Polysialylation of the Synaptic Cell Adhesion Molecule 1 (SynCAM 1) Depends Exclusively on the Polysialyltransferase ST8Siall \textit{in Vivo}^*\[5\]

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\textbf{Background:} Polysialic acid is a developmentally regulated posttranslational modification.

\textbf{Results:} Loss of the polysialyltransferase ST8Siall but not ST8SialIV abolished polysialylation of SynCAM 1 in the mouse brain.

\textbf{Conclusion:} Polysialylation of SynCAM 1 is mediated by ST8Siall throughout postnatal mouse brain development.

\textbf{Significance:} Studying the molecular requirements for protein polysialylation is crucial for understanding how this process is regulated.

Polysialic acid is a unique carbohydrate polymer specifically attached to a limited number of glycoproteins. Among them is synaptic cell adhesion molecule 1 (SynCAM 1), a member of the immunoglobulin (Ig) superfamily composed of three extracellular Ig-like domains. Polysialylation of SynCAM 1 is cell type-specific and was exclusively found in NG2 cells, a class of multifunctional progenitor cells that form specialized synapses with neurons. Here, we studied the molecular requirements for SynCAM 1 polysialylation. Analysis of mice lacking one of the two polysialyltransferases, ST8Siall or ST8SialIV, revealed that polysialylation of SynCAM 1 is exclusively mediated by ST8Siall throughout postnatal brain development. Alternative splicing of the three variable exons 8a, 8b, and 8c can theoretically give rise to eight transmembrane isoforms of SynCAM 1. We detected seven transcript variants in the developing mouse brain, including three variants containing exon 8c, which was so far regarded as a cryptic exon in mice. Polysialylation of SynCAM 1 was restricted to four isoforms in perinatal brain. However, cell culture experiments demonstrated that all transmembrane isoforms of SynCAM 1 can be polysialylated by ST8Siall. Moreover, analysis of domain deletion constructs revealed that Ig1, which harbors the polysialylation site, is not sufficient as an acceptor for ST8Siall. The minimal polypeptide required for polysialylation contained Ig1 and Ig2, suggesting an important role for Ig2 as a docking site for ST8Siall.

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Synaptic cell adhesion molecule 1 (SynCAM 1) is a member of the immunoglobulin superfamily that was identified in the nervous system as a potent inducer of synapse formation (1). SynCAM 1 is prominently expressed in the developing and mature brain, mediating Ca\(^{2+}\)-independent homo- and heterophilic interactions across the nascent and mature synaptic cleft (1–3). In developing neurons, SynCAM 1 shapes migrating growth cones, assembles at axo-dendritic contacts, and participates in adhesive \textit{trans} interactions that induce presynaptic specializations (4, 5). Moreover, studies on genetic mouse models with increased or no SynCAM 1 expression demonstrated a crucial role of this synapse-organizing molecule in regulating the number and plasticity of excitatory synapses (6).

SynCAM 1 is a single-spanning membrane protein with three extracellular Ig-like domains and a short cytosolic part (1). The first Ig-like domain provides the binding interface for homo- and heterophilic \textit{trans} interactions, whereas Ig2 and Ig3 were shown to drive \textit{cis} oligomerization of SynCAM 1 (5, 7). The three Ig-like domains contain six potential N-glycosylation sites, and the presence of N-glycans at Asn-80 and Asn-104 in Ig1 was demonstrated to be essential for synapse induction by promoting adhesive \textit{trans} interactions of SynCAM 1 (7).

Genetic and bioinformatic characterization of the human and murine SynCAM 1 gene revealed that they are composed of 12 and 11 exons, respectively. Alternative pre-mRNA splicing results in the formation of several transmembrane isoforms and a secreted form that encompasses only the Ig-like domains of SynCAM 1 (8–11). In the case of human SynCAM 1, differential usage of three alternative exons, here termed exons 8a, 8b, and 8c, can theoretically lead to eight membrane-bound isoforms, which differ only in a short juxtamembranous extracellular stem region. In mice, however, the variable exon 8c has been described as cryp-

