A brain atlas of synapse protein lifetime across the mouse lifespan

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A brain atlas of synapse protein lifetime across the mouse lifespan

Graphical abstract

**Highlights**

- Brain excitatory synapses differ widely in postsynaptic protein lifetime
- Synapses with diverse protein lifetimes are spatiotemporally distributed
- Age-resilient synapses have long protein lifetimes
- Synapse protein lifetime increases in a model of autism and schizophrenia

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**In brief**

Protein lifetime in synapses is important for brain development and aging, learning, and memory. The authors show that synapses with diverse protein lifetimes are differentially distributed on neurons and in brain regions and that this architecture changes throughout the lifespan and is impacted in a model of autism and schizophrenia.
SUMMARY

The lifetime of proteins in synapses is important for their signaling, maintenance, and remodeling, and for memory duration. We quantified the lifetime of endogenous PSD95, an abundant postsynaptic protein in excitatory synapses, at single-synapse resolution across the mouse brain and lifespan, generating the Protein Lifetime Synaptome Atlas. Excitatory synapses have a wide range of PSD95 lifetimes extending from hours to several months, with distinct spatial distributions in dendrites, neurons, and brain regions. Synapses with short protein lifetimes are enriched in young animals and in brain regions controlling innate behaviors, whereas synapses with long protein lifetimes accumulate during development, are enriched in the cortex and CA1 where memories are stored, and are preferentially preserved in old age. Synapse protein lifetime increases throughout the brain in a mouse model of autism and schizophrenia. Protein lifetime adds a further layer to synapse diversity and enriches prevailing concepts in brain development, aging, and disease.

INTRODUCTION

The lifetime of a protein at its subcellular location is a fundamental determinant of its role in homeostatic and adaptive processes. Physiological and pathological events trigger protein modifications, and the subsequent replacement with a newly synthesized protein reinstates cellular functions to their original state, a process known as proteostasis (Kaushik and Cuervo, 2015; Labbadia and Morimoto, 2015). In addition to homeostatic roles, protein turnover is essential for adapting the levels of proteins to developmental and adaptive changes in gene expression; for example, a reduction in an mRNA can only lead to reduced levels of protein if there is ongoing protein removal. The speed of the underlying mechanisms that produce, deliver, and remove proteins controls how long it takes for these proteostatic and adaptive processes to complete their task. This in turn is directly related to the lifetime of the protein at its site of action, with a short protein lifetime indicative of rapid homeostasis and adaptation.

In brain synapses, protein turnover controls adaptive processes including synaptic transmission, plasticity, and learning (Colledge et al., 2003; Ehlers, 2003; Kato et al., 2005; Patrick et al., 2003). Francis Crick highlighted the importance of synaptic protein lifetime for the duration of memory, reasoning that protein removal erased learning-induced protein modifications (Crick, 1984). An imbalance in the rate of synaptic protein production and removal is thought to underlie learning and other cognitive deficits in genetic forms of autism and intellectual disability (Labbadia and Morimoto, 2015; Louros and Osterweil, 2015);
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A reduction in the rate of protein removal is also thought to underlie the synaptic and behavioral deficits in old age (Kaushik and Cuervo, 2015; Sabath et al., 2020; Santra et al., 2019) and contribute to the accumulation of toxic protein species, such as aggregates of β-amyloid and Tau, that contributes to synaptic pathology in Alzheimer’s, Parkinson’s, and other neurodegenerative diseases (Helton et al., 2008; Kaushik and Cuervo, 2015; Tai et al., 2012; Vilchez et al., 2014).

Excitatory synapses are highly diverse in molecular composition and morphology (Broadhead et al., 2016; Cizeron et al., 2020; Zhu et al., 2018). The synapse types and subtypes have different spatial distributions on dendrites and between neuron types and are found in characteristic compositions in brain regions and subregions, forming a highly organized synaptic architecture that changes throughout the lifespan (Cizeron et al., 2020). The spatiotemporal differences in synapse molecular composition raise the possibility that synapses may also differ in their protein lifetime.

The lifetime of endogenous proteins has been measured in bulk brain synapse preparations, which lack information about synapse diversity (Cohen et al., 2013; Dörbaum et al., 2018; Ehlers, 2003; El-Husseini et al., 2002; Fornasiero et al., 2018; Heo et al., 2018; Price et al., 2010). Measurements of protein lifetime at single-synapse resolution have been limited to small populations of neurons and exogenously expressed proteins (Gray et al., 2006; Steiner et al., 2008; Villa et al., 2016). To overcome these limitations, we have adapted HaloTag self-labeling protein tag technology to visualize and quantify the lifetime of endogenous synaptic proteins in individual excitatory synapses in all parts of the mouse brain at any age. HaloTag technology (Los et al., 2008) has been widely used to label different proteins in many cellular and tissue contexts, including studies of protein turnover and lifetime. The system involves two components: the genetically encoded HaloTag protein domain and the HaloTag ligand, a small molecule that binds to the HaloTag domain. The ligand forms a highly stable covalent bond that is irreversible under physiological and non-physiological conditions (Encell et al., 2012; England et al., 2015; Hoelzel and Zhang, 2020; Los et al., 2008; Xue et al., 2015). The binding site for a chloralkane linker is located within a deep pocket (~15 Å) away from the protein surface, protected from the cellular milieu (Los et al., 2008). Coupling the HaloTag ligand to fluorophores enables irreversible labeling and visualization of tagged proteins, and observing the reduction in fluorescent signal over time enables the lifetime of the protein to be measured.

We engineered the mouse genome to express the HaloTag protein domain fused to endogenous PSD95 (PSD95-HaloTag mice) (Figures 1A, S1A, and S1B). PSD95 (also known as DLG4) is a highly abundant scaffold protein of excitatory synapses that interacts with and organizes more than 100 other postsynaptic proteins, including NMDA (N-Methyl-D-aspartic acid) and AMPA (α-Amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid) subtypes of neurotransmitter receptors (Fernández et al., 2017, 2009; Frank et al., 2016; Husi et al., 2000). PSD95 is essential for synaptic plasticity and learning (Carlisle et al., 2008; Fernández et al., 2017; Fitzgerald et al., 2015; Frank et al., 2016; Migaud et al., 1998; Nithianantharajah et al., 2013). Injecting a fluorescent HaloTag ligand into PSD95-HaloTag mice labels excitatory synapses with a fluorescent date-stamp. The lifetime of PSD95 in a synapse is then measured by visualizing the duration of the labeled protein. We combined this labeling strategy with our synaptome mapping pipeline technology (SYNMAP), which is capable of quantifying the fluorescent signals from billions of individual synapses across the mouse brain (Cizeron et al., 2020; Zhu et al., 2018). We report the creation of a single-synapse resolution atlas (Bulovaite et al., 2021a) of PSD95 protein lifetime across the brain in young, mature, and old mice and in a model of neurodevelopmental disorders.

RESULTS

Genetic labeling and visualization of postsynaptic protein lifetime

In PSD95-HaloTag mice, PSD95 expression, protein complex assembly, and the physiology of synaptic transmission and synaptic plasticity are normal (Figure S1). The bond between PSD95-HaloTag and a fluorescent HaloTag ligand (tetramethyl-rhodamine-Halo, or TMR-Halo) proved to be extremely stable (Figure S1G), consistent with previous studies (Encell et al., 2012). To irreversibly label and visualize the PSD95-HaloTag in vivo, we injected a cell- and blood-brain-barrier-permeable fluorophore ligand (silicon-rhodamine-Halo, or SiR-Halo) (Lukinavicius et al., 2013; Masch et al., 2018) into the tail vein of PSD95-HaloTag mice and prepared sections of brain tissue for imaging on a spinning disk confocal microscope (pixel resolution 84 nm and optical resolution, as measured by full width at half maximum, ~330 nm in x-y and ~620 nm in z) (Figure S2). By examining the persistence of labeling after injection, we could identify the synapses and brain regions with different PSD95 lifetimes (Figure 1B).

Imaging the brain of 3-month-old PSD95-HaloTag mice 6 h (day 0) after injection revealed widespread labeling (Figures 1C and S3A), whereas none was detected in injected wild-type mice (Figure S3B). 300 nmol of injected SiR-Halo was sufficient to saturate the majority of HaloTag-binding sites (Figure S3C). Higher doses of SiR-Halo were not feasible due to toxic effects of the DMSO solvent (Thackaberry et al., 2014). We confirmed that the SiR-Halo ligand quantitatively labeled all brain regions by injecting it into compound heterozygous PSD95HaloTag/EGFP mice, quantifying the SiR-Halo and EGFP synaptic puncta density in 110 brain regions, and then testing their correlation (R = 0.976, p < 0.0001) (Figures S3D and S3E). To better understand how long the SiR-Halo ligand remains in the body after injection, we performed a pharmacokinetic analysis (STAR Methods; Figure S4). The peak concentration of SiR-Halo was detected 1 min after injection (blood, 4.6 μM; brain, 0.08 μM), and by 15 min the concentration was 50% of the peak value in both blood and brain. By 24 h, SiR-Halo was undetectable in blood and brain (~10 nM detection threshold).

Spatial diversity in synaptic protein lifetime

Images collected over the course of 2 weeks revealed a loss of labeling in all brain regions, with different regions showing different rates (Figures 1C and S3A). Synapses with the longest PSD95 protein lifetime were concentrated in the isocortex (neocortex) and hippocampal formation (HPF) (Figures 1C and S3A). Examination of images at single-synapse resolution

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Figure 1. Visualizing synapse protein lifetime across the mouse brain

(A) Experimental flow. Homozygous PSD95-HaloTag mice were injected with a Halo ligand, sagittal brain sections were imaged using spinning disk confocal microscopy, and synaptic puncta were analyzed and mapped.

(B) Schematic of PSD95-HaloTag labeling in the postsynaptic terminal of synapses with short or long protein lifetime.

(C) SiR-Halo fluorescence labeling at five time points post-injection, showing whole brain, HPF, and layers of the isocortex. Key shows regions and layers (Table S1).

(D and E) High-magnification representative images of puncta fluorescence decay in layers of the motor cortex (D) and in stratum radiatum of CA1sr, CA2sr, and CA3sr subregions of the HPF (E). For brain region/subregion abbreviations, see Table S1. Scale bars: 2 mm (C, whole brain), 500 μm (C, HPF, and CTX), 5 μm (D and E).
revealed that each brain region is composed of populations of synapses with different PSD95 lifetimes (Figures 1D, 1E, and S5), some with long (weeks) and others with short (hours or days) protein lifetimes, which we refer to as LPL and SPL synapses, respectively. These synapse populations were spatially organized. For example, within the laminar organization of the isocortex there was a clear gradient from superficial (layer 1) to deep (layer 6) layers, with synapses with the longest PSD95 lifetime located in layer 1 (Figures 1C, 1D, S3A, and S5). In the HPF, the dendritic fields of pyramidal neurons in the CA1 exhibited synapses with longer PSD95 lifetimes than those in CA2 and CA3 (Figures 1C, 1E, S3A, and S5). This pattern was evident in both basal and apical dendrites, indicating that synaptic protein lifetime is determined, at least in part, by cell-wide mechanisms that differ between pyramidal neuron subtypes (Figures 1C and S5). Synapses in the dendritic fields of granule cell neurons in the dentate gyrus (DG) and olfactory bulb had short PSD95 protein lifetimes, further suggesting cell-type-specific turnover mechanisms (Figures 1C and S5). Comparison of the large CA3 thorny excrescence synapses and those in the polymorphic layer of the DG with the smaller CA1 synapses showed that the largest synapses of the HPF (Broadhead et al., 2016; Cizeron et al., 2020; Harris and Weinberg, 2012; Zhu et al., 2018) do not have the longest protein lifetimes (Figure S5).

