Expression of Synaptotagmin in *Drosophila* reveals transport and localization of synaptic vesicles to the synapse

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**SUMMARY**

Synaptotagmin is a synaptic vesicle-specific integral membrane protein that has been suggested to play a key role in synaptic vesicle docking and fusion. By monitoring Synaptotagmin’s cellular and subcellular distribution during development, it is possible to study synaptic vesicle localization and transport, and synapse formation. We have initiated the study of Synaptotagmin’s expression during *Drosophila* neurogenesis in order to follow synaptic vesicle movement prior to and during synapse formation, as well as to localize synaptic sites in *Drosophila*. In situ hybridizations to whole-mount embryos show that *synaptotagmin* (*syt*) message is present in the cell bodies of all peripheral nervous system neurons and many, if not all, central nervous system neurons during neurite outgrowth and synapse formation, and in mature neurons. Immunocytochemical staining with antisera specific to Synaptotagmin indicates that the protein is present at all stages of the *Drosophila* life cycle following germ band retraction. In embryos, Synaptotagmin is only transiently localized to the cell body of neurons and is transported rapidly along axons during axonogenesis. After synapse formation, Synaptotagmin accumulates in a punctate pattern at all identifiable synaptic contact sites, suggesting a general role for Synaptotagmin in synapse function. In embryos and larvae, the most intense staining is found along two broad longitudinal tracts on the dorsal side of the ventral nerve cord and the brain, and at neuromuscular junctions in the periphery. In the adult head, Synaptotagmin localizes to discrete regions of the neuropil where synapses are predicted to occur. These data indicate that synaptic vesicles are present in axons before synapse formation, and become restricted to synaptic contact sites after synapses are formed. Since a similar expression pattern of Synaptotagmin has been reported in mammals, we propose that the function of Synaptotagmin and the mechanisms governing localization of the synaptic vesicle before and after synapse formation are conserved in invertebrate and vertebrate species. The ability to mark synapses in *Drosophila* should facilitate the study of synapse formation and function, providing a new tool to dissect the molecular mechanisms underlying these processes.

Key words: synapse, synaptic vesicle, neurogenesis, fruitfly, neuromuscular junction, *Drosophila*, Synaptotagmin

**INTRODUCTION**

Neurogenesis in both vertebrate and invertebrate species requires a complex maturation of groups of neurons that form highly organized synaptic contacts with a variety of target tissues. Communication at these synapses involves release of neurotransmitters from synaptic vesicles that fuse with the presynaptic membrane following the influx of extracellular calcium. Synaptic vesicle proteins are then retrieved from the presynaptic membrane and recycled into synaptic vesicles for further rounds of neurotransmitter release (for review, see Sudhof and Jahn, 1991). The biochemical events underlying neurotransmitter release and the trafficking of synaptic vesicles may be similar in both vertebrate and invertebrate synapses, since electrophysiological data and morphological analyses at the light and electron microscopy level indicate conserved functional and structural properties of the synapse (Jan and Jan, 1976; Budnik et al., 1990). Hence, proteins of the synaptic vesicle that are conserved throughout evolution are likely to play similar functions in distantly related organisms. One such family of proteins are the Synaptotagmins, integral membrane proteins of synaptic vesicles, which exhibit 57% overall identity between rat (when discussing activities and homology we will be referring to rat Synaptotagmin I) and *Drosophila* (Perin et al., 1991a). Synaptotagmin has been identified as part of a multi-gene family in several vertebrate species (Geppert et al., 1991; Wendland et al., 1991). All isoforms contain a single transmembrane region and a cytoplasmic carboxy-terminal moiety with two repeats homologous to a domain known to be involved in calcium-dependent membrane interactions (Perin et al., 1990).
addition, Synaptotagmin has been shown to bind acidic phospholipids in a calcium-dependent manner and to mediate the bridging of membranes (Perin et al., 1990; Brose et al., 1992). Synaptotagmin has also been reported to interact with ω-conotoxin-sensitive calcium channels (Leveque et al., 1992), the presynaptic receptor for latrotoxin (Petrenko et al., 1991), and another presynaptic membrane protein, syntaxin (Bennett et al., 1992). These data implicate Synaptotagmin both in docking synaptic vesicles to their presynaptic release sites, and in the calcium-dependent fusion process.

We have chosen to investigate Synaptotagmin’s role in neurotransmission in vivo and have therefore initiated this study in Drosophila. The expression pattern of Synaptotagmin in embryos may demonstrate the distribution of synaptic vesicles before and after synaptogenesis and the timing and location of synapse formation. Several general neuronal markers have been used to identify neurons and their processes in Drosophila. Using antibodies against horse radish peroxidase and glutamate, synaptic sites in the bodywall musculature of embryos (Johansen et al., 1989b) and larvae (Johansen et al., 1989a; Budnik et al., 1989, 1990) have been identified. In addition, neuronal dye fills and electron microscopy have identified some neuronal terminations in embryos (Sink and Whittington, 1991), larvae (Johansen et al., 1989a) and adults (Koenig et al., 1989; Costello and Wyman, 1986). However, relatively little is known about the location of many synapses, particularly those outside the neuromuscular junction and primary sensory centers in the adult brain. In particular, it is unclear when and where synapses form between neurons in the CNS of the embryo. Moreover, it is unclear in Drosophila whether synaptic vesicles are only made and transported after synapse formation, or whether synaptic vesicles are also made during axonogenesis and can subsequently localize to synaptic contact sites. In addition, the lack of an easily identifiable synaptic marker in invertebrates has slowed the identification of proteins that are involved in synaptic targeting and synapse formation.