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The abbreviations used are: SynCAM 1, synaptic cell adhesion molecule 1; Cadm1, cell adhesion molecule 1; endoN, endosialidase; FnIII, fibronectin type III domain; NCAM, neural cell adhesion molecule 1; Neu5Ac, 5-n-acylneuraminic acid; pAb, polyclonal antibody; Pn, postnatal day n; PNGase F, peptide-N-glycosidase F; polysia, polysialic acid; polyST, polysialyltransferase.
targets for polysialylation, and (iii) the protein parts of SynCAM in perinatal brain development, (ii) the SynCAM 1 isoforms that are functional role of polySia in regulating SynCAM 1-mediated interactions of NG2 cells (13). Moreover, this fraction of polySia-SynCAM 1 was found to be restricted to a particular cell population called NG2 cells (13). These multifunctional progenitor cells form unique synaptic associations with neurons through which they receive excitatory synaptic inputs (28–30). On SynCAM 1, polySia is attached selectively to N-glycans at Asn116 located in the first Ig-like domain (13). Polysialylation completely abolished homophilic SynCAM 1 binding in vitro, suggesting a functional role of polySia in regulating SynCAM 1-mediated interactions of NG2 cells (13).

In the present study, we investigated the molecular requirements for SynCAM 1 polysialylation by determining (i) the enzyme responsible for SynCAM 1 polysialylation during postnatal brain development, (ii) the SynCAM 1 isoforms that are targets for polysialylation, and (iii) the protein parts of SynCAM 1 that serve as major acceptor structure.

**Experimental Procedures**

**Antibodies, Enzymes, and Reagents**—The anti-polySia monoclonal antibody (mAb) 735 (IgG 2a)(31) and endosialidase (endoN) were purified as described previously (32, 33). The rabbit polyclonal antibody (pAb) directed against the C-terminal domain of SynCAM 1 was purchased from Sigma-Aldrich. Chicken anti-SynCAM 1 mAb 3E1 (IgY) was obtained from MBL, and mouse anti-SynCAM 1 mAb L45/30 (IgG 1) was obtained from the University of California Davis/National Institutes of Health NeuroMab Facility. Rabbit anti-human IgG-Fc pAb was purchased from Bethyl Laboratories. Horseradish peroxidase-conjugated antibodies were from Southern Biotech. Peptide-N-glycosidase F (PNGase F) was purchased from Roche Applied Science.

**Animals—St8sia2–/–, St8sia4–/–, and Ncam–/–** mice (22, 23, 34) have been backcrossed to the C57BL/6j genetic background for six generations. St8sia2–/– mice were provided by Jamey Marth (University of California, Santa Barbara, CA), and Ncam–/– mice were provided by Harold Cremer (Developmental Biology Institute of Marseille Luminy, Campus de Luminy, Marseille, France). All protocols for animal use were in compliance with the German law for the protection of animals and approved by the local authorities.

**Reverse Transcription-PCR Analysis**—Total RNA from mouse brains was extracted using TRIzol reagent (Invitrogen) according to the manufacturer’s instructions. 2.5 μg of total RNA of C57BL/6j mouse brain was reverse transcribed using RevertAid reverse transcriptase (Fermentas) according to the manufacturer’s recommendations. Amplification of SynCAM 1-specific cDNA was performed by PCR with the primers SK4s (5’-GATGAAGTACTCTTTCTTTTCTTCG-3’) and SK5as (5’-AAAAATAGCGGCCCCAGAATGATGAGCACA-3’). PCR products were separated on 3% agarose gels and stained with ethidium bromide. To verify the identity of the PCR products, individual bands were excised from the gel. Extracted DNA was subcloned into the TOPO-TA cloning vector (Invitrogen) and analyzed by sequencing.

**Expression Plasmids**—Full-length cDNA of murine SynCAM 1 was transcribed from 1 μg of total RNA of perinatal C57BL/6j mouse brain using the SuperScript first-strand synthesis system (Invitrogen). SynCAM 1 cDNAs of the transcript variants 8–, 8a, 8b, 8b, 8bc, and 8abc were amplified by PCR with the primer pair MR68s (5’-GATCGGTACCATGGCGAGTGCT-GAAGTACTCTTTCTTTTCTTCG-3’) and MR69as (5’-GCCATGCGGCCGCGCTTGTTTGTTTCTTTCTCAG-3’). After digestion with KpnI and NotI, the obtained PCR products were ligated into the corresponding sites of pcDNA3.1-Zeo (Invitrogen). The identity of each PCR product was confirmed by sequencing and sequences of variants 8–, 8a, 8b, 8bc, and 8abc were identical to the reported sequences with GenBank™ accession numbers NM_001025600, NM_018770, NM_207675, NM_207676, and NM_018772.