A brainwide synaptome atlas of protein lifetime in excitatory synapses

To quantify PSD95 protein lifetime in brain subregions we calculated its half-life using single-synapse resolution data and SYNMAP synaptome mapping technology (Zhu et al., 2018). PSD95-HaloTag mice were injected with SiR-Halo ligand and parasagittal brain sections imaged at post-injection time points (6 h, 1.5, 3, 7, and 14 days) (Figures 1C and S3A). The density and intensity of SiR-Halo-labeled synaptic puncta were quantified in 12 overarched brain areas (isocortex, olfactory areas, HPF, cortical subplate, striatum, pallidum, thalamus, hypothalamus, midbrain, pons, medulla, and cerebellum) and 110 subregions registered to the Allen Reference Atlas (Table S1). By fitting a single-phase exponential decay function to the data for every subregion we calculated the average PSD95 puncta density (PSD95 density\(_{1/2}\)) and intensity (PSD95 intensity\(_{1/2}\)) half-lives and plotted the values in brain maps (Figures 2A, 2B, S6A, and S6B). PSD95 density\(_{1/2}\) is derived from the reduction in the number of SiR-Halo-positive puncta, whereas PSD95 intensity\(_{1/2}\) is derived from the reduction in SiR-Halo signal within labeled synapses. The two half-life measures were highly correlated (R = 0.9088, p < 0.0001; Figure S6C). We also tested if SiR-Halo labeling and puncta detection efficiency in brain subregions influenced protein lifetime measurements and found no correlation (Pearson’s correlation, R = 0.1337, p = 0.1638). The half-life density (PSD95 density\(_{1/2}\)) and intensity (PSD95 intensity\(_{1/2}\)) datasets were used to create the Protein Lifetime Synaptome Atlas (Bulovaite et al., 2021a).

Dendritic and subregional distribution of synapses with different PSD95 lifetimes

We observed a 10-fold difference in PSD95 density\(_{1/2}\) between the subregions with the longest (layer 1 of the motor cortex, 12.1 days) and shortest (olfactory bulb glomerular layer, 1.2 days) half-life (Figures 2A, 2C, S6A, and S7A). Comparison of PSD95 density\(_{1/2}\) for the 12 overarched brain areas showed that cortical structures are populated by synapses with the longest protein lifetimes, in contrast to subcortical structures which predominantly comprise synapses with the shortest protein lifetimes (Figures 2A, 2C, S6A, S6B, and S7B–S7D). All regions of the isocortex exhibited a gradient, with a 2.4-fold range in PSD95 density\(_{1/2}\) from layer 1 (10.6 days) to layer 6 (4.5 days) (Figures 2A, S6A, S7E, and S7F). Similarly, the subregions within the HPF with the shortest PSD95 density\(_{1/2}\) were in initial pathways of the trisynaptic circuit (DG, 3.5 days; CA2, 1.9 days; CA3, 3.0 days), whereas those with the longest PSD95 density\(_{1/2}\) were in the latter part of the circuit in the CA1 field (5.5 days) and subiculum (6.4 days) (Figures 2B, S7G, and S7H).

Although dendrites contain protein synthesis and degradation machinery (Biever et al., 2020; Bingol and Schuman, 2006; Ostroff et al., 2002; Steward and Levy, 1982), it is not known whether synapse protein lifetime varies across the dendritic tree. We quantified PSD95 density\(_{1/2}\) in a series of parallel windows from the pyramidal neuron soma in the CA1 to the distal basal dendrites in stratum oriens (so) and to the distal apical dendrites in stratum radiatum (sr), stratum lacunosum (sl), and stratum molecular (sm) (Figure 2D). In apical dendrites, a gradient of increasing PSD95 protein lifetime was observed (Figure 2D, a–d: linear regression, R^2 = 0.1571, p = 0.007). We next examined segments of the apical dendrite including the sr, sl, and sm, which each receive distinct afferents (from CA3, entorhinal cortex layer 2 and entorhinal cortex layer 3, respectively). A positively sloped gradient in PSD95 lifetime was found in sr (Figure 2D, a and b: linear regression, R^2 = 0.5110, p < 0.0001). There was a significant reduction in PSD95 lifetime between distal sr and the sl (Figure 2D, b and c: Welch’s t test, p = 0.0151), which was not due to differences in synapse size (Figure S8). From the sl to the distal end of the sm we observed a positively sloped gradient (Figure 2D, c and d: linear regression, R^2 = 0.6234, p = 0.0005). PSD95 puncta density followed a similar gradient to that of PSD95 lifetime, but since half-life measure is independent of absolute puncta numbers at day 0, differences in puncta density cannot affect half-life values (Figure S8). In the basal dendrites of so there was an initial gradient in PSD95 protein lifetime before leveling and reversing in distal dendrites. These results show that synapse protein lifetime differs within the dendritic tree of neurons and that synapse protein lifetime is influenced by distance from the soma and by afferent inputs.

PSD95 turnover occurs in stable dendritic spines

Dendritic spines can be structurally stable for long or short durations (Grutzendler et al., 2002; Holtmaat et al., 2005; Trachtenberg et al., 2002; Zuo et al., 2005). We asked whether protein turnover occurs within stable spines or whether the loss of HaloTag labeling might reflect spine elimination. To visualize dendritic spines housing PSD95 in individual apical dendrites, we dye-filled CA1 pyramidal neurons from PSD95-HaloTag mice injected with SiR-Halo (Figures 3A and S9A). Consistent with previous results that show PSD95 is expressed in a subset of larger, more stable synapses (Cizeron et al., 2020; Fortin et al., 2014; Santuy et al., 2020; Zhu et al., 2018), a subset of spines...
expressed PSD95 at day 0, and labeled spines were still visible 10 days after SiR-Halo injection. If PSD95 were being turned over in synapses, we reasoned that we could observe both “old” and “new” protein using a two-step labeling procedure.

Three days after injecting mice with SiR-Halo we labeled brain sections with TMR-Halo and found that the two labels were colocalized in many synapses (Figures 3B and S9B).

To simultaneously visualize PSD95 turnover and spine turnover in the brain of living PSD95-HaloTag mice, we monitored individual synapses in layer 1 of somatosensory cortex through a brainwide atlas of synapse protein lifetime (Figures 2A and S9A).
Figure 3. PSD95 turnover in dendritic spines

(A) 3D reconstructed dye-filled CA1 neurons and SiR-Halo-positive puncta. (i–iv) CA1 apical dendrites (stratum radiatum) intracellularly injected with Alexa 488 (green) at day 0 (i and ii) and at day 10 post-injection of SiR-Halo (red) (iii and iv). (v and vi) Higher-magnification of (i) and (iii), respectively, showing dendritic spines with SiR-Halo-positive puncta (white arrows) and without SiR-Halo signal (blue arrows). Scale bar: 5 µm (i–iv), 1.5 µm (v and vi).

(B) Colocalization (orange) of SiR-Halo (red) injected at day 0 and post-fixation-applied TMR-Halo (green) at day 3 in synapses across layers of the motor cortex. Scale bars: 5 µm.

(C) Schematic of the cranial window implant above the somatosensory cortex.

(D) Bright-field image (left) of the superficial vasculature and, superposed, a fluorescence image of YFP-labeled axons and dendrites in cortical layer 1 that are used as fiducial points for tracking SiR-Halo ligand puncta over time, as shown for day 0 (center) and day 7 (right).

(E) A YFP-labeled dendrite (top) containing SiR-Halo-labeled puncta (bottom), repeatedly imaged over days to weeks. SiR-Halo puncta can be tracked in stable and dynamic spines (e.g., P1–P5) and in the immediate surrounding of the dendrite (e.g., P6).

(F) The mean ± SD fluorescence intensities of individual SiR-Halo puncta (n = 295 from 3 mice Psd95HaloT/HaloTag mice) normalized to the first imaging time point. Red line, a single exponential fit of the means (λ = 0.12/day; t1/2 = 5.8 days).

(G) The measured fluorescence intensity over time of the SiR-Halo puncta examples in (E).
cranial window (Holtmaat et al., 2009) over weeks after injection of SiR-Halo (Figure 3C). Viral vectors expressing YFP were used to visualize dendritic spines and estimate spine size (STAR Methods; Figures 3D and 3E) (Holtmaat et al., 2005). In individual synapses we observed decay profiles that were similar (Figure 3F) to the population data (Figure 2). A comparable overall decay of fluorescence was observed in regions imaged only once over the course of the experiment (Figure S9C), excluding photobleaching as a main factor in driving the fluorescence decay. In line with the dye injection study (Figure 3A), not all spines expressed PSD95 to a similar extent (Figures 3E and 3G), although in general SiR-Halo fluorescence intensities correlated well with spine size (Figure S9C) (Gray et al., 2006). Even within single dendrites, spines variably retained the SiR-Halo label. Label retention was weakly related to spine size (Figure S9C), and spines observed to disappear often had low levels of PSD95 or rapid loss of labeling (Figures 3E and 3G), consistent with published observations (Cane et al., 2014; Gray et al., 2006). Together, these data indicate that structurally stable spines differ in their PSD95 content and lifetime, even within the same dendrite.

**Synapse classification by protein lifetime**

The differences in protein lifetime between individual synapses raise questions about the diversity of synapses that comprise the synaptome. In previous work we used a data-driven approach to classify synapses from across the whole brain and lifespan based on their “static” protein composition (Cizeron et al., 2020; Zhu et al., 2018). Using two markers (PSD95eGFP and SAP102mKO2), each of which assembles a set of postsynaptic proteins into distinct molecular complexes (Frank et al., 2016), we classified synapses into three types: type 1 express PSD95, type 2 express SAP102, and type 3 express both markers. Each of these types was further classified using morphological criteria into a total of 37 subtypes (Zhu et al., 2018). To determine whether the different types and subtypes of synapses have characteristic protein lifetimes, we combined synaptome mapping of PSD95-HaloTag with PSD95eGFP and SAP102mKO2 using triple knockin mice (Psd95HaloTag/eGFP; Sap102mKO2/y). 6-month-old triple knockin mice were injected with SiR-Halo and brain sections collected at day 0 and 7. To test that the estimation of half-life using two time points accurately reflected measurements made with five time points (Figure 2), we correlated the values obtained in all brain subregions and found a very high correlation (puncta density, R = 0.9789, p < 0.0001; puncta intensity, R = 0.9506, p < 0.0001) (Figure S10A). SiR-Halo-positive synapses were categorized using PSD95eGFP and SAP102mKO2, allowing us to determine the PSD95 lifetime of two synapse types (type 1 and type 3) and 30 of the 37 known subtypes (all except subtypes 12–18, which do not express PSD95) in all brain areas. Ranking the synapse subtypes by their PSD95 protein lifetime from brainwide data (Figure 4A) and individual regions (Figure S10B) showed that both type 1 and type 3 synapses contain subtypes with short and long protein lifetimes. Thus, populations of synapses with distinct molecular compositions are further divisible into subtypes with varying protein lifetimes. Synapse subtype 2 maintained the most (96.7%) and subtype 6 the least (23.3%) label after 7 days (Figure 4A). To visualize the potential duration that any synapse subtype could retain copies of PSD95 we plotted exponential decay functions (Figure 4B). This revealed that several synapse subtypes in the adult mouse could retain copies of PSD95 for many months and, potentially, as long as the natural lifespan of the mouse (Burt, 1940; Hamilton, 1937).

**Building blocks of synaptome architecture**

The protein lifetime of each synapse subtype was independent of its location in the brain, with subtype lifetime rankings maintained across brain regions (Figures 4C and S10B). This indicates that synapse subtypes with different protein lifetimes are general building blocks of whole-brain architecture. Furthermore, these findings suggest that the spatial distribution of synapse subtypes with different protein lifetimes underpins the half-life measurements from individual brain regions. To confirm this, we generated a self-organizing feature map (SOM), which reduces the high dimensionality of synaptome maps of the 37 subtypes to a two-dimensional representation (Figures 4D and S10C). The SOM shows a systematic organization in which anatomically related regions become closely clustered owing to their similarity in subtype density (Figure S10C). Consistent with our hypothesis, we observe a systematicity in half-lives when subregion half-life values are shown (Figure 4D).

LPL synapses are particularly interesting because they could potentially maintain PSD95 modifications for long durations. When grouped together, the five subtypes with longest protein lifetimes (subtypes 2, 3, 34, 35, and 5) represented 6.7% of PSD95-positive excitatory synapses in the whole brain; however, there were marked regional and subregional differences (Figures 4E and S10D), with a 15-fold range across the main brain areas: isocortex (11.9%), striatum (5.9%), cortical subplate (4.4%), cerebellum (3.9%), HPF (3.8%), olfactory areas (3.0%), midbrain (2.4%), hypothalamus (2.2%), medulla (2.2%), pons (2.1%), pallidum (1.8%), and thalamus (0.8%). The subregion with the highest percentage of these LPL synapses was layer 2-3 of the somatosensory cortex (17.2%), while the lowest percentage was found in the CA2 sl-moleculare (0.2%). Isocortex
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regions had the highest percentages of LPL synapses, with superficial layers showing 4- to 5-fold more LPL synapses than the deepest layers. Within the HPF, the subiculum had the highest percentage (9.6%) of LPL synapses of all subregions, followed by CA1sr (5.8%). These results show that the neocortex (isocortex), which is the most recently evolved region of the cortex as compared with the ancient allocortex (olfactory areas and HPF) (Rakic, 2009), contains higher numbers of LPL synapses, with the superficial layers of the neocortex being particularly well populated.

