Motor dysfunction in cerebellar Purkinje cell-specific vesicular GABA transporter knockout mice

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The L7-VGAT mice are a novel model of ataxia without PC degeneration, and would also be useful for studying the role of PCs in cognition and emotion. Keywords: cerebellum, Purkinje cells, VGAT, knockout mice, ataxia, mouse model

INTRODUCTION

γ-Aminobutyric acid (GABA) is a major inhibitory neurotransmitter in the adult mammalian central nervous system and plays modulatory roles in neural development. The vesicular GABA transporter (VGAT) is an essential molecule for GABAergic neurotransmission due to its role in vesicular GABA release. Cerebellar Purkinje cells (PCs) are GABAergic projection neurons that are indispensable for cerebellar function. To elucidate the significance of VGAT in cerebellar PCs, we generated and characterized PC-specific VGAT knockout (L7-VGAT) mice. VGAT mRNAs and proteins were specifically absent in the 40-week-old L7-VGAT PCs. The morphological characteristics, such as lamination and foliation of the cerebellar cortex, of the L7-VGAT mice were similar to those of the control littermate mice. Moreover, the protein expression levels and patterns of pre- (calbindin and parvalbumin) and postsynaptic (GABA-A receptor α1 subunit and gephyrin) molecules between the L7-VGAT and control mice were similar in the deep cerebellar nuclei that receive PC projections. However, the L7-VGAT mice performed poorly in the accelerating rotarod test and displayed ataxic gait in the footprint test. The L7-VGAT mice also exhibited severer ataxia as VGAT deficits progressed. These results suggest that VGAT in cerebellar PCs is not essential for the rough maintenance of cerebellar structure, but does play an important role in motor coordination. The L7-VGAT mice are a novel model of ataxia without PC degeneration, and would also be useful for studying the role of PCs in cognition and emotion.

Abbreviations: BSA, bovine serum albumin; CB, calbindin; CNS, central nervous system; DCM, deep cerebellar nuclei; DCN, deep cerebellar nuclei; DIG, digoxigenin; ECL, enhanced chemiluminescence; EDTA, ethylenediaminetetraacetic acid; GABA, γ-aminobutyric acid; GABA_B receptor α1 subunit; GAD, glutamate decarboxylase; GAD_3, GABA transporters X; GABA_A receptor α2 subunit; IPSC, inhibitory postsynaptic current; ISH, in situ hybridization; KO, knockout; PB, phosphate buffer; PBSTx, Triton X-100 in PB; PC, Purkinje cell; PFA, paraformaldehyde; PMSE, phosphomolybdofluorsilicic acid; PV, parvalbumin; P40W, postnatal 40 weeks of age; SGH, sodium-glucose cotransporter 1; TBO, thionin; VN, vestibular nuclei; VGAT, vesicular GABA transporter; VN, vestibular nuclei; y-Aminobutyric acid (GABA) is a major inhibitory neurotransmitter in the adult mammalian central nervous system and plays modulatory roles in neural development. The vesicular GABA transporter (VGAT) is an essential molecule for GABAergic neurotransmission due to its role in vesicular GABA release. Cerebellar Purkinje cells (PCs) are GABAergic projection neurons that are indispensable for cerebellar function. To elucidate the significance of VGAT in cerebellar PCs, we generated and characterized PC-specific VGAT knockout (L7-VGAT) mice. VGAT mRNAs and proteins were specifically absent in the 40-week-old L7-VGAT PCs. The morphological characteristics, such as lamination and foliation of the cerebellar cortex, of the L7-VGAT mice were similar to those of the control littermate mice. Moreover, the protein expression levels and patterns of pre- (calbindin and parvalbumin) and postsynaptic (GABA-A receptor α1 subunit and gephyrin) molecules between the L7-VGAT and control mice were similar in the deep cerebellar nuclei that receive PC projections. However, the L7-VGAT mice performed poorly in the accelerating rotarod test and displayed ataxic gait in the footprint test. The L7-VGAT mice also exhibited severer ataxia as VGAT deficits progressed. These results suggest that VGAT in cerebellar PCs is not essential for the rough maintenance of cerebellar structure, but does play an important role in motor coordination. The L7-VGAT mice are a novel model of ataxia without PC degeneration, and would also be useful for studying the role of PCs in cognition and emotion.

Thus, VGAT is crucial for the vesicular GABA release, and VGAT deficits cause the nearly complete disappearance of inhibitory postsynaptic currents (IPSCs) in dissociated cultures or slices derived from VGAT knockout (KO) mice (Wosiek et al., 2006; Saito et al., 2010). However, because VGAT KO mice die during the perinatal period, the study of VGAT function in the adult CNS has been limited (Tong et al., 2009). The cerebellum is important for motor function, cognition, and emotion (Strata et al., 2011; Lisberger and Thach, 2013). Cerebellar Purkinje cell neurons (PCs) are the sole output neurons of the cerebellar cortex and project to the deep cerebellar nuclei (DCN) and vestibular nuclei (VN) neurons. The DCN and VN are involved in motor control in animals through their communication with the nuclei of the thalamus and brainstem. Because PCs are GABAergic neurons, it is thought that the existence of VGAT in PCs is critical for PC function. However, the significance of VGAT in PCs has not been demonstrated.

