLDs of Bucky ball and Xvelo are required for targeting to the Balbiani bodies in oocytes. This experiment also provides strong evidence that Bucky ball and Xvelo are functional homologs.

(E) Stage I oocytes were injected with mRNA coding for Xvelo-mCherry and incubated overnight. The oocytes were incubated in 10 μ M ThT, washed twice, and imaged by confocal microscopy. Bottom: zoomed in images. Line scans showing the co-localization of Xvelo-mCherry and ThT stain from five Balbiani bodies were plotted. Each color represents the line scan of a different Balbiani body. We speculate that the outer rim Xvelo-mCherry signal belongs to the newly translated Xvelo-mCherry protein that has just started to form a new, immature matrix and does not yet stain with ThT.

(F) SDD-AGE detects SDS-resistant Xvelo aggregates in vivo. Equal amounts of cytoplasmic extracts of stage I oocytes and mature eggs were loaded onto SDS-PAGE. A five times more amount of egg extracts was loaded for SDD-AGE gels to make Xvelo concentrations comparable between the oocyte and egg extract lanes. Xvelo was detected by an anti-Xvelo antibody.

See also Figure S5.

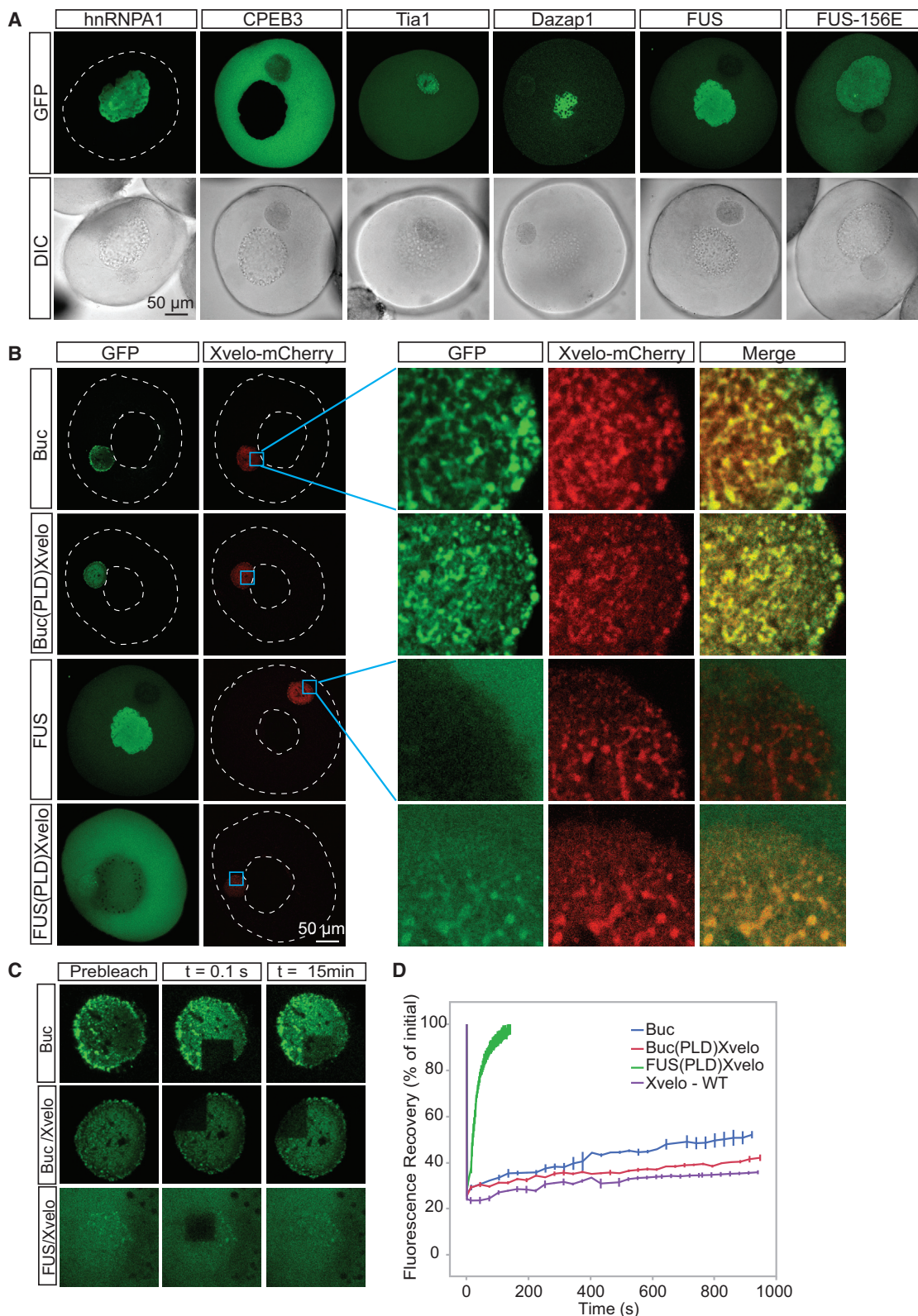


Figure 6. Xvelo Has Unique Properties for Forming a Stable Matrix

(A) mRNAs encoding for GFP-tagged hnRNPA1, CPEB3, Tia1, Dazap1, FUS, and FUS156E were in vitro synthesized and microinjected into the oocytes. After overnight incubation, the oocytes were imaged by laser scanning confocal microscopy.

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To further investigate the specificity of the Xvelo PLD for targeting to the Balbiani body, we swapped the PLD of Xvelo with the PLD of FUS, an unrelated prion-like RNA binding protein, and with the PLD of Bucky ball. The resulting chimeric proteins were named FUS(PLD)Xvelo and Buc(PLD)Xvelo, respectively (Figure S6C). Buc(PLD)Xvelo localized to the Balbiani body, with a FRAP time in between Xvelo-WT and Bucky ball-WT (Figures 6C and 6D). FUS(PLD)Xvelo localized to the cytoplasm and weakly to the Balbiani body (Figure 6B). The Balbiani body-localized