**Synapse protein lifetime increases throughout the lifespan**

Synapse protein synthesis and turnover are thought to be important during brain development and aging (Hipp et al., 2019; Kaushik and Cuervo, 2015; Labbadia and Morimoto, 2015; Yerbury et al., 2016); yet, to the best of our knowledge, there have been no studies of synaptic protein turnover comparing childhood, mature adulthood, and old age (Cizeron et al., 2020).

These lifespan studies revealed that PSD95 protein lifetime increases in the majority of brain regions from birth to old age (Figures 5A, 5B, S11A, and S11B). At 3 weeks there was a paucity of subregions showing long protein lifetimes, with only the most superficial layers of the neocortex and CA1 sr/sm showing a $\text{PSD95}^{\text{density}_{1/2}}$ above that of other regions (Figure 5A). Between 3 weeks and 3 months, the $\text{PSD95}^{\text{density}_{1/2}}$ increased in all subregions, with the greatest increases in subregions of the isocortex and cerebellum granular layer (Figures 5A and 5B). We also examined age-dependent changes in PSD95 protein lifetime in the dendritic tree of CA1 pyramidal cells (Figure 5C). At 3 weeks there were nascent gradients in both basal and apical dendrites formed from synapses with short protein lifetimes, and their protein lifetime increased ~5-fold by 3 months. To uncover which synapse subtypes contribute to this increase we measured the effect size (Cohen’s $d$) for the change in density for all subtypes between 3 weeks and 3 months. This identified subtype 2, which is the LPL subtype with the longest protein lifetime, as significantly increased across many brain regions (Figures S11Ci and S11Cii). By contrast, no increase was observed for subtype 34 (a type 3 LPL subtype; Figure S11Ciii), indicating that LPL synapses with different molecular compositions are non-redundant and have distinct developmental trajectories. During development, brain regions differentiate from each other by the acquisition of sets of diverse synapses (Cizeron et al., 2020), and we asked whether brain regions differentiate with respect to their synapse protein lifetime compositions. Comparison of the $\text{PSD95}^{\text{density}_{1/2}}$ between all subregions showed a high similarity at 3 weeks that dramatically decreased by 3 months (Figure 5D). Together, these data show that from weaning to maturity the mouse brain accumulates synapses with long protein lifetimes and that brain regions acquire a distinct and thus specialized compositional signature of synapses with different protein lifetimes.

We next examined how the brain changes with old age. Between 3 and 18 months, the regional $\text{PSD95}^{\text{density}_{1/2}}$ increased in most areas of the brain (Figures 5A and 5B). By contrast, aging did not influence the spatial organization of synapses on the dendritic tree of CA1 pyramidal neurons (Figure 5C). Previously, we showed an overall loss of excitatory synapses between 3 and 18 months but with a subtype specificity to the retention or loss (Cizeron et al., 2020). Here, we found that the increased regional $\text{PSD95}^{\text{density}_{1/2}}$ in old age reflects the preservation of LPL synapses. Although most synapse subtypes decrease between 3 and 18 months (Cizeron et al., 2020), the LPL synapses (subtypes 2, 34, 3, and 5) were retained (Figures 5E and S11Civ–vi). Subtype 2 was preferentially retained in neocortex (isocortex), whereas in allocortex (olfactory cortex and HPF) subtypes 34, 3, and 5 were retained, revealing that LPL synapses with different molecular compositions are resilient to aging and populate distinct brain regions.

**Increased synaptic protein lifetime in a model of autism and schizophrenia**

Mutations in the DLG2 gene cause autism and schizophrenia in humans (Fernández et al., 2009; Fromer et al., 2014; Ingason et al., 2015; Kirov et al., 2012; Nithianantharajah et al., 2013; Purcell et al., 2014; Reggiani et al., 2017; Ruzzo et al., 2015; Walsh et al., 2008), and the Dlg2 mutant mouse model recapitulates the impairments in attention, learning, and cognitive flexibility found in humans (Nithianantharajah et al., 2013). Dlg2 encodes PSD93, a postsynaptic scaffold protein that binds to PSD95, and together these proteins assemble a ternary complex with the NMDA receptor subunit GluN2B (Frank et al., 2016). To test if Dlg2 mutations affect the synaptome map of PSD95 lifespan we bred a cohort of PSD95-HaloTag mice containing heterozygous (Dlg2$^{+/+}$;Psd95$^{HaloTag/+}$) or homozygous (Dlg2$^{−/−}$;Psd95$^{HaloTag/+}$) null alleles of Dlg2 (Figures 6 and S12). Synaptome maps showed a robust increase in PSD95 protein lifetime across all 110 brain regions of both heterozygous and homozygous Dlg2 mutant mice as compared with PSD95-HaloTag control mice, with a gene-dosage-dependent effect. PSD95 half-life

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**Figure 5. Synapse protein lifetime changes across the lifespan**

(A) PSD95 puncta half-life across 110 mouse brain subregions in PSD95-HaloTag mice at 3 weeks, 3 and 18 months for the whole brain (top row) and HPF (bottom row). $n = 7–10$ Psd95$^{HaloTag/+}$ mice at each time point and age group.

(B) Percentage increase in puncta half-life between 3 weeks and 3 months (left brain map) and between 3 weeks and 18 months (right brain map).

(C) Mean fraction of puncta remaining at day 7 compared with day 0 (± SD) is plotted across basal and apical dendrites of the CA1 dendritic tree at 3 weeks, 3 and 18 months ($n = 3$ mice per group).

(D) Matrix of half-life similarities between pairs of subregions (rows and columns) for the 3 weeks and 3 months age groups. Brain subregions are listed in Table S1.

(E) Heatmap showing change (Cohen’s $d$ effect size) in synapse subtype (subtype number shown at top) density across brain subregions between 3 and 18 months. Subtypes are ranked with longest lifetime on the left and shortest on the right. * $p < 0.05$ (Benjamini–Hochberg correction). See Figure S11C for detailed heatmaps.
increased up to 210% in heterozygous and 380% in homozygous Dlg2 mutants. Most affected were subregions of the isocortex and HPF CA1, which are areas that show dysfunction in schizophrenia (Bänner and Meyer-Lindenberg, 2017; Heinz et al., 2019), intellectual disability and autism (Carper and Courchesne, 2005; Postema et al., 2019; Weston, 2019).

**DISCUSSION**

We have systematically measured protein lifetime at single-synapse resolution across the brain and lifespan of the mouse using the HaloTag self-labeling system and synaptome mapping technology. We focused on PSD95 because it is a highly abundant protein in the postsynaptic terminal of excitatory synapses that plays a crucial role in assembling glutamate receptors and other proteins into signaling complexes that control synaptic strength, synaptic plasticity and learning (Carlisle et al., 2008; Fernández et al., 2017; Fitzgerald et al., 2015; Frank et al., 2016; Migaud et al., 1998; Nithianantharajah et al., 2013). Between the ages of 3 and 6 months, when the amounts of PSD95 in individual excitatory synapses remain stable (Cizeron et al., 2020), we found that the vast majority of synapses in all brain areas replaced their PSD95 content within a few days or weeks. This indicates that proteostasis is constantly rebuilding the excitatory synapse proteome and does so many times throughout the life of the mouse in most brain synapses.

Brain synapses differed widely in their PSD95 protein lifetimes, and these had distinct spatial distributions. We designated the synapse subtypes with the longest protein lifetimes as LPL synapses and those with the shortest protein lifetime as SPL synapses. It is striking that the neocortex is the brain region most enriched in LPL synapses. Moreover, there is a gradient of enrichment across the layers, with the greatest density of LPL synapses in the superficial layer and the lowest density in the deepest layer. Some LPL synapses in the adult mouse brain showed a potential to retain PSD95 for many months, indicative of a remarkable proteome stability. A progressive lengthening of PSD95 lifetime occurs across the whole brain between 3 weeks and 18 months of age. In the aging mouse brain, there is a general loss of synapse number and it was the LPL synapses that are preferentially retained in old age. In a mouse model of autism and schizophrenia, there was a brainwide increase in PSD95 lifetime, indicating that germline disease-relevant mutations that cause childhood and young-adult disorders impact on synapse protein lifetime.

**Protein composition and lifetime contribute to synapse diversity**

Synapse diversity arises by differences in protein composition (Zhu et al., 2018), and here, we show that it also arises by differences in protein lifetime. To further understand the relationship between these two parameters we combined synaptome mapping using two markers that inform on protein composition (PSD95eGFP and SAP102mKO2) with the PSD95-HaloTag that allows quantification of protein lifetime. This enabled us to assign a protein lifetime measure to 30 of the previously defined synapse subtypes (Zhu et al., 2018). We found that synapses with different protein compositions (type 1, expressing PSD95 only; type 3, expressing PSD95 and SAP102) included both LPL and SPL synapses with a wide range of PSD95 protein lifetimes. Thus, synapse protein composition and lifetime are two orthogonal parameters that define the synapse proteome, adding another layer to the known diversity of brain synapses. The protein lifetime of synapse subtypes was independent of brain region, strongly supporting the conclusion that synapses with different protein lifetimes are building blocks that can be employed in any area of the brain, where they underpin the synaptome architecture of neurons, regions, and, ultimately, the overall brain.

**Synaptome architecture arises from the spatial distribution of synapses with different protein lifetimes**

LPL and SPL subtypes are distributed in all brain regions at differing abundance in a non-random manner. Within individual dendrites, protein lifetime differed between adjacent synapses, was longer in distal dendrites and appeared to be influenced by synaptic inputs at different locations on the dendritic tree. Synapses within hippocampal subregions and layers showed different protein lifetimes, indicating projection- and cell-type-specific regulation of PSD95 lifetime. It will be of interest to ask if synapse subtypes with different protein lifetimes are in spatial proximity to dendritic translational machinery and proteosomes (Biever et al., 2020; Bingol and Schuman, 2006; Ostroff et al., 2002; Steward and Levy, 1982) and whether the influence of inputs on postsynaptic protein lifetime is controlled by protein.
interactions across the synaptic cleft or by activity-dependent mechanisms.

**A lifespan trajectory of synapse protein lifetime**

PSD95 lifetime is shortest in juvenile mice and longest in old mice. This lifespan trajectory of increasing protein lifetime occurs across almost all brain regions. Between 3 and 4 weeks of age, there is an increase in the amount of PSD95 in individual synapses (Cizeron et al., 2020), indicating that PSD95 synthesis must exceed its degradation. Yet at this age, synapses have the shortest PSD95 lifetime and the highest rate of protein degradation. This indicates that in juvenile mice, which are at the point of weaning and beginning an independent life, there is a very high rate of synapse proteome construction and remodeling, which is well suited to the adaptive changes programmed by the transcriptome (Skene et al., 2017) and any superimposed activity-dependent changes induced by environmental experience.

We previously showed that specific synapse subtypes are preferentially retained in old age whereas others are lost: “age-resilient” and “age-sensitive” synapse subtypes, respectively (Cizeron et al., 2020). Here, we report that the age-resilient synapse subtypes correspond to LPL synapses and the age-sensitive subtypes correspond to SPL synapses. Thus, in old age, synapses with rapid proteostasis and adaptive properties are depleted from the brain, whereas those with slow protein turnover and slow adaptive properties are retained.