We detect the presence of only one Drosophila synaptotagmin (syt) gene that is transcribed throughout development and is expressed by most, if not all, neurons. Based on the developmental expression pattern of Synaptotagmin during neurite outgrowth and synaptogenesis, we conclude that synaptic vesicles or their precursors are made and transported along the axon prior to synapse formation. Once synaptic contacts have been established, synaptic vesicles localize to these sites within a brief time period. Synaptotagmin immunocytochemistry also shows that most synapses are highly concentrated in the CNS of embryos. These antisera are the first to label most if not all synapses in Drosophila, and should be of general use to study synapse formation and function.

MATERIALS AND METHODS

cDNA cloning and sequencing
A 9- to 12-hour-old embryonic cDNA library (Zinn et al., 1988) was screened with 32P-labeled oligos and random-primed labeled cDNA fragments derived from the previously identified Drosophila syt cDNA isolated from adult flies (Perin et al., 1991a). Library filters were hybridized overnight at 42°C in 25% formamide. Filters were washed with 6x SSC, 0.1% SDS (oligos) or 2x SSC, 1% SDS (labeled fragments) twice at 50°C for 50 minutes. Sequencing of the positive cDNAs was performed using the dyeex nucleotide chain termination method (Sanger et al., 1977).

RNA blotting
Poly(A)+ RNA was prepared from developmental stages of Canton-S flies. 5 µg of RNA was electrophoresed in formaldehyde gels and blotted to nylon. Blots were probed with uniformly 32p-labeled cDNA probes corresponding to the entire coding region or smaller fragments of Drosophila syt (Perin et al., 1991a). Blots were then washed at high stringency (0.1x SSC, 0.1% SDS) and exposed for one week at −70°C with intensifying screens.

Generation of Synaptotagmin antibodies
Two rabbit polyclonal antibodies were raised against Drosophila Synaptotagmin. Antibody DSYT1 was generated against a synthetic peptide, corresponding to the 15 amino-terminal amino acids of Synaptotagmin, coupled to keyhole limpet hemocyanin. Antibody DSYT2 was prepared against a bacterial produced protein containing residues 134-474, which corresponds to the complete cytoplasmic sequence following the transmembrane region. The production of bacterial recombinant proteins was as described by Perin et al. (1991b). After induction, the bacterial pellet was resuspended, run on a preparative SDS-PAGE gel, and the appropriate band excised. 100 µg of protein was injected 5 to 6 times subcutaneously in New Zealand white rabbits before bleeding. The production of RSYT1 (prepared against rat recombinant Synaptotagmin) and RSYT2 (prepared against a peptide containing the 12 amino-terminal residues of rat Synaptotagmin) has been previously described (Perin et al., 1991b). Polyclonal antisera RSYT3 was generated against the carboxy-terminal 34 amino acids of rat Synaptotagmin 1.

In situ hybridization and immunocytochemical staining
Whole-mount in situ hybridizations were performed as described by Tautz and Pfaffle (1989). Immunocytochemical staining of embryos was carried out as described by Bellen et al. (1992). Antibody staining of third instar larvae was as described by Johansen et al. (1989a). Immunocytochemical staining of adult heads was performed as described by Han et al. (1992). Primary and secondary antibodies were preabsorbed to 0- to 9-hour-old embryonic overnight at 4°C. DSYT2 was used at a final concentration of 1:500 in embryos and larvae, and 1:1000 in adult head sections. DSYT2 staining is highly reproducible under these conditions; however, staining in the embryonic CNS is restricted after cuticle formation (after stage 17) due to poor penetration of the antibodies. This limitation can be readily overcome by disrupting the cuticle of late embryos, thus providing greater access for the antibodies. Monoclonal antibody (mAb) BP104 (Hortsch et al., 1990) was used at 1:10, and mAb. 22 C10 (Zipursky et al., 1985) at 1:10. Secondary anti-mouse and anti-rabbit antibodies were used at 1:200.