For generation of the plasmid encoding a SynCAM 1-Fc chimera composed of all three Ig-like domains of SynCAM 1 fused to the Fc part of human IgG1, the region encoding the corresponding extracellular domain of SynCAM 1 (amino acids 1–349) was amplified by PCR with primers MR91s (5’-GACT-
RESULTS

Polysialylation of SynCAM 1 during Postnatal Mouse Brain Development Depends Strictly on ST8SiaII—To dissect the contribution of the two polysialyltransferases, ST8SiaII and ST8SiaIV, to SynCAM 1 polysialylation, we compared the polysialylation level of SynCAM 1 in the postnatal brain of wild-type, ST8sia2−/−, and ST8sia4−/− mice. For each genotype, whole brains were collected at eight different time points between postnatal day 1 (P1) and P21 and from adult mice. SynCAM 1 was immunoprecipitated from brain lysates by a polyclonal antibody directed against the intracellular C-terminal domain of SynCAM 1. The isoform of interest was analyzed by Western blotting either with the polysialylation-specific mAb 735 to detect polysialylated SynCAM 1 (Fig. 1A) or anti-SynCAM 1 mAb 3E1 to visualize the complete fraction of immunoprecipitated SynCAM 1 (Fig. 1B). In brain homogenates of wild-type mice, analysis with mAb 735 revealed a broad band with an apparent molecular mass ranging from ~80 to ~110 kDa (Fig. 1A, top), representing the polysialylated form of SynCAM 1 (13). Polysialylated SynCAM 1 was detected at an almost constant level from P1 to P9, followed by a slight decrease toward P11. Between P11 and P15, however, a sharp drop in the polysialylation of SynCAM 1 was observed. At P15, minimal amounts of polysialylated SynCAM 1 were still visible, but the signal reached the detection limit thereafter. In contrast, the overall expression level of SynCAM 1 stayed nearly constant (see top of Fig. 1B for SynCAM 1 analyzed after immunoprecipitation and supplemental Fig. S1A for total SynCAM 1 expression monitored by direct Western blot analysis of brain lysates). The total fraction of SynCAM 1 appeared as a diffuse band centering at 85 kDa throughout the first postnatal week. From P7 on, the SynCAM 1 signal started to extend toward the lower molecular mass range, and from P15 to adulthood, two prominent signals at ~95 and ~50 kDa were observed. Age-dependent variations in the SynCAM 1 protein pattern have been described previously and may reflect developmental changes in the iso- and glycoform pattern (1, 10).

Compared with wild-type animals, polysialylation and overall expression level of SynCAM 1 were unchanged in ST8sia4−/− mice. Moreover, down-regulation of polysialylated SynCAM 1 followed the same time course (bottom panels in Fig. 1, A and B, respectively). Conversely, loss of ST8siaII resulted in a complete absence of polysialylated SynCAM 1 at all time points analyzed (see Fig. 1A, middle, and supplemental Fig. S1B for data obtained after prolonged exposure time). In contrast, no changes in the overall expression of SynCAM 1 were observed (middle panel of Fig. 1B and supplemental Fig. S1A). These data highlight that throughout postnatal brain develop-
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FIGURE 2. SynCAM 1 transcript variants expressed during postnatal mouse brain development. A, schematic representation of the murine SynCAM 1 gene (Cadm1). Exons that are subject to alternative splicing were designated as 8a, 8b, and 8c. The dashed lines indicate which protein parts are encoded by particular exons. Peptides encoded by the variably spliced exons 8a, 8b, and 8c are shown as white, gray, and black boxes, respectively. Their peptide sequences are given together with the number of predicted O-glycosylation sites in parentheses. The arrows indicate the binding sites of the exon 5-specific forward primer SK4 and the exon 9-specific reverse primer SK5 used for RT-PCR in B. B, RT-PCR analysis of SynCAM 1 transcript variants during postnatal mouse brain development. Transcript variants encoding membrane-bound isoforms of SynCAM 1 were amplified by RT-PCR from total RNA of mouse brains collected at P1, P3, P5, P7, P9, P15, P21, and adult stage. PCR products were separated on a 3% agarose gel and assigned to individual transcript variants according to size (see arrows on the right). TMD, transmembrane domain.