**Functional implications of synapse protein lifetime diversity**

The adaptive and homeostatic properties of synapses have been of intense interest, particularly in the context of how synaptic proteins are modified in response to neuronal activity and environmental experience. Central among these proteins is PSD95 which assembles glutamate receptors with other proteins into supercomplexes (Fernández et al., 2009; Frank et al., 2016, 2017; Husi et al., 2000). Although these proteins and their modifications are necessary for the initial encoding of memories, the mechanisms controlling the duration of memories and mechanisms of forgetting are less clear. Our discovery of LPL synapses, which could retain copies of PSD95 for many months, challenges Crick’s assumption (Crick, 1984) that synaptic protein lifetime is far shorter than long-term memory duration. Several findings suggest that LPL synapses may be involved with retaining memories for longer durations than SPL synapses. LPL synapses are heavily enriched in the neocortex where long-term memories are stored (Squire and Dede, 2015). Young mice, which have fewer LPL synapses than mature mice, have shorter-lasting memories than adult mice (Akers et al., 2012). The lifetime of PSD95 in CA1, visual, and piriform cortex corresponds well with the duration of spatial, visual, and olfactory memory representations measured using calcium imaging of individual neurons (Driscoll et al., 2017; Rubin et al., 2015; Schoonover et al., 2021) and with the duration of protein-synthesis-independent memories in DG engram cells (Roy et al., 2017). A hallmark of old age is the retention of long-term memories and a reduction in short-term and working memory capacity (Li et al., 2004; Reinert, 1970; Tucker-Drob, 2009), which is paralleled by the preferential retention of LPL but not SPL synapses in old age.

This framework also fits with the anatomical locus of innate behaviors, which are genetically programmed and “hard wired.” Basal brain structures that contain innate behaviors are highly enriched in SPL synapses. This would endow these circuits with rapid proteostatic mechanisms capable of erasing and reversing protein modifications and thereby ensuring stable genetically programmed synaptic functions. In support of the notion that the rapid proteostatic properties of SPL synapses are a mechanism of homeostatic synaptic plasticity, studies in CA1 pyramidal neurons show that synapses closest to the soma, which have the shortest PSD95 lifetime, contribute more to homeostasis than distal synapses (Lee et al., 2013), which have the longest PSD95 lifetime.

**Synaptic protein lifetime and mechanisms of disease**

Our findings have relevance for brain diseases that arise at different ages. Childhood neurodevelopmental disorders, such as autism, are caused by mutations affecting protein synthesis and degradation (Lourois and Osterweil, 2016). In the Dlg2 mutant mouse model of autism and schizophrenia, we have shown that PSD95 lifetime is severely affected in the majority of brain regions. We speculate that the mechanism involves the physical interactions of PSD95 and PSD93 (which is encoded by Dlg2) with NMDA receptors and other proteins in the postsynaptic terminal (Frank et al., 2016). The loss of PSD93 may influence PSD95 interactions with other proteins and its anchoring in the synapse. Many neurodevelopmental disorders are caused by mutations in genes that encode PSD95-interacting proteins (Bayés et al., 2011; Fernández et al., 2009; Fromer et al., 2014; Husi et al., 2000; Kirov et al., 2012; Purcell et al., 2014; Skene et al., 2017), and it will be of interest to establish whether these affect PSD95 protein lifetime.

The longer PSD95 protein lifetime in synapses of old mice could potentially be explained by the progressive reduction in protein clearance that underpins senescence and longevity (Kelmer Sacramento et al., 2020; Koyuncu et al., 2021). This reduction in protein clearance contributes to the accumulation of toxic and misfolded proteins in neurodegenerative disorders (Labbadia and Morimoto, 2015; Yerbury et al., 2016) and is thought to result from decreased proteasomal activity in aging (Carrard et al., 2003; Dasuri et al., 2009; Hayashi and Goto, 1998; Kelmer Sacramento et al., 2020; Saez and Vilchez, 2014; Vilchez et al., 2014; Zeng et al., 2005). Our findings suggest that SPL synapses would be more efficient at removing toxic proteins than LPL synapses and, consequently, that LPL synapses would be more vulnerable to damage. The accumulation of LPL synapses in old age, together with their enhanced vulnerability to toxic proteins, might at least in part explain the proclivity of Alzheimer’s disease to target synapses in old age and impair memory.

**Technical considerations for HaloTag studies of synapse protein lifetimes**

Our pharmacodynamic experiments show that the injected SiR-Halo ligand is rapidly cleared from the body, leaving negligible amounts 24 h after injection. The ligand is therefore only available for labeling the PSD95-HaloTag protein in the brain on the day of injection and does not linger long enough to label protein synthesized on subsequent days. The rapid clearance of ligand
from the circulation raises the possibility of using multiple pulses of ligand labeled with different fluorophores to examine PSD95 dynamics with greater temporal precision, exploiting the capacity of super-resolution methods to explore nanoscale lifetime architecture (Masch et al., 2018). Our mice can also be used to examine endogenous PSD95 lifetime in living primary neuronal cultures (Kratschke, 2018) and could be used to study the dwell time of endogenous PSD95 at synapses and any diffusion, trafficking, or exchange of PSD95 that occurs between synapses. Endogenous PSD95 can be conditionally tagged for cell-type-specific studies of synapse protein composition and turnover (Zhu et al., 2020). Our biochemical, electrophysiological, and anatomical analyses failed to expose an impact of the HaloTag; however, in the absence of a method to measure the lifetime of endogenous PSD95 at single-synapse resolution we cannot exclude the possibility that the tag might have some influence on the lifetime of proteins to which it is attached.

A number of studies have shown that proteins that interact with each other in multiprotein complexes have correlated half-lives (Martin-Perez and Villén, 2017; Mathieson et al., 2018; Price et al., 2010). This implies that PSD95 lifetime could be a “sentinel” for the lifetime of many other postsynaptic proteins that it interacts with. If all proteins in a given dendritic spine had correlated lifetimes this would imply that the whole spine is governed by the same proteostatic mechanisms, such as localized dendritic protein translation and degradation machinery. Our methods could be applied to determine whether pre- and postsynaptic protein lifetimes are correlated.

The protein lifetime synaptome atlas resource
The atlas provides a versatile reference resource for understanding and modeling memory and molecular and cellular anatomy in health and disease. It provides a framework that can be integrated with anatomical, molecular, and phenotypic datasets. Data from the mouse atlas can also be used to generate hypotheses for validation in other species; for example, it will be interesting to ask whether the human neocortex is populated with synapses with longer protein lifetimes than those in the mouse. The value of the foundation dataset, which is already linked to the Lifespan Synaptome Atlas (Cizeron et al., 2020), will increase as it is expanded to encompass lifetime studies of further synaptic proteins, both pre- and postsynaptic and inhibitory as well as excitatory. Understanding the spatiotemporal similarities and differences between these proteins will be highly informative as to underlying proteostasis mechanisms. The atlas will take on significant health discovery value as neurological disease models are incorporated, and common underlying themes may emerge.

STAR METHODS
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**RESOURCE AVAILABILITY**
- **CONTACT FOR REAGENT AND RESOURCE SHARING**
  - Sarah Lempiere, Cathy McLaughlin, Emma Sigfridsson, Rand Dahan, Gabor Varga, Emily Robson, Theresa Wong, Bev Notman, Ian Hawes, Dimitra Koukaroudi, William Mungall, and Fabio De Moliner for technical assistance. Colin Davey for editing. Debbie Maiuezis for artwork. Trevor Robbins, Szu-Han Wang, Tim Bussey for comments on the manuscript. Funding: S.G.N.G.: The European Research Council (ER) under the European Union’s Horizon 2020 Research and Innovation Programme (695668 SYNNOVA). Simons Foundation Autism Research Initiative (529085), and the Wellcome Trust (Technology Development Grant 202932). J.D.: Interdisciplinary Platform Cajal Blue Brain (CSIC, Spain). P.M.-S.: Spanish Ministerio de Ciencia e Innovación (IJC-2016-27658). T.J.O.: National Institute of Mental Health Grant (R01MH060919-15). A.H.: Swiss National Science Foundation (grant 31003A_173125), the Swiss National Centre Competence in Research (NCCR) Synapsy (grant 51NF40-185897), and a gift from a private foundation with public interest through the International Foundation for Research in Paraplegia (chair Alain Rossier). For the purpose of open access, the author has applied a CC-BY public copyright license to any author accepted manuscript version arising from this submission.

**AUTHOR CONTRIBUTIONS**
E.B. developed methods; planned the experiments; collected, imaged, and analyzed brain samples; and performed image and data analysis. Z.Q.
developed software and performed image analyses. M.K. analyzed protein extracts and developed methods in primary cultures. A.Z. imaged Dlg2 brain samples and performed image and data analysis. N.H.K., D.G.F., and E.J.T. constructed the PSD95-HaloTag mice. R.G. built the Protein Lifetime Synaptope Atlas website and performed statistical analysis. B.K. developed software for image and data analysis. S.A.J. and T.J.O. performed electrophysiology. P.M.-S. and J.D. performed neuron filling. E.H. and A.H. performed in vivo neuronal imaging. E.F., L.M.-T., and M.V. provided advice on experimental design and analyzed data. S.G.N.G. conceived the project and wrote the paper with input from all authors.

**DECLARATION OF INTERESTS**

The authors declare no competing interests.

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## KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Mouse monoclonal anti-HaloTag | Promega | Cat# 9211 RRID: AB_2688011 |
| Mouse anti-PSD95 (clone 16/PSD-95) | BD Biosciences | Cat# 610496 RRID: AB_397862 |
| **Bacterial and virus strains** |        |            |
| AAV9.CamKII.Cre.SV40 | Addgene | Cat# 105558 |
| AAV2.EF1a.DIO.eYFP | UNC GTC vector | Cat# 42585 |
| **Chemicals, peptides, and recombinant proteins** | | |
| 4',6 diamidino-2-phenylindole (DAPI) | Sigma-Aldrich | Cat# D9542 |
| Alexa Fluor 488 dye solution | Life Technologies | Cat# A10436 |
| Pluronic F-127 (20% in DMSO) | Thermo Fisher | Cat# P3000MP |
| ProLong Gold Antifade mounting medium | Life Technologies | Cat# P36930 |
| OCT embedding matrix | CellPath | Cat# KMA-0100-00A |
| TMR-Halo | Promega | Cat# G8251 |
| SiR-COOH | Spirochrome | Cat# SC004 |
| Amine (O2) HaloTag | Promega | Cat# P6711 |
| **Deposited data** | | |
| Allen mouse brain reference atlas | Allen Brain Institute | https://mouse.brain-map.org/static/atlas |
| Synaptome data, including data for time points, individual animals and half-life values | This paper | https://doi.org/10.7488/ds/3070 |
| **Experimental models: Cell lines** | | |
| Embryonic stem cell line used for gene targeting: E14Tg2a | | 08021401 |
| **Experimental models: Organisms/strains** | | |
| Mouse: PSD95-HaloTag/HaloTag | This paper | N/A |
| Mouse: PSD95-eGFP/eGFP | Zhu et al., 2018 | N/A |
| Mouse: SAP102mKO2/mKO2 | Zhu et al., 2018 | N/A |
| Mouse: Dlg2-/- | McGee et al., 2001 | N/A |
| **Oligonucleotides** | | |
| Mouse genotyping primers PSD95-HaloTag: exon F, GTCACTAGTCCTTTGTGACCTTG | This paper | N/A |
| Mouse genotyping primers PSD95-HaloTag: 9UTR F, GATACTAGTCAGAGGAGTGTC | This paper | N/A |
| Mouse genotyping primers PSD95-HaloTag: Halogen F, CTGACCTGAAATGGAGATGGAC | This paper | N/A |
| **Recombinant DNA** | | |
| HaloTag coding sequence | Promega | https://www.promega.co.uk/resources/technologies/halotag/ |
| **Software and algorithms** | | |
| MATLAB | MathWorks | https://uk.mathworks.com/products/matlab.html |
| ImageJ | NIH | https://imagej.nih.gov/ij/ |
| Python | Python Software Foundation | https://www.python.org/ |
| Illustrator | Adobe | https://www.adobe.com/products/illustrator.html |
| Imaris 7.6.5 | Bitplane | https://imaris.oxinst.com/ |

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RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Seth Grant (seth.grant@ed.ac.uk).

Materials availability
Newly generated materials are available from the lead contact upon request and, where appropriate, provision of a materials transfer agreement.