Immunoblots and trypsin digestion
Crude synaptic vesicle preparations were made by freezing 10,000-20,000 flies in liquid N2 and separating heads from bodies using a sieve. Frozen heads were crushed in liquid N2 and resuspended in 0.32 M sucrose with addition of protease inhibitors (5 µg/ml Leupeptin, 5 µg/ml Pepstatin, 5 mM Benzamidine, 0.25 mM PMSF, 10 mM EDTA, 1 mM EGTA; final concentrations) and homogenized. A 20 minute, 15,000 g spin was used to pellet cell debris and cuticle. A fraction enriched in synaptic vesicles was pelleted with a 1 hour 150,000 g spin. A similar procedure with
mammalian brain tissue yields 70-80% pure synaptic vesicles. SDS-PAGE and immunoblotting were performed as described by Perin et al. (1988). Abs. DSYT1 and DSYT2 were used at 1:200 on westerns. Antibody-reactive bands were visualized with peroxidase-labeled secondary antibodies and enhanced chemiluminescence detection (Amersham). Synaptic vesicles (100 µg of protein) were digested with 10 ng trypsin in 100 mM NaCl, 20 mM Hepes-NaOH for 1 hour at 37°C. Reactions were stopped by boiling in electrophoresis sample buffer (62 mM Tris-HCl, pH 6.8, 5% 2-mercaptoethanol, 2% SDS).

Fly stocks
Flies were raised at 24°C on standard medium with addition of baker’s yeast. All flies used in this work are Canton-S.

RESULTS

Synaptotagmin is transcribed exclusively in neurons
Two to three different syt genes have been identified in several vertebrate species (Geppert et al., 1991; Wendland et al., 1991). To determine whether different genes encoding Synaptotagmin-like proteins exist in Drosophila and to determine whether adult syt transcripts encode the same protein as in embryos, we have carried out low-stringency cDNA screens, genomic southern analysis and in situ hybridizations to embryos and polytene chromosomes. We have screened a 9-12 hour embryonic cDNA library (Zinn et al., 1988) at low stringency with the previously characterized syt cDNA from adult flies (Perin et al., 1991a). Eleven positive plaques were picked at random, cloned, restriction enzyme mapped and partly sequenced. All embryonic clones examined have similar restriction enzyme maps and identical sequences over large regions of the open reading frame (ORF) when compared to those of the adult syt cDNA (data not shown). In addition, genomic Southern blots using the embryonic and adult cDNAs as probes reveal identical bands (data not shown). The cDNAs of both adults and embryos hybridize to cytological bands 23A6-23B1 of third instar larvae polytene chromosomes. These observations suggest that all isolated cDNAs are derived from a single locus. We have also failed to identify additional Synaptotagmin isoforms with western analysis (see below). These observations suggest that a single syt gene is present in Drosophila and that the expression pattern described below is specific for syt.

To determine the complexity and size of syt transcripts, we performed a developmental northern analysis. As shown in Fig. 1, syt transcripts can be detected at all stages of development after germ band retraction. Expression of a 4.5 kb syt message appears between 8 and 12 hours of embryogenesis and is present at all subsequent developmental stages. A 7.0 kb message appears in 12-20 hour embryos and is present throughout development. An approximately 1.5 kb transcript is seen in third instar larvae and pupae. We have previously reported the presence of adult transcripts of 7 and 4.5 kb (Perin et al., 1991a). To determine the relationship of these transcripts to the previously published syt cDNA (Perin et al., 1991a), northern blots were probed with different fragments from the cDNA. Probes derived from the 5′ region of the cDNA, including both coding (bps 435-795) and noncoding sequences (bps 0-270, 0-430), identify all three transcripts. However, probes derived from the 3′ untranslated region (bps 1600-1900), as well as the 3′ coding region (1220-1900), only hybridize to the 4.5 and 7 kb messages, indicating that the 1.5 kb message may represent a transcript generated by alternative splicing or premature termination. This transcript may thus only contain approximately half of the ORF and is possibly nonfunctional. The 4.5 and 7 kb transcripts are recognized by small fragments spanning the entire coding region, suggesting that these messages may represent transcripts that differ in their untranslated region. These data indicate that syt is transcribed at most stages of development, including stages during which active synaptogenesis and neuronal modeling occur.

To study the tissue distribution of syt in embryos, whole-mount in situ hybridization experiments were performed using digoxigenin-labeled syt cDNA probes. As shown in Fig. 2, syt is first expressed in the CNS in a few neurons in each segment of the ventral nerve cord (VNC) in late stage 13 embryos. During stage 14, the pattern in the VNC becomes more complex, and the cell bodies of more neurons are labeled. At stage 15, intense staining can be found in the supra- and subesophageal ganglia, as well as in the VNC. In addition, a subset of cells of the PNS are stained. As shown in the schematic drawing and PNS in situ inset of Fig. 2E, all the cells of the PNS that stain correspond to neurons. Individual neurons from the lateral PNS cluster are labeled in Fig. 2E, demonstrating that none of the support cells including glia are labeled. Although it is difficult to determine if all neurons in the CNS express syt, many

![Fig. 1. Developmental Northern using syt cDNA as a probe. syt messages first appear in 8-12 hour embryos and are present during all stages of development. Three transcripts of approximately 7.0, 4.5 and 1.5 kb can be detected. The 4.5 kb message appears in 8-12 hour embryos and is present throughout the life cycle. The 7 kb message appears in 12-20 hour embryos and is present at all other stages of development. The 1.5 kb message is only present in third instar larvae and pupae and may represent an alternative spliced product or premature termination that results in a message that lacks significant portions of the ORF. These messages all seem to be derived from a single syt gene (see text).](image-url)
neurons in the VNC and brain are stained. During embryogenesis, we observed no syt expression in non-neuronal cells. We conclude that syt is a neuronal-specific marker, which is expressed in all PNS and many, if not all, CNS neurons during the final embryonic differentiation stages and in the fully developed embryonic nervous system.