For informative designation of SynCAM 1 transcripts and isoforms, we named exons and variable peptides as shown in Fig. 2A. The N-terminal signal peptide of SynCAM 1 is encoded by exon 1, and the three Ig-like domains are encoded by exons 2–7. The three variable exons that encode short peptides located in the juxtamembranous extracellular stem region were designated as 8a, 8b, and 8c. The transmembrane region is encoded by exon 9 and the C-terminal intracellular domain by exon 10. For unequivocal designation of each transcript variant, we named them according to the absence (variant 8−) or presence of variable exons (e.g. variant 8b contains only exon 8b, whereas variant 8abc contains all variable exons). Consistent with the current view that the variable exon 8c is a cryptic exon in mouse (8, 10), we found no exon 8c-containing murine sequences in the databases when we searched specifically for expressed sequence tags. However, by general BLAST search analysis, we found partial and/or complete coding sequences for the exon 8c-containing murine transcript variants 8abc (GenBank™ accession number EU541476), 8bc (accession numbers EU541477 and AB37719), and 8c (accession number EU541478).

To investigate which transcript variants of SynCAM 1 are expressed during postnatal mouse brain development, we performed RT-PCR analysis with an exon 5-specific forward and an exon 9-specific reverse primer as described by Hagiyama et al. (10). Based on the calculated fragment size, five of the obtained PCR products could be attributed to transcript variants 8−, 8b, 8c, 8ab, and 8abc (see Fig. 2B). Transcript variants 8a and 8bc differ only by 3 bp, and the corresponding PCR products could not be separated. However, sequencing of the PCR products demonstrated that during postnatal brain development, all possible combinations of the three variable exons were expressed with the single exception of variant 8ac, which was not found at any developmental stage.
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Notably, transcript variant 8abc showed a constantly low expression level throughout postnatal brain development, which might have hampered the identification of this variant in previous studies (10, 11). Transcript variant 8c was hardly detectable before P7/P9 but was constantly expressed thereafter. The most prominent age-dependent changes were seen for variant 8−, which was strongly up-regulated between P1 and P21 and appeared as the predominant transcript variant in the adult mouse brain.

Polysialylation of SynCAM 1 in Perinatal Mouse Brain Was Restricted to Four Isoforms—To analyze which SynCAM 1 isoforms serve as targets for polysialylation in vivo, we isolated the polysialylated fraction of SynCAM 1 from P1 brain by immunoprecipitation with the anti-polySia mAb 735. Because more than 95% of all polySia is attached to NCAM in the perinatal brain of wild-type mice (13), we used brains of Ncam−/− mice for efficient isolation of polySia-SynCAM 1. As shown previously, polysialylation of SynCAM 1 occurs independently of NCAM polysialylation and is identical in Ncam−/− and wild-type mice. The isolated polysialylated fraction of SynCAM 1 was characterized before and after specific removal of polySia by endoN or after de-N-glycosylation by PNGase F. After separation by SDS-PAGE, the material was analyzed by Western blotting using two different anti-SynCAM 1 antibodies: mAb 3E1 directed against the extracellular part of SynCAM 1 shared by all isoforms and mAb L45/30 directed against the variable peptide encoded by exon 8c (see Fig. 3B for schematic representation of the antibody binding sites).

In the case of mAb 3E1 (Fig. 3A, left), a broad signal with an apparent molecular mass of 90–125 kDa was detected. Removal of polySia by endoN resulted in a moderate increase in electrophoretic mobility, whereas four discrete bands appeared after de-N-glycosylation by PNGase F. Their apparent molecular masses of about 50, 60, 70, and 80 kDa exceeded the masses of 42–48 kDa calculated for the unmodified SynCAM 1 isoforms, suggesting the presence of O-glycans. Moreover, the bands at 60 and 80 kDa were also detected by mAb L45/30 (Fig. 3A, right) which is specific for the peptide encoded by exon 8c (supplemental Fig. S2, bottom).