Data and code availability
- Data are available at the Protein Lifetime Synaptome Atlas website (Bulovaite et al., 2021a) and Edinburgh DataShare (Bulovaite et al., 2021b).
- Synaptome map data have been deposited in Edinburgh DataShare and are publicly available as of the date of publication. The DOI is listed in the key resources table.
- All original code has been deposited at GitHub and is publicly available as of the date of publication. DOIs are listed in the key resources table.
- Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Gene targeting and mouse generation
Animal procedures were performed in accordance with UK Home Office regulations and approved by Edinburgh University Director of Biological Services. The genetic targeting strategy adapted from Fernández et al. (2009) was used to fuse HaloTag protein to the C-terminus of endogenous PSD95. HaloTag coding sequence (Promega) together with a short linker were inserted into the open reading frame of the mouse Psd95 gene, followed by insertion of a loxP floxed PGK-EM7-neo-pA cassette, using recombination in Escherichia coli. E14Tg2a ES cells (from 129P2 ola) were used for gene targeting. Following identification of positive targeting clones using PCR, E3.5 blastocysts from C57BL/6 mice were injected with ES cells containing the target gene. Male chimeras were crossed with C57BL/6 females to produce first generation heterozygous animals. These mice were crossed with CAG Cre recombinase-expressing mice in order to remove the loxP floxed neo cassette in vivo. Animals were then cross-bred to produce a homozygous PSD95-HaloTag colony.

PSD95-HaloTag mice were genotyped by PCR using the following primers (5’-3’): exon F, GTCACATGCTTTTGACCTTG; 95UTR F, GATACATGCAGAGAGGAGTGTC; and Halogen F, CTGACTGAAGTCGAGATGGAC. Generation and characterization of the Psd95eGFP/eGFP;Sap102mKO2/mKO2 knock-in mouse line was described previously (Zhu et al., 2018). To establish the Psd95HaloTag/eGFP;Sap102mKO2/mKO2 line, double homozygous Psd95KGFPeGFP;Sap102mKO2/mKO2 mice were crossed with Psd95HaloTag/HaloTag mice. Generation and characterization of the Dlg2 mutant mouse line was described previously (McGee et al., 2001). To establish the Dlg2+/–;Psd95HaloTag/+ line, Dlg2–/– mice were crossed with Psd95HaloTag/HaloTag mice.

The mice were group housed and littermates were randomly assigned to experimental groups. The following groups of mice (m, male; f, female) were used for the study of PSD95 lifetime: 3 month old animals, day 0 n=9 (5m, 4f), day 1.5 n=8 (4m, 4f), day 3 and day 14 n=8 (3m, 5f), day 7 n=7 (2m, 5f); 3 week old animals, day 0 n=8 (5m, 3f), day 7 n=8 (6m, 2f); 18 month old animals,
day 0 n=10 (7m, 3f), day 7 n=9 (8m, 1f). For triple colocalization the groups were: day 0 and day 7, n=8 (8f). For studies in Dlg2 mutant animals, 3- to 4-month-old animals were used and the groups were: Dlg2+/–;Psd95HaloTag/+ animals, day 0 n=5 (2m, 3f), day 7 n=6 (2m, 4f); Dlg2+/–;Psd95HaloTag/+ animals, day 0 n=8 (6m, 2f), day 7 n=8 (3m, 5f); Dlg2–/–;Psd95HaloTag/+ animals, day 0 n=7 (3m, 4f), day 7 n=7 (2m, 5f). Sample size was estimated based on Cizeron et al. (2020).

**METHOD DETAILS**

**Western blotting**

PSD95 protein expression and abundance were tested in whole-brain synaptosome extracts from wild-type, Psd95+/–HaloTag and Psd95HaloTag/HaloTag animals. Three extracts per genotype were loaded onto 4–12% gradient Bis-Tris gels, with 20 μl loaded per well. The gels were subjected to SDS-PAGE and western blotting. The membrane was probed with antibodies for PSD95, PSD95-HaloTag and alpha-Tubulin (loading control) and bands detected using the Li-COR Odyssey imaging system.

**SDS-PAGE analysis**

Synaptosome preparations from Psd95+/–HaloTag and wild-type mice as well as cell lysates from HEK293 cells expressing cytosolic HaloTag protein alongside wild-type controls were incubated with 1 μM tetramethyl-rhodamine HaloTag ligand (TMR-Halo) for 1 hour at 4°C. The preparations were then subjected to SDS-PAGE analysis and imaged using the Li-COR Odyssey imaging system.

**Electrophysiological recordings**

Hippocampal slices obtained from the dorsal third of the hippocampus were prepared and maintained in vitro using previously described techniques (Babiec et al., 2017) approved by the Institutional Animal Care and Use Committee at the University of California, Los Angeles, USA. In experiments using extracellular recording techniques, slices were maintained at 30°C in an interface-slice type recording chamber perfused (2-3 ml/min) with an oxygenated (95% O2/5% CO2) artificial cerebrospinal fluid (ACSF) containing 124 mM NaCl, 4 mM KCl, 25 mM NaHCO3, 1 mM NaH2PO4, 2 mM CaCl2, 1.2 mM MgSO4, 10 mM glucose (all Sigma-Aldrich). Field EPSPs (fEPSPs) evoked by Schaffer collateral/commisural fiber stimulation were recorded in stratum radiatum of the CA1 region using low-resistance (5-10 MΩ) glass recording electrodes filled with ACSF (basal stimulation rate = 0.02 Hz). LTP was induced using two, one-second-long trains of 100 Hz stimulation (inter-train interval = 10 seconds). Whole-cell voltage-clamp recordings of evoked excitatory postsynaptic currents (EPSCs) and miniature EPSCs (mEPSCs) were performed using slices maintained in submerged-slice type recording chambers perfused with a modified ACSF containing 2.4 mM KCl, 3.0 mM CaCl2, 2.4 mM MgSO4, 100 mM picrotoxin. In these experiments, recording electrodes were filled with a solution containing 102 mM Cs-glucuronate, 20 mM CsCl, 10 mM K-gluconate, 10 mM TEA-Cl, 5 mM QX-314, 2.5 mM Mg-ATP, 0.2 mM Na-GTP, 20 mM HEPES (pH 7.3, 290 mOsm). Evoked EPSCs (at 0.2 Hz) were recorded at membrane potentials of ±80 mV and ±40 mV and AMPAR-mediated and NMDAR-mediated components of the EPSCs were estimated by the amplitude of EPSCs 5 and 50 milliseconds after EPSC onset, respectively. Spontaneous mEPSCs were recorded at ±80 mV in the presence of 1.0 μM TTX (Alomone Labs). Data collection and initial analysis were performed blind to genotype.

**HaloTag ligand generation**

SiR-Halo coupling and purification were performed as previously described (Lukinavicius et al., 2013). 50 mg (0.159 mmol) SiR-COOH dye (Spirochrome) and 36 mg (0.161 mmol, 1.5 eq) Amine (O2) HaloTag (Promega) building block were dissolved in 950 μl dimethylformamide (DMF). 70 mg (0.517 mmol, 5 eq) N,N-diisopropylethylamine (DIPEA) were added into the solution. 70 mg (0.135 mmol, 1.3 eq) benzotriazol-1-yl-oxytrityloldinophosphonium hexafluorophosphate (PyBOP) dissolved in 230 μl DMF was then added to the SiR-COOH and Amine (O2) HaloTag solution. The solution was stirred at room temperature for 2 hours protected from light. Prior to purification, the compound was phase-separated from DMF. The compound was collected with the organic phase (diethyl ether) while the aqueous phase (brine) was discarded. Two to three tablespoons of dried magnesium sulphate were added to the organic phase solution, mixed well and filtered to remove any aqueous remains. Diethyl ether was evaporated, and the crude product was purified using silica gel column chromatography (2% MeOH in dichloromethane) to give the SiR-Halo ligand (45 mg, 63% yield).

**In vivo application of HaloTag ligands**

SiR-Halo ligand was dissolved in DMSO to a stock concentration of 5 mM. HaloTag ligand solution for injections was prepared as described (Grimm et al., 2017) but at 1.5 mM dye concentration and the total injection volume was adjusted by average weight for the animal group examined. 3 week old mice received 70 μl (21 μl HaloTag stock solution, 7 μl Pluronic F-127, 42 μl saline), 3 month old mice received 200 μl (60 μl HaloTag stock solution, 20 μl Pluronic F-127, 120 μl saline) and 18 month old mice received 300 μl (90 μl HaloTag stock solution, 30 μl Pluronic F-127, 180 μl saline) of HaloTag dye solution. Prior to injection, animals were placed in a heat box for 5-10 minutes to allow the blood vessels of the tail to dilate and become more visible. The mice were placed in a rodent restrainer for the injection. A bolus injection of HaloTag ligand solution was delivered into the lateral tail vein. Following injection, the animals were monitored for any adverse effects twice daily for the length of the experiment. The experimenter was blinded to time points and genotypes at the time of injections.
Tissue collection and section preparation

Mice were fully anesthetized by intraperitoneal injection of 0.10–0.20 ml (according to age) pentobarbital (Euthatal). The thorax was opened and an incision made to the right atrium of the heart. A fine needle (26G, 0.45 x 100 mm) was inserted into the left ventricle and the animal was transcardially perfused with 10–15 ml 1xPBS followed by 10–15 ml fixative (4% paraformaldehyde (PFA)) and left at 4°C for 3–4 hours (according to age). Samples were transferred to 30% sucrose solution and incubated for 48–72 hours at 4°C. Brains were then embedded in OCT (embedding matrix for frozen sections) solution (CellPath) inside a plastic mould (Sigma-Aldrich), and the moulds placed in beakers containing isopentane (Sigma-Aldrich), and the beakers moved to a container containing liquid nitrogen for freezing. Frozen brains were stored at -80°C for up to 4–5 months. Frozen brain samples were cut at 18 μm thickness using a cryostat (NX70 Thermo Fisher) to obtain sagittal sections referring to 12-13/21 bregma level from the Allen Mouse Brain Atlas (sagittal, https://mouse.brain-map.org/experiment/thumbnails/100042147?image_type=atlas). Cut brain sections were placed on Superfrost Plus glass slides (Thermo Fisher). A drop of 1xPBS was placed on a glass slide prior to picking up the brain sections to ensure the brain tissue lay flat. After cutting, brain sections were left to dry in the dark at room temperature overnight and were then stored at -20°C. Frozen brain sections were placed in a dark chamber and incubated at room temperature for 1 hour prior to mounting the coverslips. Sections were washed with 800 μl 1xPBS to remove any remaining OCT on or around the brain tissue and allowed to dry. A 12 μl drop of MOWIOL solution was applied on top of the brain section. A glass coverslip (18 mm diameter, thickness #1.5, VWR) was carefully lowered on top of the sample to avoid any bubbles forming in between the glass slide and the coverslip. The sections were left to dry in the dark at room temperature overnight and then stored at 4°C for up to 1 week. The experimenter was blinded at the stage of tissue sectioning and section preparation for imaging.

Post-fixation labeling with HaloTag ligands

Brain sections were first washed with 800 μl PBS to remove any remaining OCT and left to dry in the dark. Hydrophobic marker pen was used to draw around sections to contain the solutions. 50 μl 10 μM TMR-Halo in 1xPBS was added to each brain section and samples incubated for 1 hour at room temperature in a wet dark chamber. Brain sections were then washed twice for 10 minutes each with 1xTBS (Tris buffered saline) containing 0.2% Triton X-100 detergent to remove any unbound TMR-Halo ligand and once with 1xTBS for 10 minutes.

Pharmacokinetic analysis

Wild-type C57BL/6 mice (n=21, males and females, ~2.5-10 months old) were injected with 200 μl 1.5 mM SiR-Halo solution and anaesthetised using isoflurane at different time points post-injection (1, 5, 15, 30 minutes, 1, 6, 24 hours). Additional n=6 mice did not receive an injection of SiR-Halo but were processed as per below and were used as controls. The thorax was opened and a ~0.5 ml blood sample was drawn from the left ventricle. The animals were then perfused with 12 ml 1xPBS and decapitated. The brains were dissected and snap frozen in liquid nitrogen. The collected blood samples were left to clot for 20 minutes at room temperature and a further 5 minutes on ice and then centrifuged at 10,000 rpm for 10 minutes. The supernatant was collected, and the resulting serum samples as well as brain samples were stored at -80°C until further analysis. Each brain sample was homogenized in 7 ml buffer solution containing 1% sodium deoxycholate, 50 mM Tris pH 9 and 100 mM NaCl. Following ~20 strokes with a homogenizer, the solution was left on ice for 30 minutes after which another 10 strokes were performed and the solution was incubated for another 30 minutes. The samples were then centrifuged at 50,000 g (20,000 rpm) for 30 minutes at 4°C. The supernatant was collected, aliquoted and stored at -80°C.