To determine the cellular and subcellular distribution of Synaptotagmin during embryonic development, we generated polyclonal antibodies against a peptide containing the 15 amino-terminal amino acids of Synaptotagmin.

Fig. 2. Embryonic whole-mount in situ hybridizations using a digoxigenin-labeled syt cDNA. syt is expressed in the later phases of the differentiating embryonic nervous system. Embryos are shown anterior to the left, dorsal up. All views are ventrolateral views. Staging is as described by Campos-Ortega and Hartenstein (1985). (A) Stage 13-14 embryo. Note that specific cells along the midline of each segment of the ventral nerve cord (VNC) are stained. (B) Stage 15 embryo. Many more cells in the VNC and brain express syt in a segmentally repeated pattern. (C) Stage 16 embryo. There is intense staining in the supra- and suboesophageal ganglion as well as in the VNC. PNS neurons also express syt weakly. (D) Stage 17 embryo. syt is expressed abundantly in the CNS and PNS. E. A schematic diagram (redrawn from Bodmer et al. (1989)) in which all cells that express syt in the PNS are labeled in black. The photograph shows an abdominal segment of a Drosophila embryo from our in situ hybridizations with focus on the lateral cluster (lc). Cells that express syt correspond to PNS neurons. The support cells of the PNS do not express syt. Individual neurons of the dorsal cluster (dc) and ventral cluster (vc) can also be individually identified, but are not in focus in this photograph. Only the v' cluster is shown in the photograph. Abbreviations: es, external sensory organ; ch, chordotonal organ; md, multiple dendrite neurons; n, neuron; th, thecogen cell; tr, trichogen cell; to, tormogen cell; li, ligament cell; s, scolopale cell; c, cap cell; da, dendritic arbor neuron; bd, bipolar dendrite neuron; g, glial cell; td, trachea innervating cell; le, lateral external sensory neuron; lm, lateral multiple dendrite neuron; vch, v' chordotonal neuron; lch5, the five lateral chordotonal neurons. From dorsal to ventral: the dorsal cluster, the lateral cluster, and the two ventral clusters.
Synaptotagmin expression in *Drosophila* coupled to keyhole limpet hemocyanin (Ab. DSYT1). In addition, a bacterial recombinant protein containing the cytoplasmic portion of *Drosophila* Synaptotagmin (Perin et al., 1991a) was prepared and used as an immunogen (Ab. DSYT2). Polyclonal antibodies against similar domains of the rat protein (Ab. RSYT1 against recombinant rat Synaptotagmin and Ab. RSYT2 against an amino-terminal peptide) were also used in order to compare the *Drosophila* and rat proteins. As shown in Fig. 3, Ab. DSYT1 recognizes a $69 \times 10^3 \, M_r$ protein from *Drosophila* synaptic vesicles, but fails to recognize proteins of rat synaptic vesicles. This is expected because the amino-terminal domain of Synaptotagmin is highly divergent between rat and *Drosophila*. Ab. DSYT2 recognizes a protein of the same molecular weight as DSYT1 in *Drosophila* synaptic vesicles, as well as rat Synaptotagmin (see Fig. 3). Ab. DSYT2 also recognizes a protein of $55 \times 10^3 \, M_r$ in *Drosophila* synaptic vesicles, which probably represents a breakdown product of Synaptotagmin (see below). Abs. RSYT1 and RSYT2 recognize a $65 \times 10^3 \, M_r$ protein in rat synaptic vesicles. Ab. RSYT1 occasionally shows faint reactivity with a $69 \times 10^3 \, M_r$ protein of *Drosophila* synaptic vesicles. These results indicate that antisera to Synaptotagmin from *Drosophila* and rat are specific for Synaptotagmin.