protein recovered quickly after photobleaching, with a half-life of ~ 1 min, much faster than that observed for Buc(PLD)Xvelo (~ 2 hr) (Figures 6C and 6D). Taken together, these data provide strong evidence that the prion-like domains of functional homologs Xvelo and Bucky ball have unique features that target and form a stable matrix in the Balbiani body.

We next examined the specificity of PLD interactions in vitro with an aggregation assay that compared the assembly properties of Xvelo and FUS. We repeated previous reports showing that FUS-WT forms liquid droplets in vitro, whereas the aggregation prone mutant FUS-156E forms aggregates (Patel et al., 2015). We mixed Xvelo-RFP with either FUS-GFP or FUS-156E-GFP in vitro in a high-arginine, high-salt buffer and then diluted out the arginine and salt to initiate aggregation. Xvelo-RFP networks and FUS-WT-GFP droplets or FUS-156E-GFP aggregates formed in the vicinity of each other but clearly were separate and did not interact with one another (Figure S6D). Thus, we conclude that PLDs do not aggregate with one another randomly, even when they are in close proximity at high concentration.

Xvelo Binds RNA and Clusters Mitochondria

A key biological function of Xvelo during Balbiani body formation is likely to be the binding and concentration of organelles and RNA through its amyloid-like self-assembly. We looked for direct evidence that Xvelo can form a network that is sufficient to bind and concentrate mitochondria and RNA. To test whether Xvelo-GFP networks can bind RNA, we used a non-specific control mRNA coding for mCherry and an RNA that is enriched in Balbiani bodies, the *Xenopus* homolog of *nanos*, *xcat-2* (Zhou and King, 1996). Both of the RNAs bound to the networks (only mCherry-RNA results are shown). In contrast, the F1 fragment that lacks the C-terminal motif did not bind to either of the RNAs. Thus, Xvelo networks can sequester RNA in a manner that requires a putative RNA-binding domain at the C terminus of Xvelo (Figure 7A), but apparently without any strong RNA sequence preference in vitro.