The fluorescence of SiR-Halo in blood serum and brain homogenate samples was detected using BioTek CYTATION 3 spectrophotometer (600 nm excitation wavelength). To increase the fluorescence readout (Lukinavičius et al, 2013), blood serum samples were diluted in a ratio of 1:2 in 10% SDS solution (in 1xPBS). Brain homogenates were diluted in a ratio of 1:1 in 10% SDS solution (in 1xPBS). Three mice per time point and a triple replicate per sample were performed.

CA1 pyramidal neuron filling

Mice (n=14) were perfused intracardially with 1xPBS followed by 4% PFA and, after removal, brains were post-fixed in PFA for 24 hours. After washing in PBS, brains were cut in coronal sections (200 μm thick) with a vibratome, and to identify cell nuclei the sections were prelabeled with 4',6-diamidino-2-phenylindole (DAPI) (Sigma). Then, CA1 pyramidal neurons were individually injected with Alexa Fluor 488 dye solution (Life Technologies) by continuous current until the dendrites were filled completely as previously described (Benavides-Piccione et al., 2013). After injections, sections were mounted in ProLong Gold Antifade mounting medium (Life Technologies). Imaging was performed with a Zeiss LSM 710 confocal microscope. The fluorescence of DAPI, Alexa 488 and 633 (SiR-Halo-positve puncta) was recorded through separate channels. Stacks of images were acquired at high magnification (63x oil immersion; pixel size 0.057 x 0.057 μm; z-step 0.14 μm) and no pixels were saturated within the spines. For intracellular injection experiments, the animals were randomly assigned into groups for different time points for sacrifice.

Spinning disk confocal microscopy

Imaging was performed using an Andor Revolution XDI spinning disk microscope equipped with CSU-X1 (pinhole size 50 μm) and 2x post-magnification lens. Images of 512 x 512 pixels in size and 16-bit depth were obtained using Andor iXon Ultra back-illuminated EMCCD camera and Olympus UPlanSAPO 100x oil-immersion lens (NA 1.4). To cover the whole area of the sagittal brain section,
multi-tile single-plane image acquisition was arranged with no overlap between adjacent tiles. Focus in z-plane was achieved by selecting and recording desired z-positions for four points on the brain section, which were then used to calculate the intermediate z-positions for in-between image tiles. The optical resolution was estimated based on reconstructed 3D point spread functions (PSFs) and full width at half maximum (FWHM) measurements on images of near-IR FluoSpheres (diameter: 40 nm, excitation/emission wavelength: 660/680 nm; Invitrogen). 100 full width at half maximum (FWHM) measurements on images of near-IR FluoSpheres (diameter: 40 nm, excitation/emission wavelength: 660/680 nm; Invitrogen). 100 µl FluoSpheres solution (1:1000 in 1xPBS) was placed on a glass slide with a brain section from a wild-type mouse and incubated for 30 min. After a wash with 1xPBS, the slides were mounted with #1.5 thickness coverslips using MOWIOL as a mounting medium. Z-stack images were acquired with 150 nm step size and 15-20 image planes were obtained for each bead sample. 3D PSFs from 392 beads were reconstructed to obtain average FWHM values.

Imaging parameters
High-magnification (×100) images at single-synapse resolution covering the whole brain section were obtained using 250 EM gain, 2-frame averaging and 5000 millisecond acquisition speed. SiR-Halo was excited at 640 nm, TMR-Halo at 561 nm, eGFP at 488 nm and mKO2 at 561 nm. Emitted light was filtered with a QUAD filter (BP 440/40, BP 521/21, BP 607/34 and BP 700/45).

Long-term imaging in vivo
5-month-old male (n=3) and female (n=1) homozygous PSD95-HaloTag mice were used in accordance with the guidelines of the Federal Food Safety and Veterinary Office of Switzerland and in agreement with the veterinary office of the Canton of Geneva (license number GE12219A). To generate sparse expression of YFP in the supragranular layers of cortex, a mix of AAV9.CamKII.Cre,SV40 (2.4x10^13 GC/ml; Addgene #105558) and AAV2.Ef1a.DIO.eYFP (4.6x10^12 GC/ml; UNC GTC vector #42585) (100 nl in a ratio 1:1500) was injected in the primary somatosensory cortex. Cranial windows were implanted as described previously (Holmata et al., 2009) using a circular (3 mm diameter) coverglass pressed down onto the intact dura and glued to the skull using dental acrylic cement. 2-3 weeks after window implantation, SiR-Halo ligand was injected in the tail vein, as described above. Images were acquired in head-fixed mice under anesthesia (0.1 mg/kg Medetomidin and 5 mg/kg Midazolam, i.p.) using a custom-built, two-photon laser-scanning microscope (2PLSM) (Holmata et al., 2009) controlled by custom software written in MATLAB (Scanimage, Vidrio Technologies) (Pologruto et al., 2003). A tunable Ti:sapphire laser (Chameleon ultra II, Coherent) was used as a light source, tuned to λ=840 nm for simultaneous excitation of YFP and SiR-Halo ligand, or λ=910 nm for YFP alone. Excitation power was kept constant over time. Fluorescent images were collected using a 20x, 0.95 numerical aperture water-immersion objective (Olympus) and GaAsP photomultiplier tubes (10770PB-40, Hamamatsu). Emitted light was spectrally separated using a 565 nm dichroic mirror (Chroma) and two band-pass emission filters (ET510/50 nm, Chroma; 675/67 nm, Semrock). Field-of-views (FOVs) with YFP-labeled dendrites were first imaged at 6 hours post-injection, and then re-imaged after 1, 2, 3, 7, 10 and 14 days. Additional FOVs were imaged only once over the course of the experiment to measure population fluorescence decay similarly to the in situ experiments (Figure S9C). For each FOV, two separate image stacks (typically 10-30 planes, separated by 1 μm) were acquired at 2 milliseconds/line (image size, 1024 x 1024 pixels; pixel size, 0.06 x 0.06 μm), one at excitation λ=840 nm and one at λ=910 nm. The dim YFP signal that was collected at λ=910 nm served to align the dendrites in both image stacks.

QUANTIFICATION AND STATISTICAL ANALYSIS

Analysis of electrophysiological data
Acquisition and analysis of electrophysiological data were performed using pCLAMP software (Molecular Devices). Results are reported as mean ± SEM.

Detection of synaptic puncta
Synaptic puncta detection from fluorescence images was performed using a machine learning-based ensemble method developed in-house by Dr Zhen Qiu (Cizeron et al., 2020; Zhu et al., 2018). The classifiers/detectors of PSD95-eGFP and SAP102-mKO2 were trained in a previous study (Zhu et al., 2018). The training of the image detector for PSD95-HaloTag was required. We first collected a training set of 249 images (10.8 m) by randomly sampling across 12 main brain regions using bootstrapping. All puncta in the training set were manually located by three independent individuals of varying scientific expertise. Manual annotation of each punctum was then weighted for each individual. The ground truth was generated by weighted averaging over all individuals; an average weight greater than 0.7 was considered as true puncta. The ensemble learning was applied for training and a K-folder method was applied to validate and test the trained detector.

Measurement of synaptic parameters
Upon detection and localization, synaptic puncta were segmented by applying an intensity threshold. The threshold for each detected punctum was set adaptively at 10% of the height of its fluorescence intensity profile. A set of parameters were then quantified for each segmented punctum, including mean pixel intensity, size, skewness, kurtosis, circularity and aspect ratio (Zhu et al., 2018). Additionally, quantification of puncta density per unit area was performed. For calculations of PSD95 lifetime, primarily the measurements of puncta density and total fluorescence intensity content (punctum size x punctum mean intensity) were used.
Comparison of SiR-Halo and eGFP puncta populations

Owing to different imaging and puncta detection parameters, fewer SiR-Halo puncta were detected compared with eGFP (Figure S3E). To assess whether SiR-Halo labeled a representative sample of the diverse population of PSD95-positive synapses, total fluorescence intensity distributions for SiR-Halo and eGFP from injected Psd95−/−;eGFP/HaloTag mice were compared. A scaling factor of 0.65 was applied to eGFP puncta distribution to overlay the outlines of the two frequency distributions. The outlines closely overlapped indicating that eGFP and SiR-Halo labeled comparable populations of synaptic puncta, including comparable proportions of puncta of different intensities. These results demonstrate that injection of a fluorescent ligand into PSD95-HaloTag mice efficiently and quantitatively labels PSD95 across all regions of the brain.

SiR-Halo-positive puncta 3D reconstruction

The z-stacks of SiR-Halo and dye-filled neuron images were analyzed with 3D image processing software Imaris 7.6.5 (Bitplane). SiR-Halo-positive puncta were manually reconstructed along apical dendrites (stratum radiatum). For detailed information regarding 3D reconstruction of SiR-Halo-positive puncta, see Figure S9A.

Analysis of long-term imaging

The images from λ=840 nm and λ=910 nm acquisitions were overlaid using the Fiji pairwise stitching algorithm (BigStitcher) (Hörler et al., 2019). The images at different time points were aligned using the ‘Correct_3D_Drift’ plugin in Fiji (Parslow et al., 2014). To correct for slight variations in excitation efficiency and image quality at different time points, the images were normalized using the YFP fluorescence signal from the dendrites, under the assumption that YFP levels did not substantially change over time, and background subtracted using the blood vessel interior as the baseline signal. For single SiR-Halo punctum decay analysis, the total fluorescence of each punctum at every time point was measured from the plane that yielded the brightest signal. For spine SiR-Halo puncta, the spines were selected in the YFP channel. YFP and SiR-Halo fluorescence intensity (from 910 nm and 840 nm excitation, respectively) was integrated in the spine head. To correct for bleaching due to repeated imaging of the puncta, on the first day of imaging a series of seven images were taken of a control FOV in rapid succession (within 5 minutes). For each point in the series the level of bleaching was estimated (~3%) and applied as a correction factor to the corresponding time point in the long-term imaging series.

Colocalization of synaptic puncta

Object-based colocalization method was employed to assess double (between SiR-Halo and eGFP) and triple (between SiR-Halo, eGFP and mKO2) colocalization of markers of synaptic puncta. The double colocalization between two channels (SiR-Halo and eGFP) was obtained by minimizing the summation of the total centroid distance between all eGFP and SiR-Halo using linear programming (Zhu et al., 2018). The triple localization is computationally much more complex and an NP-hard problem, thus it cannot be resolved efficiently using linear programming. New multiple-channel localization algorithms were developed by adapting the multiple hypothesis tracking algorithm (Chenouard et al., 2013) to the particle tracking problem: the image channel of different markers were considered as temporal image frames in tracking, but more image matching between different markers was added because there was no temporal sequence in multilocalization. A distance threshold of 400 nm was applied to identify markers that were colocaled or triple localized. 400 nm is within the range of the previously measured diameters of postsynaptic density as observed by electron microscopy (Harris et al., 1992).

Classification of synaptic puncta

Using classifiers trained and built in a previous study (Cizeron et al., 2020), Psd95−/−;eGFP/+;Sap102mKO2/y synapses were classified into 37 subtypes based on the parameters measured in the previous steps. A small percentage of puncta (~0.002%) that did not get classified into any subtype group were denoted ‘other subtype’. The classification results were then combined with the triple localization result of PSD95-HaloTag to quantify the lifetime of different synapse subtypes over all brain regions/subregions at time points post SiR-Halo injection.