Since Ab. DSYT2 recognizes two bands in synaptic vesicle preparations, we further defined the specificity of the antisera. Trypsin digestion of *Drosophila* synaptic vesicles was performed in order to examine the appearance of breakdown products of Synaptotagmin. As shown in Fig. 4A, Ab. DSYT1 recognizes a major amino-terminal trypsin fragment of approximately $31 \times 10^3 \, M_r$, while Ab. DSYT2 recognizes a carboxy-terminal fragment of approximately $38 \times 10^3 \, M_r$ in *Drosophila* synaptic vesicles. Both Ab. DSYT2 and Ab. RSYT1 recognize the same $36 \times 10^3 \, M_r$ breakdown product in rat synaptic vesicles, indicating that the DSYT2 antibody recognizes a conserved region of the cytoplasmic domain of Synaptotagmin (see Fig. 4B). The presence of a single hypersensitive trypsin site in rat Synaptotagmin (as shown in Fig. 4B) has been previously reported (Perin et al., 1991a). We find the presence of a hypersensitive site in a similar location in *Drosophila* Synaptotagmin. The additional band at $55 \times 10^3 \, M_r$ recognized by Ab. DSYT2, but not by Ab. DSYT1, appears to be an alternate step in the breakdown of Synaptotagmin. Indeed, trypsin digestion eliminates the $55 \times 10^3 \, M_r$ band with a concomitant increase in the $38 \times 10^3 \, M_r$ product and without production

**Fig. 3.** Immunoblots of partially purified synaptic vesicles of *Drosophila* and rat using antibodies against the amino-terminal amino acids, DSYT1 and RSYT2, and the remainder of the protein, DSYT2 and RSYT1, are shown. The antibodies recognize a $69 \times 10^3 \, M_r$ protein in *Drosophila* and a $65 \times 10^3 \, M_r$ protein in rat. Note that the DSYT2 antibody recognizes Synaptotagmin of both species. DSYT2 consistently recognizes two protein bands in *Drosophila* synaptic vesicles (see Fig. 4).

**Fig. 4.** Western analysis of synaptic vesicle preparations digested with trypsin. The DSYT1 and DSYT2 antibodies recognize independent domains of Synaptotagmin. (A) Upon trypsin digestion of synaptic vesicles DSYT1 identifies a amino-terminal $31 \times 10^3 \, M_r$ moiety, whereas DSYT2 recognizes a $38 \times 10^3 \, M_r$ carboxy-terminal fragment, as well as a $55 \times 10^3 \, M_r$ protein that is probably an intermediate in the breakdown process (see text). (B) The DSYT2 and RSYT1 antibodies both recognize the same $36 \times 10^3 \, M_r$ trypsin breakdown product in rat synaptic vesicles, providing additional evidence that these antibodies specifically recognize Synaptotagmin.
of other trypsin products. This $55 \times 10^3 \, M_r$ protein lacks the amino-terminal epitope recognized by Ab. DSYT1 and probably represents a breakdown product lacking the terminal portion of the luminal domain. Additional evidence indicating that the $55 \times 10^3 \, M_r$ protein is indeed Synaptotagmin, and not a cross-reacting protein, was obtained from immunocytochemical staining of embryos with a small deficiency (Df (2L) C144; 23A1-23C; Jeff Sekalsky and Bill Gelbart, personal communication) that uncovers the syt locus. Embryos homozygous for this deficiency showed no staining with DSYT2, suggesting that the protein is encoded by a gene contained within this deficiency, and hence most likely corresponds to Synaptotagmin. To confirm this hypothesis, an antibody against the carboxy-terminal 34 amino acids of rat Synaptotagmin I, a region highly conserved between rat and Drosophila, was prepared and tested on synaptic vesicle preparations. This antisera (RSYT3) also recognized proteins of $69 \times 10^3$ and $55 \times 10^3 \, M_r$ in Drosophila synaptic vesicles, as well as a protein of $65 \times 10^3 \, M_r$ in rat synaptic vesicles (data not shown). Although we believe this product to be an alternate step in the degradation of Synaptotagmin, we cannot completely rule out that the $55 \times 10^3 \, M_r$ protein is a splicing variant of Synaptotagmin.

To determine whether Synaptotagmin is present throughout the life cycle of Drosophila, we prepared crude synaptic vesicle extracts from 0-20 hour embryos, first and third instar larvae, pupae and adults. Western analysis with Abs. DSYT1 and DSYT2 shows that a $69 \times 10^3 \, M_r$ protein and the above described breakdown products are present at all stages of development (data not shown). This observation, combined with the previous sequencing and in situ hybridization data, suggests that Synaptotagmin is present as a single isoform at all stages of development. In addition, no other bands are detected in western blots of whole extracts, indicating that the antisera do not recognize additional proteins from Drosophila.

The sequence of the syt ORF predicts a protein of $50 \times 10^3 \, M_r$ in Drosophila and $48 \times 10^3 \, M_r$ in rat. Part of the difference between observed and predicted $M_r$ in the rat protein has been shown to be due to glycosylation of the mammalian form (Perin et al., 1991b). Endoglycosylase F digestion of