To assign the four bands obtained after de-N-glycosylation to particular isoforms, we compared their electrophoretic mobilities with the mobility of individual de-N-glycosylated SynCAM 1 isoforms. To obtain these marker proteins, we generated full-length cDNAs for each transmembrane isoform and expressed them individually in CHO cells. After immunoprecipitation by an anti-SynCAM 1 pAb recognizing the C-terminal domain, all isoforms were analyzed by Western blotting before and after PNGase F treatment (supplemental Fig. S2). All isoforms were modified by N-glycans as demonstrated by a significant increase in electrophoretic mobility after de-N-glycosylation. With the exception of isoforms 8− and 8c, the apparent molecular masses of the de-N-glycosylated isoforms exceeded the mass calculated for the unmodified polypeptide. This probably reflects the presence of O-glycans. The most prominent mass difference was seen for the peptide 8a-containing isoforms (supplemental Fig. S2), which is in agreement with 17, 4, and 2 O-glycosylation sites predicted for peptide 8a, 8b, and 8c, respectively.

Defined, de-N-glycosylated isoforms were then used as markers to assign the isoforms of SynCAM 1 that served as polySia carrier in P1 mouse brain. PolySia-SynCAM 1 was again immunoinsolated from perinatal Ncam−/− brain using the anti-polySia mAb 735. After de-N-glycosylation, the electrophoretic mobilities of the four bands were directly compared with the mobility of the de-N-glycosylated forms of each individually expressed isoform (Fig. 3C). The bands at 80 and 70 kDa co-migrated with isoforms 8abc and 8ab, respectively. The band at 60 kDa co-migrated with the upper band of isoform 8bc, which represents most likely the fully O-glycosylated form. The fourth band at ~48 kDa showed an electrophoretic mobility similar to that of isoforms 8b and 8c. However, due to the fact that this band was not recognized by the peptide 8c-specific mAb L45/30 (Fig. 3A, right), it was assigned to isoform 8b. Based on this approach, the isoforms underlying the polysialylated fraction of SynCAM 1 at P1 were identified as 8abc, 8ab, 8bc, and 8b, all characterized by the presence of an exon 8b-encoded peptide (Fig. 3E). For polySia-SynCAM 1 expressed at P7, we observed the same set of four isoforms, with 8b and 8bc being the major polySia carriers (Fig. 3D). At P11, when polysialylation of SynCAM 1 starts to decrease (Fig. 1A), less SynCAM 1 was pulled out by anti-polySia mAb 735 (Fig. 3F). For this material, only isoforms 8b and 8bc were detected as underlying polySia carriers (Fig. 3D). Isoforms 8ab and 8abc, which had been observed as minor polySia carriers at P1, might have been below the detection limit at P11 due to the observed overall decrease in SynCAM 1 polysialylation at this time point.

ST8SiaII Can Polysialylate All Transmembrane-bound Isoforms of SynCAM 1—Aimed at understanding the impact of the variable peptides for efficient polysialylation of membrane-bound SynCAM 1, we studied polysialylation of individual isoforms in cell culture experiments. However, all of our initial attempts to generate polysialylated SynCAM 1 by transient or stable co-expression of ST8SiaII and SynCAM 1 failed. After we had established which isoforms of SynCAM 1 were polysialylated in vivo, we selected one of them, namely isoform 8abc, and optimized co-expression conditions in the murine cell line LMTK−. This fibroblast cell line lacks endogenous expression of SynCAM 1 (supplemental Fig. S3) and is negative for NCAM and both polySTs (32). Under standard transfection conditions, SynCAM 1 was extremely well expressed, whereas only small amounts of polySia-SynCAM 1 were generated (supplemental Fig. S3). Decreasing the amount of SynCAM 1 plasmid resulted in decreased expression of SynCAM 1 but in parallel resulted in increased levels of polySia-SynCAM 1, suggesting a detrimental effect of increased SynCAM 1 expression on the polysialylation of this acceptor molecule. Optimal polysialylation of SynCAM 1 was achieved by transient co-transfection of low but equal amounts of SynCAM 1 and ST8SiaII cDNA (see supplemental Fig. S3, right).