Segmentation of brain regions and subregions

An overview montage of each imaged sagittal brain section was stitched from ~45,000-50,000 image tiles using a custom MATLAB script and downsized by a factor of 16. Quantification of PSD95-Halo labeling was performed on anatomical brain regions as defined by the Allen Mouse Brain Atlas. A combination of manual and semi-automated delineation methods was employed for defining 110 anatomical regions in each sagittal brain image. Manual delineations of subcortical brain regions (subregions of the olfactory areas, cortical subplate, striatum, pallidum, hypothalamus, thalamus, midbrain, cerebellum, pons, medulla) were performed using a polygon selection tool in Fiji/ImageJ (Schindelin et al., 2012; Schneider et al., 2012). Semi-automated delineations of isocortex and hippocampal formation were performed with in-house custom-built software. The program provides a user interface for a freeform deformation of a surface on which we map a specially prepared image that contains a multitude of delineated regions (Sederberg and Parry, 1986). The specially prepared images were, in most cases, delineation templates obtained from the Allen Mouse Brain Atlas. The delineation templates mapped on a digital surface are subdivided into a uniform grid so that each cell contains a subdivision of the delineation template. The software allows the user to manipulate the grid points, which results in the deformation of the grid cells and in turn leads to the deformation of the delineation template (Catmull, 1974). The software superimposes the delineation
template on top of the brain image being delineated. The user can deform the delineation template to ‘fit’ the underlying montage image of the brain section. The software has a function allowing the user to paint over any damaged areas on the tissue to exclude them from the finalized delineation. The final delineation is saved as a set of .roi files representing individual brain regions, which are used in the later steps of the mapping pipeline.

**Decay rate estimation**

PSD95 decay rate was estimated by two alternative methods: (a) puncta count-based estimation and (b) total punctum intensity-based estimation. Method (a) relies purely on the presence or absence of fluorescent synaptic puncta, whereas method (b) considers the fluorescence content within each punctum. For both methods, the mean at day 0 was considered a reference point to which all the subsequent values were normalized. Single-phase exponential decay function was fitted to fractions of puncta/synapse fluorescence remaining:

$$N(t) = N_0 e^{-\lambda t}$$

where \(N(t)\) denotes the puncta density/fluorescence intensity at time \(t\), \(N_0 = N(0)\) is quantity at \(t = 0\) and \(\lambda\) is the decay rate constant.

**Puncta count-based estimation**

The detected number of fluorescent puncta per 100 \(\mu m^2\) (puncta density) was estimated for each ROI from every brain examined using the SYNMAP pipeline. Normalized puncta densities (y-axis) for each animal were plotted against time (x-axis) and an exponential decay function was fitted to estimate half-life and \(\pm\) 95% confidence interval.

**Total puncta intensity-based estimation**

For each detected punctum within an ROI, mean fluorescence intensity was multiplied by punctum size to obtain the total fluorescence content for each punctum. The resulting fluorescence intensities were summed to obtain the total synaptic fluorescence content within a given brain region. Normalized fluorescence intensities (y-axis) for each animal were plotted against time (x-axis) and an exponential decay function was fitted to estimate half-life and \(\pm\) 95% confidence interval.

**PSD95 half-life calculations**

PSD95 half-life was defined as the time required for the PSD95-Halo puncta density/fluorescence intensity to fall to half its initial value. Half-life was calculated based on the decay constant obtained from exponential curve fitting via the following formula:

$$t_{1/2} = \frac{\ln(2)}{\lambda}$$

**PSD95 fraction remaining calculation**

In order to compare the decay of puncta number/fluorescence intensity between two time points, for each ROI we estimated the fraction of puncta/fluorescence intensity remaining at day 7 compared with day 0. The mean values for fraction remaining were calculated using:

$$f = \frac{A}{B}$$

where \(A\) represents the mean of day 7 measurements and \(B\) is the mean of day 0 measurements. The error in the measurement of fraction remaining was estimated by following the statistical rules for propagation of uncertainty. The formula used for estimating the standard deviation of a ratio of means was:

$$\sigma_f = \left| f \right| \sqrt{\left(\frac{\sigma_A}{A}\right)^2 + \left(\frac{\sigma_B}{B}\right)^2}$$

where \(\sigma_f\) is a standard deviation of the function \(f\) (ratio of mean (day 7)/mean (day 0)), \(\sigma_A\) is the standard deviation of \(A\) and \(\sigma_B\) is the standard deviation of \(B\).

**Statistical analysis**

Covariance of pairs of synaptic parameters (density, half-life) was assessed using Pearson’s correlation function in MATLAB (Figures S3E and S6C). Correlation coefficient (R) values are presented alongside P-values that determine the statistical significance (P < 0.05) of the correlation between the two variables.

**Bayesian analysis**

Changes in subtype densities between 3 month old and 18 month old mice were established in a previous study (Figure S13 in Cizeron et al, 2020) and the heatmaps (Figures 5E and S11C) were then permuted column-wisely based on the ranking order of the subtype lifetime in Figure 4A. The changes in subtype densities (Figures 5E and S11C) were tested using Bayesian estimation (Kruschke, 2013). Bayesian testing was also used to compare PSD95 lifetime between the Dlg2 mouse mutant and control (Figure S12). The input data were modelled assuming a t-distribution, and a Markov chain Monte Carlo algorithm was then performed to estimate the posterior distribution of the changes. P-values were then calculated for the distribution to complete the test. Each data point in each subregion was modelled and tested separately. The results were finally corrected over all subregions using the Benjamini-Hochberg procedure.
Cohen’s d effect size
The difference between two groups was measured by calculating a Cohen’s d effect size (Figures 5E, S11C, and S12), which is based on the difference between two means divided by their pooled standard deviation:

\[ d = \frac{x_1 - x_2}{s} \]

The pooled standard deviation was calculated as follows:

\[ s = \sqrt{\frac{(n_1 - 1)s_1^2 + (n_2 - 1)s_2^2}{n_1 + n_2}} \]

Self-organizing feature maps
Self-organizing feature maps (SOMs) (Kohonen, 1982) were constructed based on the subtype density of 110 subregions. SOMs preserve the local neighborhood so that subregions which are similar (neighbors) in the 37-dimensional subtype space become neighbors in the 2-dimensional output space. Network nodes were connected on a 2-dimensional rectangular grid of size 35 x 35 nodes. Input consisted of the 37 subtype densities of a subregion, where each subtype was normalized to [0, 1]. Number of training iterations was set to 45 and starting neighborhood size was set to 15 and reduced by a factor 0.9 for each iteration and weights were initialized to [0, 1]. After training, the strongest responding node was identified for each subtype and the position of this node was used for positioning the name of the subregion (Figure S10C, panel i) (half-height of left-hand side of first letter of subregion name) as well as the position of the disc (disc center) representing the subregion (Figures 4D and S10C, panel ii). Radius of discs was proportional to the intensity half-life \( t_{1/2}^{\text{PSD95 intensity}} \) of the subregion represented by the disc. Colors of subregions used for presentations were the same as in Figure 2A.

Similarity matrices
Each row/column in the matrix represents one delineated brain subregion (Figure 5D). Elements in the matrix are the synaptome similarities between two subregions quantified by differences in the PSD95 lifetime. A Gaussian kernel function was applied to convert the differences into similarities (Zhu et al., 2018).

Protein lifetime synaptome atlas website
The Protein Lifetime Synaptome Atlas website (Bulovaite et al., 2021a) provides users with detailed information on PSD95 lifetime in regions and subregions of the mouse brain across the lifespan. The main panel shows the heatmap of protein half-life for the selected punctum parameter (density, intensity) and age or age comparison. The brain region navigator provides a collapsible tree structure that allows the user to pinpoint any subregion, its half-life values, and linked punctum parameter and synapse type/subtype prevalence data from the Lifespan Synaptome Atlas (Cizeron et al., 2020). Half-life data for further synaptic proteins will be added as they become available.

ADDITIONAL RESOURCES
All brain maps generated for the study are freely available at (Bulovaite et al., 2021a).
Supplemental information