Fig. 5. Immunocytochemical staining of whole-mount embryos using the DSYT2 antibody. Synaptotagmin is rapidly transported to the synapses and localizes synaptic contact sites. Anterior is to the left, dorsal is up. (A) Ventrolateral view. Stage 14 embryo. Note that many cell bodies of each segment of the ventral nerve cord (VNC) are stained. The cell bodies of the neurons are only intensely labeled for a very short period. (B) Stage 15 embryo. Lateral view, focal plane is on the center of the embryo. Note that most protein has accumulated in the dorsal part of the VNC and the brain. (C) Stage 16 embryo. Ventrolateral view. Synaptotagmin is mostly localized along the longitudinal tracts but is also present in many other areas of the CNS (see Fig. 6B). Staining can also be observed transiently in the axons of the motorneurons and the PNS neurons during this stage. The commissures of the VNC stain only weakly. (D) Stage 17 embryo. Lateral view, focal plane is on the CNS of the embryo. Synaptotagmin is mainly localized along the two longitudinal tracts of the CNS. Synaptotagmin is also present at synapses in the bodywall musculature at this stage (not in focal plane). Ab. DSYT2 also shows faint reactivity with an antigen present at low levels in a subcellular compartment of the pole cells (not shown). These cells, however, lack syt message. No other non-neuronal cells are labeled by Ab. DSYT2.
Synaptotagmin expression in *Drosophila* did not cause a \( M_r \) shift of the protein on SDS-PAGE gels (data not shown), indicating that Synaptotagmin may not be N-linked glycosylated in *Drosophila*. However, when the cytoplasmic domain of Synaptotagmin is expressed in *E. coli*, a recombinant protein of \( 41 \times 10^3 \) \( M_r \) is made, only slightly larger than the predicted molecular mass of \( 39 \times 10^3 \) (Perin et al., 1991a). Taken together, these observations suggest that other secondary modifications or secondary structure of the full-length protein may account for the molecular weight differences that we observe.

**Synaptotagmin expression reveals trafficking of synaptic vesicles and localizes sites of synaptic contact**

Ab. DSYT2 was used for immunocytochemical staining of whole-mount embryos using a secondary antibody coupled to horseradish peroxidase. A developmental profile of the expression pattern of Synaptotagmin is shown in Fig. 5. This expression pattern correlates well with the cellular distribution of *syt* seen in whole-mount in situ hybridization experiments, yet reveals a very different subcellular distribution of the protein compared to the message. Expression of Synaptotagmin is detected at low levels in late stage 13 or early stage 14 embryos in a subset of neurons of the CNS (see Fig. 5A). As shown in Fig. 6A, Synaptotagmin is localized to the cell body of clusters of neurons in the ventral part of the VNC. Some punctuate or granular staining can be seen in these neurons close to the cell membrane (see Fig. 6A). Although it is difficult to determine the precise subcellular localization at the light microscope resolution, this staining may correspond to the Golgi apparatus. At later stages of embryonic development, very little staining can be observed in the cell bodies of neurons, except transiently in the neurons of the PNS. During stage 15 and later, most Synaptotagmin accumulates along the dorsal longitudinal tracts of the VNC and brain (Fig. 5B–D). The commissures in the VNC stain only weakly (Fig. 5C). As shown in Fig. 6B, staining in the CNS is not only confined to the area along the longitudinal tracts. Punctate clusters of staining can be

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**Fig. 6.** Immunocytochemical staining of whole-mount embryos using Ab. DSYT2. Subcellular localization of Synaptotagmin during embryonic development. (A) Detailed view of the VNC of the CNS of a stage 14 embryo. Note the clusters of neurons in each segment that express Synaptotagmin. Synaptotagmin is mostly localized to a subcellular compartment associated with the membrane, possibly the Golgi complex. (B) Detailed view of the CNS of a stage 16 embryo. Most staining accumulates along the longitudinal tracts and at the sites of incoming peripheral neurons. (C) Detail of the chordotonal organs of the lateral cluster of the PNS of a stage 16 embryo (see also Fig. 1 E). Note that Synaptotagmin is transiently present in the dendrites of the chordotonal neurons (arrow head) and in the scolopales (arrow). (D) Detail of the ventral region of a stage 17 embryo with focus on the surface of the ventral muscles. The tape-like structures are muscles. The arrowheads point to multiple synapses of neuromuscular junctions of two segments.
seen lateral to the longitudinal tracts, and larger and smaller clusters alternate at the positions were the intersegmental and segmental nerves connect to the CNS. Thus, we conclude that most synapses in the CNS are located along the longitudinal tracts of the VNC and brain, and that they are non-randomly distributed.