Germ cells receive a pool of organelles from cyst cells to form Balbiani bodies and become oocytes in mice (Lei and Spradling, 2016). In early-stage oocyte development, these organelles are

clustered into Balbiani bodies by an unknown mechanism. We asked whether Xvelo can cluster organelles on its own in a cell-free system and thus stimulate aspects of Balbiani body reconstruction in vitro. For this, we used an established cell-free system, cytoplasmic extracts from *Xenopus* eggs with intact actin, which contain abundant organelles and RNA, like the germ cell environment prior to Balbiani body formation (Field et al., 2014; Lei and Spradling, 2016). We labeled the mitochondria with MitoTracker and added Xvelo-GFP to the *Xenopus* egg extracts to 1.5 μ M final concentration. Xvelo-GFP self-assembled as expected, and induced co-clustering of mitochondria (Figure 7B). Mitochondrial clustering activity depended on the PLD of Xvelo (Figure 7B, second panel). When we treated these clustered mitochondria with 2 M KCl, mitochondria were still stably bound to, and entrapped by, the Xvelo network (Figure 7B, third panel). We used line-scans to quantify co-clustering of Xvelo assemblies and mitochondria over five frames, in an area extending 2 mm² (Figure 7C). Note the similar line-scans for Xvelo-WT and mitochondria, demonstrating co-clustering, and the lack of mitochondrial clustering with the 4D mutant, which does not self-assemble. Although fragment 1 can still form networks similar to full-length networks in egg extracts, these networks cannot recruit mitochondria (Figure S7A). This allows us to speculate that entrapment of mitochondria is not based on the geometrical properties of Xvelo networks.

As a control, we introduced FUS-WT and FUS-156E to egg extracts. Both FUS-WT and FUS-156E behaved as expected in the egg extracts; WT formed droplets, and 156E formed aggregates. Strikingly, FUS structures were avoided by mitochondria in the egg extracts, as opposed to the clustering we see by Xvelo (Figure 7D). Thus, we conclude that mitochondrial clustering in egg extracts is a specific property of Xvelo that requires its amyloid-like self-assembly.

DISCUSSION

Here, we show that Xvelo, a highly enriched protein in *Xenopus* Balbiani bodies, forms a mitochondria-embedding, SDS-resistant matrix in vivo pervading the entire volume of the Balbiani body in early-stage oocytes. Xvelo has a prion-like domain in its N terminus, which is sufficient and necessary to target and incorporate into the Balbiani body. Pure protein experiments further support our in vivo data that Xvelo has amyloid-like properties. Our key functional experiment, namely clustering of mitochondria by Xvelo in egg extracts, is a reconstruction of aspects of Balbiani body formation. The fact that Xvelo can cluster mitochondria but the prion-like domain mutant does not strongly suggests that this clustering is dependent on its prion-like domain. Thus, these data suggest that the organelle-rich Balbiani body

(B) mRNAs encoding for Xvelo-mCherry and GFP-tagged Bucky ball (Buc), FUS, and the PLD-swap versions of Xvelo, in which the PLD of Xvelo was replaced either by the PLD of Bucky ball (BucPLDXvelo) or the PLD of FUS (FUSPLDXvelo), were injected into the oocytes at equal concentrations and imaged after overnight incubation.

(C) Internal rearrangement of fluorescent Bucky ball-GFP (Buc), and PLD-swap versions of Xvelo, Buc(PLD)Xvelo-GFP and FUS(PLD)Xvelo-GFP, after photobleaching over time.

(D) The fluorescent recovery of photobleached constructs in Balbiani bodies in (C), as well as Xvelo-WT and two other biological replicates for each, is shown by quantification of fluorescence in a bleached region over time normalized by an unbleached neighboring region.

See also Figure S6.