A brain atlas of synapse protein lifetime
across the mouse lifespan

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Fig. S1. Related to Fig. 1. Generation and characterization of the PSD95-HaloTag knock-in mouse. (A) PSD95-HaloTag protein domain structure. PSD95 contains three PDZ domains followed by SRC homology 3 (SH3) and guanylate kinase (GK) domains. The HaloTag is fused at the C-terminus. (B) Targeting strategy used for the genomic Psd95 (Dlg4) locus. The Dlg4 allele was targeted with the HaloTag sequence inserted before the stop codon (asterisk). The neomycin resistance cassette (neo) was deleted between loxP sites (triangles) by crossing the PSD95-HaloTag mouse with a transgenic Cre recombinase-expressing mouse. (C) Western blot analysis on crude synaptosome preparations from wild-type (WT), Psd95\textsuperscript{HaloTag/+} and Psd95\textsuperscript{HaloTag/HaloTag} mice using antibody against PSD95. PSD95 and PSD95-HaloTag fusion protein are indicated. (D) The same set of synaptosome preparations as in (C) were probed with anti-HaloTag antibody. Only PSD95-HaloTag protein is detected. (E,F) PSD95-HaloTag is assembled into 1.5 MDa postsynaptic supercomplexes. Complexes were separated using Blue native PAGE of the synaptosome preparations described in (C,D) and immunoblotted with antibodies to PSD95 (E) and HaloTag (F). (G) SDS-PAGE analysis of crude synaptosome preparations from PSD95-HaloTag mice and cell lysate from HEK cells expressing HaloTag-domain both labeled with TMR-Halo ligand. (H) Fiber volley/field excitatory postsynaptic potential (fEPSP) input/output curves generated by eliciting fEPSPs with intensities of presynaptic stimulation corresponding to 100, 75, 50, and 25% of max. (I) Paired-pulse facilitation is normal in PSD95-HaloTag mutants. Results in (H) and (I) from 5 WT (20 slices), 5 heterozygous (21 slices), and 4 homozygous mutant (18 slices) mice. (J) Example mEPSCs (five superimposed 1-second-long recordings for each) showing no difference between WT and mutant. (K,L) Cumulative probability plots of mEPSC amplitudes (K) and inter-event intervals (L). Psd95\textsuperscript{+/-} (WT) n=4 (11 cells), heterozygous (Psd95\textsuperscript{+/-}/HaloTag) n=3 (11 cells), homozygous (Psd95\textsuperscript{HaloTag/HaloTag}) mutant n=4 (15 cells) mice. One-way ANOVA, F(2,8) = 0.049, P = 0.952 for mEPSC amplitude, and F(2,8) = 0.836, P = 0.921 for mEPSC frequency. (M) Example excitatory postsynaptic currents (EPSCs) recorded at membrane potential of -80 and +40 mV illustrate similarity of WT and homozygous mutants. (N) Ratio of NMDAR- to AMPAR-mediated currents recorded at V\textsubscript{m} = +40 mV. No difference was detected between WT and mutants (one-way ANOVA, F(2,11) = 0.345, P = 0.717). (O) Weighted decay time constants for currents elicited at V\textsubscript{m} = +40 mV. No difference in decay times was detected (F(2,11) = 3.602, P = 0.071). In (N) and (O), WT n=5 (19 cells), heterozygous n=3 (12 cells), homozygous mutant n=4 (16 cells) mice. (P) LTP induced by two 1-second-long trains of 100 Hz stimulation (delivered at time = 0, inter-train interval = 10 seconds). Sixty minutes post-100 Hz stimulation, fEPSPs were potentiated to 191 ± 4.5% of baseline in WT slices (n=10 slices from 5 mice), 179 ± 1.3% of baseline in heterozygote slices (n=10 slices from 5 mice), and 186 ± 8.2% of baseline in homozygous mutant slices (n=9 slices from 4 mice). One-way ANOVA, F(2,11) = 1.706, P = 0.226. Dashed line represents baseline synaptic strength. (H,I,K,L,N,O) report mean ± SEM.
Fig. S2. Related to Fig. 1. Optical resolution as estimated by FWHM. Frequency histograms of FWHM estimates in x-y (top) and z (bottom) from 3D PSFs of near-infrared fluorescent beads on brain tissue section from a wild-type mouse. The average FWHM for x-y is 332 nm and for z is 621 nm.
Fig. S3. Related to Fig. 1. Specific, saturating and reliable labeling with SiR-Halo ligands. (A) Representative images of SiR-Halo-labeled PSD95-HaloTag expression in the whole brain, isocortex (CTX), hippocampal formation (HPF), olfactory areas (OLF), striatum (STR), thalamus (TH) and cerebellum (CB) at the indicated days post-injection of SiR-Halo. Scale bars: 2 mm (whole brain), 500 µm (CTX, HPF, OLF), 200 µm (STR, TH, CB). (B) Fluorescence images of sagittal brain sections from a SiR-Halo-injected PSD95-HaloTag mouse (top), an uninjected PSD95-HaloTag mouse (middle), and SiR-Halo-injected wild-type mouse (bottom) reveal highly specific labeling with SiR-Halo ligand. Scale bars: 1 mm. (C) Saturation of labeling by injection of SiR-Halo. (i) HPF section from a PSD95-HaloTag mouse injected with 300 nmol SiR-Halo (6 hours post-injection). (ii) Same section as in (i) stained with TMR-Halo after fixation. The absence of labeling indicates that SiR-Halo has occupied all available Halo ligand binding sites. (iii) HPF section from a PSD95-HaloTag mouse that was not injected with SiR-Halo ligand. (iv) Same section as in (iii) stained with TMR-Halo after fixation. Note the strong labeling of TMR-Halo. Scale bars: 1 mm. (D, E) PSD95 fluorescence labeling in Psd95HaloTag/eGFP mice. (D) Representative fluorescence images of a sagittal section showing PSD95-HaloTag and PSD95eGFP expression (fluorescence) in SiR-Halo-injected Psd95HaloTag/eGFP compound heterozygous mouse. The whole-brain images are of reduced resolution (downsized by a factor of 16) for more accessible visualization. The fluorescence signal observed in the images is composed of synaptic puncta and background signal from non-synaptic sources. The background signal for different fluorophores differs between brain regions and leads to differences in the overall fluorescence observed in cerebellum and olfactory areas for SiR and eGFP channels. Scale bars: 1 mm. (E) Correlation of fluorescent puncta densities detected by PSD95eGFP (x-axis) and PSD95-HaloTag (y-axis) across 110 brain subregions (± SD). Each dot represents a brain subregion, and the color corresponds to the overarching brain area. R = 0.976 and P < 0.0001, Pearson’s correlation test. n=8 animals.
Fig. S4. Related to Fig. 1. Pharmacokinetic analysis of injected SiR-Halo in mouse blood serum and brain homogenate. (A) Calibration curves of detected mean fluorescence intensity (± SD) from different concentrations of SiR-Halo in 1xPBS, blood serum or brain homogenate from wild-type mice. Magnified graph of lower SiR-Halo concentrations (right) reveals a detection threshold at ~10^{-2} µM. N=3 biological replicates. (B) The concentration of SiR-Halo (mean ± 95% confidence interval) in blood serum (top) and brain homogenate (bottom) at 1 min, 5 min, 15 min, 30 min, 1 h, 6 h and 24 h following tail-vein injection. The concentration values were derived from interpolating the calibration curves in (A). n=3 wild-type C57BL/6 mice for each time point and 3 technical replicates per sample. (C) The mean percentage of injected SiR-Halo remaining in blood serum and brain homogenate in relation to t=1 min.
Fig. S5. Related to Fig. 1. Fluorescent puncta decay in different brain regions and subregions. High-magnification example images of SiR-Halo puncta decay across 2 weeks in regions and layers of the isocortex (somatosensory and motor cortex), subregions of the hippocampal formation, olfactory bulb, cerebellum, midbrain, thalamus, striatum and pallidum. Brain subregions are listed in Table S1. Scale bars: 5 µm.
Fig. S6. Related to Fig. 2. Regional and subregional diversity in PSD95 half-life in 3M mouse brain. (A) PSD95 puncta lifetime ($^{\text{PSD95\ density}}_{t_{1/2}}$) across 110 subregions of the 3M mouse brain (bottom). The main brain regions from which these subregions derive are illustrated at the top. (B) Synaptic PSD95 protein lifetime ($^{\text{PSD95\ intensity}}_{t_{1/2}}$) mapped for 110 mouse brain regions. For (A) and (B), n=7-9 $^{\text{Psd95\ HaloTag/HaloTag}}$ mice per time point. See Protein Lifetime Synaptome Atlas for $^{\text{PSD95\ density}}_{t_{1/2}}$ values in each subregion (Bulovaite et al., 2021a). (C) Correlation between density-based and intensity-based half-life estimates. Density-based half-life (x-axis) was plotted against intensity-based half-life (y-axis) for all brain subregions examined. Each dot represents a subregion, color-coded according to the main brain region that they belong to. Half-life is in days. $R = 0.9088$ and $P < 0.0001$, Pearson’s correlation test.
Fig. S7. Related to Fig. 2. Example fitted exponential decay curves and puncta fluorescence intensity distributions for brain regions and subregions. (A) Brain regions with characteristically long and short PSD95 punctum half-lives. Example data-fitted single phase exponential decay curves for regions with especially long-lived (top) and short-lived (bottom) PSD95 puncta (mean ± 95% confidence interval). Dots represent data from individual animals. Data from days 1.5, 3, 7 and 14 are normalized to the mean of day 0 and the fraction of puncta remaining is plotted on the y-axis. Dashed line indicates half-life. (B,C) SiR-Halo labeled puncta density (B) and intensity (C) decay curves for 12 main brain areas. (D) Total puncta intensity (punctum mean intensity x punctum size) frequency distributions at 5 time points post-injection. The distribution graphs display mean (blue line) with ± SD (shaded blue area). The frequencies are normalised to the maximum frequency within a brain area. (E,F) SiR-Halo labeled puncta density decay curves (E) and total puncta intensity frequency distributions (F) for 5 layers of the motor cortex. (G,H) SiR-Halo labeled puncta density decay curves (G) and total puncta intensity frequency distributions (H) for stratum radiatum of hippocampal CA1, CA2 and CA3. n= 7-9 Psd95HaloTag/HaloTag mice per time point.
**Fig. S8.** Related to Fig. 2. Puncta size and density gradient in hippocampal CA1. The average size (top) and density (bottom) of SiR-Halo puncta (± SD) at day 0 post-injection across the apical and basal dendrites of the CA1 pyramidal neurons. n=3 *Psd95HaloTag*/HaloTag mice. Oriens, stratum oriens; Soma, stratum pyramidale; Radiatum, stratum radiatum; Lacunosum, stratum lacunosum; Moleculare, stratum moleculare.
Fig. S9. Related to Fig. 3. PSD95 turnover inside dendritic spines. (A) Intracellular injections and SiR-Halo-positive puncta 3D reconstruction. (i-iii) Confocal microscopy images showing (i) SiR-Halo signal (red) and (ii) a CA1 pyramidal neuron intracellularly injected with Alexa 488 (green). (iii) Same neuron as in (ii) with overexposed green channel to show the total extension of the apical dendritic shaft. White box indicates the area of the analysis. (iv-ix) Representative confocal microscopy images showing the 3D reconstruction of an apical dendrite segment (iv). A solid surface was created to match the apical dendritic shaft (v, green). The red signal inside of the solid surface is selected (vi) and different intensity threshold surfaces that match the SiR-Halo-positive puncta were created (vii, red). Then, the solid surfaces that exactly matched the SiR-Halo-positive puncta were selected for each image stack (viii). In (ix), only the solid surfaces for the SiR-Halo-positive puncta are shown. Sr, stratum radiatum; sp, stratum pyramidale; so, stratum oriens. Scale bar: 20 µm (i-iii), 5 µm (iv-ix). (B) Representative high-magnification images of SiR-Halo and TMR-Halo fluorescence labeling and colocalization at day 3 post-injection of SiR-Halo. Examples presented cover subregions of isocortex, hippocampal formation, olfactory bulb, striatum, pallidum, thalamus, midbrain and cerebellum. Brain subregions are listed in Table S1. Scale bars: 5 µm. (C) In vivo live imaging of dendritic spine and PSD95 turnover. (i) Violin plots of SiR ligand intensity (arbitrary units) in 11 μm x 11 μm regions of interest (ROIs) at 0, 1, 2, 3, 7, 10 and 14 days post injection. Each ROI is imaged only once over the entire time course. (ii) Violin plots of SiR-ligand puncta intensities, normalized to the first imaging time point (day 0). Each punctum is tracked over time. Data correspond to Figure 3F. (iii) Relative SiR ligand puncta intensities versus relative parent dendritic spine YFP intensities at the first imaging time point (day 0). Each data point represents the intensities relative to the mean puncta and spine intensity in its parent field of view. Line, linear regression, indicating a positive correlation between spine intensity (size) and SiR puncta intensity. (iv) SiR ligand puncta fluorescence intensity at day 3 post injection as a fraction of their intensity at day 0, versus the relative parent dendritic spine YFP intensity. Line, linear regression, indicating a very weak correlation between the spine intensity (size) and PSD95 retention. (v) SiR ligand puncta fluorescence intensity at day 3 post injection as a fraction of their intensity at day 0, versus the relative SiR ligand puncta intensities at day 0. Line, linear regression, indicating a very weak negative correlation between the initial PSD95 content and its retention. (vi) As (v), but for the day 7 time point. No significant correlation.
Fig. S10. Related to Fig. 4. PSD95 lifetime in synapse subtypes. (A) Correlation of PSD95 puncta half-life in brain subregions estimated using 2 and 5 time points. For both puncta density (top) and puncta intensity (bottom), 5-time-point half-life estimate was derived from measurements taken at days 0, 1.5, 3, 7 and 14, whereas 2-time-point estimate was derived from measurements taken at days 0 and 7. Each dot represents a brain subregion, and the color corresponds to the overarching brain area. Dashed line represents x=y. R and P values are from the Pearson’s correlation test. (B) Synapse subtypes ranked according to their PSD95 protein lifetime in each of the 12 main brain regions. Bar height indicates the percentage of SiR-Halo-positive subtype density remaining at day 7 compared with day 0. Of the 37 synapse subtypes originally defined (Zhu et al., 2018), the 30 studied here are color-coded by their type 1 (PSD95 only) or type 3 (PSD95+SAP102) molecular composition. n=8 Psd95HaloTag/eGFP;Sap102mKO2/y mice per time point. (C) Self-organizing feature map (SOM) clustering based on subtype densities. Each region is described by its 37 densities of the subtypes and a resulting 2-dimensional SOM clustering is shown. (i) Subregion position is indicated by the lower left corner of the first letter in the subregion name. Subregion names are color-coded according to overarching brain area (see key in Figure 2A, S6A). Axes indicate node index number in the 35x35 SOM node network. (ii) Same network as in (i) but each subregion is represented by a disc, the center of which corresponds to the winning SOM node position and the radius codes for the half-life (PSD95 intensityt₁/₂). (D) The percentage of LPL synapses in brain regions and subregions. Superficial layers of the isocortex (numbered) contain the most LPL synapses. Dotted line indicates average for whole brain (6.7%). Brain subregions are listed in Table S1.
Fig. S11. Related to Fig. 5. Changes in protein lifetime and in the architecture of synapse subtypes with different protein lifetimes across the lifespan. (A) Example images of sagittal sections of PSD95-HaloTag mice injected with SiR-Halo at 3 weeks, 3 months and 18 months of age and imaged at day 0 and day 7 post injection. Scale bars: 1 mm. (B) Percentage increases in PSD95 puncta half-life ($\text{PSD95 density}_{t/2}$) across 110 brain subregions between 3 weeks and 18 months. \( n=7-10 \) Psd95$^{\text{HaloTag/HaloTag}}$ animals per time point and age group. (C) Heatmaps showing the change (Cohen’s $d$ effect size) in synapse subtype (subtype number shown at top) density in brain regions between 3 week old and 3 month old (i-iii) and between 3 month old and 18 month old (iv-vi). Subtypes are ranked with longest lifetime on the left and shortest on the right. The thirty synapse subtypes are shown together (i,iv) and separated into PSD95-only subtypes (ii,v) and PSD95+SAP102 subtypes (iii,vi). Only significant differences (\( P<0.05 \), Benjamini-Hochberg correction) are shown in color. \( n=4-5 \) Psd95$^{\text{eGFP/+};\text{Sap102}^{\text{mKO2/y}}}$ mice per age group. See Protein Lifetime Synaptome Atlas for half-life and subtype density values in each subregion (Bulovaite et al., 2021a).
Fig. S12. Related to Fig. 6. Synapse protein lifetime increases in Dlg2 mutant mice. The increase (Cohen’s d effect size) in PSD95 puncta density half-life in 110 adult brain subregions in mice carrying a heterozygous (right) or homozygous (left) Dlg2 mutation. Asterisks indicate significant differences from control (Dlg2+/+) mice (P<0.05, Benjamini-Hochberg correction). Brain subregions are listed in Table S1. n=5-8 mice for each time point and genotype (note: animals are heterozygous for PSD95-HaloTag).

Table S1. Brain regions and subregions. Column A, subregion number; column B, subregion abbreviation; column C, subregion name; column D, brain region.