Under these optimized conditions, each SynCAM 1 isoform was then individually co-expressed with ST8SiaII (Fig. 4A). After immunoprecipitation of SynCAM 1 with anti-SynCAM 1 pAb, the polysialylation status and overall expression level of SynCAM 1 was compared by Western blot analysis using anti-polySia mAb 735 (Fig. 4A, top) and anti-SynCAM 1 mAb 3E1 (Fig. 4A, bottom), respectively. Notably, polysialylated Syn-
CAM 1 was detected for all SynCAM 1 isoforms, including isoform 8 lacking all variable peptides. Thus, ST8SiaII has the ability to polysialylate all transmembrane isoforms of SynCAM 1 independent of the presence or absence of variable peptides.

**ST8SiaII Acts More Efficiently on SynCAM 1 than ST8SiaIV**—The finding that polysialylation of SynCAM 1 relies exclusively on ST8SiaII in vivo (Fig. 1A) prompted us to compare the ability of ST8SiaII and ST8SiaIV to polysialylate this acceptor molecule in a cellular context. The different SynCAM 1 isoforms were now co-expressed with ST8SiaIV (Fig. 4B). After an exposure time of 1 min, polysialylation was only seen for isoforms 8bc and 8ac (Fig. 4B, top), and prolonged exposure times of ≈10 min were required to detect faint signals for all isoforms (data not shown). By contrast, co-expression with ST8SiaII resulted in strong polySia-SynCAM 1 signals that appeared already after an exposure time of 15 s (Fig. 4A). Moreover, analysis with an anti-SynCAM 1 antibody revealed that SynCAM 1 that was co-expressed with ST8SiaII showed reduced electrophoretic mobility compared with SynCAM 1 expressed in the presence of ST8SiaIV (Fig. 4, A and B, compare bottom panels). The observed shift in mobility indicates that ST8SiaII had converted the majority of SynCAM 1 to the polysialylated form, whereas ST8SiaIV had modified only a limited portion of the SynCAM 1 pool.

**FIGURE 3. Isoform pattern of polysialylated SynCAM 1.** A, analysis of polySia-SynCAM 1 immunoprecipitated (IP) from postnatal day 1 Ncam<sup>−/−</sup> mouse brain using the anti-polySia mAb 735. Immunoprecipitated polySia-SynCAM 1 was separated by 10% SDS-PAGE before and after the removal of polySia by endoN treatment or after de-N-glycosylation by PNGase F. Western blot analysis (WB) was performed with anti-SynCAM 1 antibodies against the extracellular part shared by all SynCAM 1 isoforms (mAb 3E1), or against the variable peptide encoded by exon 8c (mAb L45/30). The positions of the four protein bands obtained after de-N-glycosylation and staining with mAb 3E1 are indicated by arrows. Bands obtained after de-N-glycosylation that were stained by both mAb 3E1 and mAb L45/30 are marked by asterisks. B, schematic representation of anti-SynCAM 1 and anti-polySia antibody binding sites. mAb 3E1 is directed against the extracellular part of SynCAM 1 encompassing the Ig domains, mAb L45/30 detects specifically the variable peptide encoded by exon 8c, and the anti-SynCAM 1 pAb used in this study is directed against the C-terminal intracellular part of SynCAM 1. mAb 735 recognizes polySia composed of at least eight α2,8-linked sialic acids. C, assignment of SynCAM 1 isoforms that serve as polySia carriers in P1 mouse brain. PolySia-SynCAM 1 was immunoisolated from postnatal day 1 Ncam<sup>−/−</sup> mouse brain followed by de-N-glycosylation as described in A. In parallel, each of the indicated SynCAM 1 isoforms was transiently expressed in CHO cells, immunoprecipitated with the anti-SynCAM 1 pAb, and de-N-glycosylated by PNGase F. Proteins were separated by 8% SDS-PAGE and analyzed by Western blotting using anti-SynCAM 1 mAb 3E1. The four bands detected after de-N-glycosylation of mouse-derived polySia-SynCAM 1 in A, D, isoform pattern of polySia-SynCAM 1 at different time points of postnatal brain development. Using the anti-polySia mAb 735, polySia-SynCAM 1 was immunoisolated from Ncam<sup>−/−</sup> mouse brain obtained at the indicated time points. The immunoprecipitated material was de-N-glycosylated and analyzed by Western blotting using anti-SynCAM 1 mAb 3E1. E, schematic representation of the four SynCAM 1 isoforms that serve as targets for polysialylation in postnatal mouse brain.
To visualize these differences on the same blot, SynCAM 1 of the isoform 8bc was expressed either alone or together with ST8SiaII or ST8SiaIV (Fig. 5). Although similar amounts of SynCAM 1 were expressed and immunoprecipitated in all three settings (Fig. 5, bottom and middle, respectively), clear differences in the extent of polysialylation were observed. Although co-expression of ST8SiaII resulted in a strong polySia-SynCAM 1 signal, only tiny amounts of polysialylated SynCAM 1 were detected in the presence of ST8SiaIV (Fig. 5, top). Analysis of the total fraction of SynCAM 1 (Fig. 5, bottom) revealed that co-expression with ST8SiaII but not ST8SiaIV resulted in a shift in electrophoretic mobility. Together, these experiments provide the first evidence that ST8SiaII and ST8SiaIV are distinct in their ability to use SynCAM 1 as an acceptor substrate.