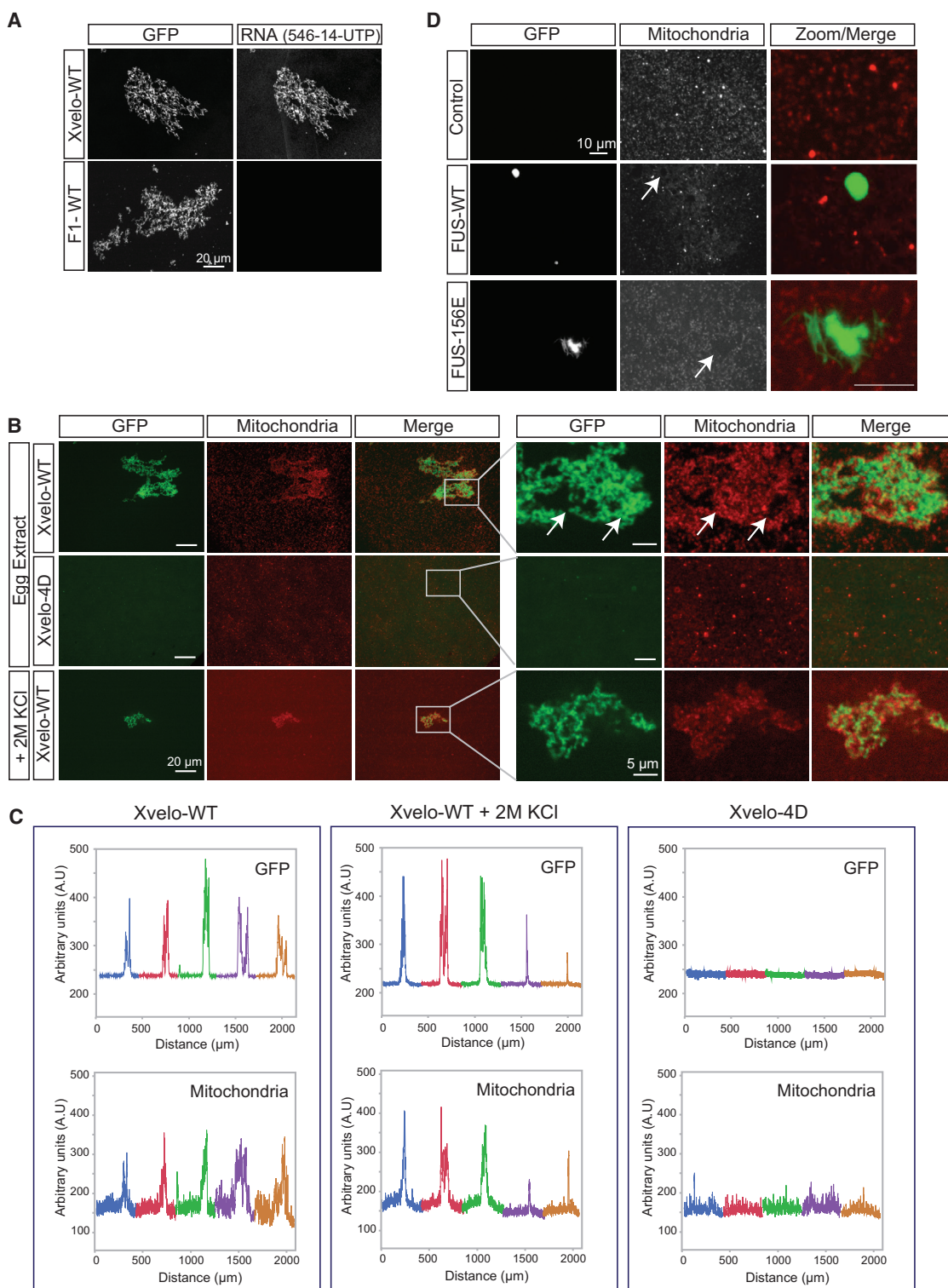


Figure 7. Xvelo-GFP Aggregates Bind to RNA and Cluster Mitochondria

(A) Labeled RNAs, *Xenopus nanos* homolog *xcat-2*, and an mRNA encoding for mCherry protein were prepared using the MEGAscript SP6 kit with ChromaTide Alexa Fluor 546-14-UTP. RNAs were added to pre-assembled networks and imaged with a spinning-disc confocal microscope.