In the periphery, expression of Synaptotagmin can be observed transiently in the afferent neurons of the sensory PNS. Expression of Synaptotagmin in these neurons is weak. During stages 15 and 16, the axons of PNS neurons also label transiently (Fig. 5C), suggesting that synaptic vesicles or their precursors are present in axons prior to synapse formation. As shown in Fig. 6C, expression of Synaptotagmin is not only restricted to axons, since the dendrites of the chordotonal neurons and their scolopales also contain Synaptotagmin transiently. In stage 17 embryos, staining in the periphery is confined to a punctate staining pattern at neuromuscular junctions. As shown in Fig. 6D, expression at these synapses is quite abundant. Double-labeling studies using immunofluorescence with MAb 22C10 (a marker for neuronal cell membranes) or MAb BP104 (a marker for neuroglian, a cell adhesion molecule) and Ab. DSYT2 indicate that all neuromuscular junctions in a particular abdominal segment express Synaptotagmin. As shown in Fig. 7A and B, the pattern of synaptic connections within a segment is highly organized and conserved throughout the segments. However, embryonic neuromuscular junctions within each segment show a variety of shapes and structures (see Fig. 7C). Some are delta-like, others are crescent shaped, and still others are straight. These synaptic contact sites increase in button number and distribution as development proceeds through the larval stages. Synaptic vesicle clustering at neuromuscular junctions of the Drosophila embryo occurs essentially simultaneously throughout each segment during stage 16. The synaptic organization of the body wall musculature innervated by the segmental and intersegmental nerve, as revealed by Synaptotagmin localization, is in agreement with that previously described by Johansen et al. (1989b).

Synaptotagmin is also expressed at synaptic contact sites
Synaptotagmin expression in *Drosophila* at later stages of development. Fig. 8 shows Synaptotagmin staining in third instar larva. Synaptotagmin can be readily detected at synapses between nerve and muscle in the periphery. Neuromuscular junctions in the bodywall musculature of *Drosophila* have been classified into two categories based on button size and axonal branching (Johansen et al., 1989a). Type I projections are characterized by large buttons with few branches. Type II processes are much longer and thinner and have many smaller synaptic terminals along the muscle surface. Both type I (Fig. 8A,C) and type II (Fig. 8B,C) processes show Synaptotagmin immunoreactivity at synaptic terminals. Our antisera also labeled synapses in physically disrupted whole-mount third instar larval brains, revealing intense punctate labeling.

Ab. DSYT2 is also useful as a marker for synapses in the adult fly (see Fig. 9). The antibody shows reactivity throughout the neuropil in adult fly head sections with intense staining in the primary sensory centers like the antennal lobes (Fig. 9A,B,G) and visual lamina (Fig. 9C,D,E,H). Fig. 9G shows an enlarged view of the antennal lobes, where one can identify the glomerular-like islets (Strausfeld, 1976) where antennal nerve fibers synapse with dendrites of interneurons. Higher order synaptic structures such as the medulla, lobula and lobular plate, fan-shaped body and mushroom body also label. Control sections processed with preimmune sera or without addition of primary antibody lacked such staining. We therefore infer that Synaptotagmin maintains its synaptic localization throughout development and can be used as a synaptic marker at all stages of development.

**DISCUSSION**

Neurotransmitter release requires the action of numerous proteins at the presynaptic nerve terminal. A substantial number of these are associated with the synaptic vesicle and many have now been isolated and biochemically characterized (Sudhof and Jahn, 1991). Although specific roles for many of these proteins have been proposed on the basis of biochemical experiments, their function has not been established in vivo. One of these proteins, Synaptotagmin, has been shown to be highly conserved evolutionarily, exhibiting 57% sequence identity between rat and *Drosophila* homologues (Perin et al., 1991a). We have therefore chosen to examine Synaptotagmin expression during *Drosophila* development.

We find no evidence that *Drosophila* contains multiple Synaptotagmin isoforms encoded by different genes as observed in several vertebrate species (Geppert et al., 1991; Wendland et al., 1991). Our results suggest that there is only a single *syt* gene that encodes a single isoform of Synaptotagmin, simplifying the interpretation of the expression patterns discussed below and facilitating the future genetic dissection of *syt*. In situ hybridizations and immunocytochemical stainings show that *syt* is expressed in most, if not all, embryonic neurons, consistent with its hypothesized neuronal function. Hence, analysis of *syt* may provide insights into many aspects of neuronal function, synaptic vesicle trafficking and synaptic localization.

It has been shown that Synaptotagmin is localized
Fig. 9. Immunocytochemical staining of adult fly head sections stained with Ab. DSYT2 using a secondary antibody coupled to horse radish peroxidase. Sections have been arranged anterior to posterior (A-F) and dorsal up. Note the accumulation of synapses in the primary sensory centers of the antennal lobes (A,B,G) and visual lamina (C,D,E,H). The glomerular-like islets where the antennal nerve synapses onto interneuron dendrites is shown in G. The layered synaptic structure of the visual lamina is shown in H. Abbreviations: l, lamina; m, medulla; lp, lobula and lobular plate; c, mushroom body calyx; f, fan shaped body; n, noduli; sog, suboesophageal ganglion; a, antennal lobe; ap, anterior prominence of the inferior medial protocerebrum.
due to transport of newly synthesized protein from the cell
muscles, we see mobilization of Synaptotagmin to discrete
body to the synapse. However, the lack of detectable Synap-
totagmin suggests that synaptic vesicle exocytosis might play a role
prior to synaptogenesis. In

projecting from and to the CNS suggest that transport of

also transiently present in the chordotonal dendrites in the

PNS. By stage 17, this staining pattern is no longer
detectable. It is thus likely that Synaptotagmin is transiently
present in dendrites, are redistributed to axons, or whether
there is simply a barrier to further synaptic vesicle
movement into dendrites is unknown.