**DISCUSSION**

In contrast to most other glycosyltransferases, which modify glycan structures irrespective of the underlying protein, the two polySTs ST8SiaII and ST8SiaIV are highly protein-specific and
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The capacity to synthesize polySia of comparable quality on NCAM (26, 47).

In contrast to NCAM polysialylation, the present study demonstrates that throughout postnatal brain development, polysialylation of SynCAM 1 is catalyzed exclusively by ST8SiaII. Loss of ST8SiaII completely abolished SynCAM 1 polysialylation, whereas no significant changes were observed in the absence of ST8SiaIV. Thus, our data highlight a unique role of ST8SiaII in polysialylation of SynCAM 1. During postnatal brain development, prominent signals for polySia-SynCAM 1 were observed between P1 and P11, whereas polysialylation of SynCAM 1 was drastically down-regulated thereafter, resulting in a sharp drop between P11 and P15. This time course followed the previously described down-regulation of ST8Sial transcript level, which dropped by 95% during postnatal brain development, with the strongest reduction between P5 and P11 (27).

Previously, we have shown that in vitro, both polySTs, ST8SiaII and ST8SiaIV, were able to polysialylate SynCAM 1. These in vitro polysialylation experiments were performed with truncated, soluble proteins that lack their transmembrane domains. In vivo, however, polySTs and SynCAM 1 are transmembrane proteins that are fixed to the Golgi membrane during the polysialylation step. The observation that ST8SiaIV cannot compensate for ST8SiaII deficiency in vivo suggests that SynCAM 1 is a poor substrate for ST8SiaIV under in vivo conditions. This was further substantiated by our co-expression experiments, demonstrating that in the cellular context, ST8SiaII is much more efficient in polysialylating SynCAM 1 than ST8SiaIV. Although ST8SiaIV was able to recognize SynCAM 1 as an acceptor molecule, the enzyme modified only a limited portion of the SynCAM 1 pool, resulting in only minute amounts of polySia-SynCAM 1. By contrast, ST8SiaII clearly outperformed ST8SiaIV in its ability to modify almost the complete pool of SynCAM 1.

Alternative splicing, representing a versatile mechanism for generating a large pool of functionally unique proteins, is particularly frequent in the nervous system (48, 49). In the case of SynCAM 1, alternative splicing of the short variable exons 8a, 8b, and 8c can, theoretically, give rise to eight transmembrane isoforms of SynCAM 1. Here, we show that all of them with the single exception of isoform 8ac were expressed during postnatal mouse brain development. So far, exon 8c has been regarded as cryptic exon in mice (8). However, based on RT-PCR and sequencing of the obtained products, we demonstrated the occurrence of three exon 8c-containing transcript variants (i.e., 8abc, 8bc, and 8c), which were not identified in similar approaches performed previously to study developmental changes in the SynCAM 1 transcript pattern in the mouse brain (10, 11).