(B) Recombinant Xvelo-WT-GFP or the prion-like domain mutant, Xvelo-4D-GFP, was added to *Xenopus* egg extracts with intact actin. Xvelo-GFP fills the gaps between mitochondria (arrows, compare to Figure 2B). Mitochondria were labeled with Mitotracker Deep Red. Images were taken with a spinning-disc confocal microscope. Bottom: the extracts were diluted with KCl so that the final KCl concentration was 2 M.

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is organized by a functional amyloid into a dense matrix that sequesters mitochondria and other organelles.

The Balbiani body changes its properties during development: it is a stable structure in the early oocyte, and it either disappears in mammals (Hertig and Adams, 1967; Pepling et al., 2007) or dissociates into small dispersed isles, called germ plasm, in the mature oocytes of germ-plasm-containing species. This suggests that its formation and dispersal are regulated. We did not detect any SDS-resistant Xvelo aggregates in egg extracts (Figure 5F), suggesting Xvelo does not form amyloid-like structures in the egg. How this transformation occurs remains unclear. Our preliminary data suggest that Xvelo is extensively phosphorylated in the egg, and most of these sites are not phosphorylated in the oocyte. Regulation by phosphorylation could be a mechanism determining the dispersal of the Balbiani body, perhaps by kinases or phosphatases that are activated at fertilization. Understanding the regulation of the physical state of Xvelo as the oocyte matures will be important to elucidate the fate of the organelles residing in the Balbiani body.

Balbani bodies are found in most vertebrates; mouse Balbiani bodies were identified only recently, but Balbiani bodies in humans were observed decades ago, although they are almost untouched in the literature (Hertig and Adams, 1967; Pepling et al., 2007). Could proteins similar to Xvelo be required for the formation of Balbiani-like structures in other organisms? In zebrafish, *bucky ball* was identified in a maternal effect mutant screen as the only gene that is essential for Balbiani body formation (Dosch et al., 2004). Although the sequence similarity between Xvelo and Bucky ball proteins are poor (Data S1B), these proteins have long patches of intrinsically disordered regions, and score positive in prion detection programs (Data S1C). Oskar is a key protein required to organize *Drosophila* pole plasm (Ephrussi et al., 1991), but no homologs of Oskar have been identified in vertebrates. We found that Oskar also is a disordered protein with a predicted PLD (Data S1D). This suggests that amyloid-like self-assembly of a disordered protein could be an evolutionary conserved mechanism for Balbiani body formation. Strikingly, a recent paper has also linked the formation of large amyloid-like aggregates to gametogenesis in yeast, suggesting amyloid-like mechanisms may be involved in germline specification across kingdoms (Berchowitz et al., 2015).

Despite the low sequence conservation of Xvelo and Bucky ball (Data S1B), key residues in their PLDs are conserved, suggesting that these residues are structurally important and underlie the observed specificity of assembly. Xvelo does not interact with other proteins with PLDs, such as FUS, upon self-assembly in vitro. This is in contrast to the promiscuous behavior of many disease-causing amyloids, which often show cross-seeding interactions and promote mutual nucleation events. We attribute this to the fact that Xvelo self-assembly does not involve an intermediate liquid-like state. We speculate that fast assembly into an inert amyloid-like structure prevents aberrant interactions with

other prion-like proteins, thus reducing the likelihood of a disease condition. We propose that rapid assembly kinetics and high specificity are important driving forces underlying the evolution of functional amyloids.

What are the potential advantages of using an amyloid-like mechanism to form the Balbiani body? One could imagine packing away germline components in amyloid-like structures is protective. The tightly packed structures could prevent other proteins from diffusing into them, such as regulators, thus keeping the organelles in a dormant state. It could act to slow down the diffusion of small toxic molecules generated by mitochondria, which could be damaging. It also provides a novel way to organize the cytoplasm, forming a rigid, giant “body,” in which the organelles are clustered together into one place and kept there for many years. Future studies are likely to provide mechanistic insight into the central question of how the germline of an organism provides young cytoplasm with its complement of organelles in every generation while the somatic cells age and die.