Localization of Synaptotagmin protein provides a general
marker allowing the localization of synaptic populations
regardless of neurotransmitter content in the

Drosophila

embryo. This can most easily be illustrated in the PNS. The
innervation pattern of bodywall muscle fibers by efferent
motor axons, as revealed by Synaptotagmin expression, cor-
responds to that previously reported using immunocyto-
chemistry with anti-glutamate and anti-HRP antibodies
(Johansen et al., 1989b). However, in the CNS, little is
known about the spatial distribution of the synapses. Inter-
estingly, the intense staining along the longitudinal tracts of
the VNC and the brain suggests that most synaptic connec-
tions are made in the vicinity of, or along these tracts. A
portion of the staining along the longitudinal tracts could be
due to transport of newly synthesized protein from the cell
body to the synapse. However, the lack of detectable Synap-
totagmin staining in the axons of stage 17 peripheral neurons
projecting from and to the CNS suggest that transport of
Synaptotagmin along axonal tracts is minimal and not easily
detected by antibody staining after synapses are formed. In
addition, the commissures of the CNS stain very faintly
when compared to the longitudinal tracts. If transport were
responsible for the strong staining along the longitudinal
tracts, then commissures should stain equally strong. An
additional possibility is that there is a preponderance of CNS
neurons that are undergoing axonogenesis during stage 17
and thus the staining could represent transport along these
tracts. This possibility is unlikely because we also detect
intense staining along the VNC in larva, when axonogen-
esis is completed. Indeed, single neuronal dye fills of
embryonic motor neurons show dramatic arborizations of
the axon along the longitudinal tracts, even after formation
of its synaptic contact with muscle fibers (Sink and Whitt-
tington, 1991). We therefore conclude that most synapses
are localized along the longitudinal tracts of the CNS. Alter-
natively, an additional Synaptotagmin protein exists which
is specific for other CNS synapses and is not detected by our
antibodies. Although we cannot rule out this possibility, we
have been unable to find evidence for other

Drosophila

Synaptotagmins.

The compartmentalization of Synaptotagmin staining also
demonstrates the timing of synapse formation in the PNS
and, by analogy, the CNS. Synaptotagmin is first detected
in stage 13 embryos in the CNS, just after axonogenesis is
initiated (Campos-Ortega and Hartenstein, 1985). During
stages 14 and 15, Synaptotagmin accumulates outside the
cell bodies in a punctate pattern along the tracts of the CNS,
indicating connections among neurons of the CNS are
formed during these stages. The targeting of synaptic
vesicles to discrete sites occurs essentially simultaneously
in all segments in the periphery during stage 16, suggesting
that the molecular signals that specify synapse formation
occur throughout the embryo in a synchronous fashion. The
events that set up specific synapses at different muscles
seem to be somewhat different for each neuromuscular
junction in each segment since the shape and length of the
synapses vary widely. However, specific synapses at similar
muscles in different segments are very similar. The
molecular mechanisms that are involved in axonal target
selection and synapse formation are largely unknown.
However, there is substantial evidence that a number of
extracellular matrix proteins, secreted proteins and receptors
may play crucial roles in synaptic targeting (Hunter et al.,
1989; Nose et al., 1992; Reichardt and Tomaselli, 1991;
Reist et al., 1992; Ushkaryov et al., 1992). It is possible that
such proteins might also play a role in synaptic vesicle clus-
tering, perhaps to cause local rearrangements of cytoskele-
tal elements at the contact site that might subsequently
sequester synaptic vesicles to release sites. Alternatively,
these proteins might participate in a signal transduction
pathway that transports newly synthesized synaptic vesicles
only to the synapse.

Synaptotagmin is also expressed in most embryonic stages
as indicated by northern and western blots. This was
confirmed by immunocytocchemical staining of third instar
larvae, which demonstrates that Synaptotagmin is present at
synapses in both type I and type II processes of motor
neurons. As expected, the CNS contains numerous synapses
which can be most readily visualized in disrupted nervous
systems. In adult fly head sections immunoreactivity is
present throughout the neuropil of the CNS, with intense
staining in the primary sensory terminations in the antennal
lobe and visual lamina, as well as in higher order brain
regions. Hence, the antisera that we have generated can be
used at all stages of development to identify synaptic contact
sites. These antisera are the first general synaptic markers to
be described in *Drosophila* and should therefore be an important tool to address questions concerning synaptic localization and function.

The temporal expression pattern and precise synaptic localization of Synaptotagmin in *Drosophila* and rat, along with the strong sequence similarity between the two proteins, suggest that Synaptotagmin plays a conserved role in synaptic function from invertebrates to vertebrates. In order to investigate the precise role of Synaptotagmin in neurotransmitter release, we have localized the *syt* gene to polytene chromosomal bands 23A6-23B1 and have begun a genetic analysis to obtain mutations in *syt*.

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