Notably, the variable peptides encoded by exon 8a, 8b, and 8c contain 17, 4, and 2 putative O-glycosylation sites, respectively, resulting in numerous glycoforms. Stretches of O-glycosylation sites are known to influence protein conformation, leading to an elongated overall structure, as described for P-selectin glycoprotein ligand-1 and ovine submaxillary mucin (50, 51). Accordingly, a highly O-glycosylated stem region in SynCAM 1 was suggested to position the Ig-like domains away from the plasma membrane, thereby regulating their accessibility for trans interactions (8). Moreover, O-glycosylation may affect cis
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Oligomerization of SynCAM 1, which was shown to precede synapse assembly and to promote trans adhesion (5).

On the transcript level, we observed the most striking developmental changes for isoform 8a, which lacks an O-glycosylated stem region. Although expressed at a low level during the first postnatal week, transcript variant 8a became the most prevalent variant in the adult brain (Fig. 2B). This suggests that SynCAM 1 isoforms with high numbers of potential O-glycosylation sites may dominate in early brain development, whereas isoform 8a, lacking any O-glycosylation sites, becomes the major isoform in the adult stage. These developmental changes might regulate the distinct functions of SynCAM 1 in the developing and mature brain (4–6, 12).

Precise knowledge on the SynCAM 1 isoform pattern and recombinant expression of individual isoforms enabled us to identify the isoforms underlying polysia-SynCAM 1 as 8abc, 8ab, 8bc, and 8b. Although polysialylation of SynCAM 1 was restricted to four particular isoforms in vitro, our data obtained by cell culture experiments demonstrated that ST8SialII is able to polysialylate all transmembrane isoforms of SynCAM 1 in vitro. Thus, the four isoforms that serve as targets for polysialylation in vitro represent most likely the SynCAM 1 isoforms expressed in NG2 cells, the cells in which polysialylation of SynCAM 1 occurs. In vivo, polysialylation of SynCAM 1 is restricted to these cells, suggesting that they provide the particular environment required for SynCAM 1 polysialylation. The key prerequisite might be a critical balance between SynCAM 1 and ST8SialII. High expression levels of the acceptor SynCAM 1 prevented its efficient polysialylation by ST8SialII, as demonstrated by our co-expression experiments. SynCAM 1 has been shown to undergo excessive lateral self-assembly, a process that precedes the engagement of SynCAM 1 in trans interactions and is mediated by the membrane proximal Ig domains 2 and 3 (5). Thus, elevated SynCAM 1 levels may favor cis oligomerization, and the resulting clustering of SynCAM 1 in the membrane might reduce its accessibility for ST8SialII.

Previously, we demonstrated that polysia is selectively attached to SynCAM 1 via N-glycans at Asn-116 located in the first Ig-like domain (13). By analysis of domain deletion variants of soluble SynCAM 1 variants, we now established the minimal structural requirements for SynCAM 1 polysialylation. Although Ig1 harbors the polysialylation site, this domain alone did not serve as an acceptor substrate for ST8SialII. However, a variant encompassing Ig1 and Ig2 was efficiently polysialylated, and no significant differences in polysialylation were observed compared with a variant containing all three Ig-like domains. These findings indicate that Ig2 provides the critical interface for the recognition of SynCAM 1 by ST8SialII.

As summarized in Fig. 7, the minimal structural requirements for polysialylation of SynCAM 1 are highly reminiscent of what was observed for NCAM. The extracellular part of NCAM is composed of five Ig-like domains followed by two fibronectin type III repeats (FnIII-1 and -2) (52). Among the six N-glycosylation sites of NCAM, only two particular sites located in Ig5 are used for polysialylation (53–55). Nevertheless, Ig5 alone did not serve as an acceptor substrate for the polysiTs, whereas a truncated NCAM variant encompassing the tandem domains Ig5 and FnIII-1 was efficiently polysialylated (56). The importance of the FnIII-1 domain as a recognition site was further substantiated by polyst-NCAM interaction studies (57). Subsequent structural analysis of FnIII-1 revealed the presence of an α-helix, an unusual feature among FnIII repeats, and an acidic surface patch essentially required for NCAM polysialylation (58, 59).

For SynCAM 1, structural information is so far only available for the C-terminal intracellular domain (60). Future work focused on the structural analysis of the extracellular part will help to identify structural features in Ig2 that provide the docking sites for ST8SialII and define SynCAM 1 as a target for polysialylation.

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