EXPERIMENTAL PROCEDURES

Detailed methods are available in [Supplemental Experimental Procedures](#).

Oocyte Handling and Injections

All experiments using *Xenopus* and zebrafish were done with approval of the Harvard Medical School (HMS) animal care review board. Ovaries were surgically removed from adult female *Xenopus laevis* frogs and treated with 2 mg/ml collagenase 1A (Sigma) in 1 × MMR by gentle rocking, until most of the oocytes were clearly dissociated. Oocytes were later injected with mRNAs encoding for indicated proteins by using a FemtoJet express microinjector (Eppendorf).

Xvelo Protein Purification from Insect Cells

Recombinant versions of MBP-Xvelo-GFP, -RFP, and no-fluorescent tag (for negative-stain electron microscopy studies) were expressed in Sf9 insect cells using the baculovirus expression system. Insect cells were harvested in lysis buffer (50 mM HEPES [pH 7.6], 100 mM KCl, and 1 M arginine). The MBP (maltose binding protein) tag was captured using dextrin Sepharose resin and cleaved off using HRV 3C protease (MPI-CBG, in-house) by incubation overnight on ice.

Microscopy

Differential interference contrast (DIC) and phase contrast microscopy for microinjections, perfusion chambers, and Balbiani body isolations were performed using a standard wide-field epifluorescence Nikon inverted microscope equipped with a Hamamatsu Orca CCD camera and 4×, 10×, and 20× dry objectives. Live confocal microscopy with oocytes was performed using Nikon A1R Laser Scanning confocal equipped with 10× dry and 40× water-immersion objectives. Spinning disc confocal images were taken with a Nikon Ti inverted microscope with Yokagawa CSU-X1 spinning disk confocal with Spectral Applied Research Aurora Borealis modification, equipped with 20× dry, 40×, and 60× oil-immersion objectives.

Semi-denaturing Detergent-Agarose Gel Electrophoresis

Stage I oocyte and egg extracts were prepared according to Field et al. (2014) with intact actin. SDD-AGE was adapted from Alberti et al. (2009). The protein concentrations of the lysates were adjusted and protein samples were mixed with 4 × sample buffer (80 mM Tris, 40 mM acetic acid, 2 mM EDTA, 20% [v/v]

(C) Line scans of Xvelo-WT-GFP and Xvelo-4D-GFP and mitochondria in *Xenopus* egg extracts. Five images were stitched together to have an area spanning larger than 2 mm². Each color represents a different field.

(D) Recombinant FUS-WT-GFP or the aggregation prone mutant, FUS-G156E-GFP, was added to *Xenopus* egg extracts with intact actin. Arrows point to the exclusion zones of mitochondria in the presence of FUS structures.

See also [Figure S7](#).

glycerol, 3% [w/v] SDS, and bromophenol blue) and incubated at room temperature for 15 min before loading onto a 1.8% agarose gel containing 1× TAE and 0.1% SDS. The gel was run in running buffer (1× TAE, 0.1% SDS) at 90 V, followed by wet transfer to nitrocellulose membranes (Amersham Biosciences). Xvelo protein was detected by an anti-Xvelo antibody.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, seven figures, one table, one movie, and one data set and can be found with this article online at <http://dx.doi.org/10.1016/j.cell.2016.06.051>.

AUTHOR CONTRIBUTIONS

E.B. and T.J.M. conceived the project together. E.B. designed the experiments with A.A.H., S.A., and T.J.M. All experiments were performed by E.B., except for mass spectrometry analysis (M.W. and S.P.G.) and electron microscopy (M.C. and M.R.). Protein expression and purification were performed by M.R., D.D., and R.L. The manuscript was written by E.B. with input from A.A.H., S.A., and T.J.M.

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