Here, to study the roles of VGAT in PCs, we generated cerebellar PC-specific VGAT KO (L7-VGAT) mice and examined the...
We generated heterozygous mice carrying one floxneo allele (VGAT\textsuperscript{floxneo/+}) in which the 5′-loxP site was introduced into the Xhd1 site in intron 1 and an frt-flanked PGK-neo cassette followed by the 3′-loxP sequence was inserted into the Kpdn site in the 3′-flanking region (Ebihara et al., 2003; Saito et al., 2010). The VGAT\textsuperscript{floxneo/+} mice were mated to FLPe transgenic mice (Takeda et al., 2005), and the male offspring were further crossed with C57BL/6 mice to eliminate the PGK-neo gene from the genome through Flpter-mediated excision. The resultant VGAT\textsuperscript{flox/+} mice were intercrossed to generate VGAT\textsuperscript{flox/flox} mice in which exons 2 and 3 of the VGAT gene were flanked by loxP sites. The generation of L7-Cre knock-in mice has been previously described (Saito et al., 2005). In these mice, the Cre recombinase gene was flanked by a PGK-neo cassette and was inserted into exon 2 of the L7/Pcp-2 (L7) gene, and the Cre recombinase is expressed under the control of the endogenous L7 promoter. VGAT\textsuperscript{flox/flx} mice were crossed with heterozygous L7-Cre knock-in mice (L7\textsuperscript{flox/+} mice) to obtain VGAT\textsuperscript{flox/flx}, L7\textsuperscript{flox/+} mice. VGAT\textsuperscript{tet/flx} L7\textsuperscript{flox/+} mice were mated to VGAT\textsuperscript{flox/flx}, L7\textsuperscript{flox/+} mice to obtain VGAT\textsuperscript{flox/flx}, L7\textsuperscript{flox/+} mice and VGAT\textsuperscript{flox/flx}, L7\textsuperscript{flox/+} mice, which were used in the subsequent experiments. We refer to the VGAT\textsuperscript{flox/flx}, L7\textsuperscript{flox/+} mice and the VGAT\textsuperscript{flox/flx}, L7\textsuperscript{flox/+} mice as L7-VGAT mice and control mice, respectively, hereafter.

EVALUATION OF VGAT KNOCKOUT IN PCs

The presence of VGAT in the L7-VGAT PCs was determined by the detection of in situ hybridization (ISH) signals for both VGAT and CB. The percentage of VGAT mRNA-positive PCs was determined in lobule III, because lobule III was uniformly stained with our ISH protocol. We analyzed two sections per animal and used two animals per genotype at 2, 4, 8, 16, and 40 weeks of age (postnatal). The number of VGAT mRNA-positive PCs was divided by the total number of PCs. Because CB is a marker of PCs (Celio, 1990), CB-positive cells were regarded as PCs.

IMMUNOHISTOCHEMISTRY

The primary antibodies used in this study included rabbit anti-VGAT (1:500, Takayama and Inoue, 2004a), mouse anti-CB (1:1,000, Swant, Bellinzona, Switzerland), mouse anti-parvalbumin (PV; 1:5,000; Sigma-Aldrich, Saint Louis, MO, USA), guinea pig anti-GABA receptor α1 subunit (GABA\textsubscript{A}Rα1;
Whole cerebella of 40-week-old control and L7-VGA T mice under deep anesthesia with diethyl ether. Brains were harvested perfused with 4% PFA in 0.1 M phosphate buffer, pH 7.4 (PB), Molecular Probes/Invitrogen).

To detect VGAT, CB, and P40W, mice were transcardially perfused with 4% PFA in 0.1 M phosphate buffer, pH 7.4 (PB), under deep anesthesia with diethyl ether. Brains were harvested from the perfused animals, post-fixed in the identical fixative for 12 h at 4 °C, and immersed in 30% sucrose in 0.1 M PB and 0.05% NaN3 for at least 48 h for cryoprotection. The frozen brains were sliced at 20 μm using a cryostat, then mounted onto slides, and dried for more than 30 min at room temperature. Immunostaining for VGAT and CB was performed following a previously published protocol (Takayama and Inoue, 2004a). Alexa568- or Alexa488-conjugated secondary antibodies were used to detect the primary antibodies. To detect P40W, the sections were blocked with 3% bovine serum albumin (BSA) in 0.3% Triton X-100 in PBS (PBSTx) for 1 h and then incubated with an anti-P40W antibody at 4 °C overnight. The primary antibody was detected with an Alexa Fluor 568 anti-rabbit IgG. To detect the colocalization of GABAAR and gephyrin, double immunofluorescence staining was performed in sections prepared from fresh-frozen tissue as previously described for the ISH. Sections were fixed in PB containing 0.5% PFA for 10 min, blocked with 3% normal goat serum in PB for 1 h, and then incubated overnight at 4 °C with anti-GABAAR, anti-gephyrin antibodies. The primary antibodies were detected using an anti-guinea pig Alexa Fluor 488 and anti-mouse Alexa Fluor 568-conjugated secondary antibodies. Fluorescence signals were obtained using an optical microscope (AxioCam, Carl Zeiss, Jena, Germany). The composite images were prepared from the digital data files using Adobe Photoshop and Illustrator.

WESTERN BLOT ANALYSIS

Whole cerebella of 40-week-old control and L7-VGAT mice were coronally cut at the most rostral level of the DCNs, and both sides of the DCNs were separated from the sectioned cerebella by cutting between the DCNs and the cerebellar cortex in ice-cold PBS. Both sides of the DCNs were separated from the sections prepared from fresh-frozen tissue as previously described for the ISH. Sections were fixed in PB containing 0.5% PFA for 10 min, blocked with 3% normal goat serum in PB for 1 h, and then incubated overnight at 4 °C with anti-GABAAR, anti-gephyrin antibodies. The primary antibodies were detected using an anti-guinea pig Alexa Fluor 488 and anti-mouse Alexa Fluor 568-conjugated secondary antibodies. Fluorescence signals were obtained using an optical microscope (AxioCam, Carl Zeiss, Jena, Germany). The composite images were prepared from the digital data files using Adobe Photoshop and Illustrator.

RESULTS

Generation of L7-VGAT mice
To investigate the role of VGAT in cerebellar PCs on cerebellar structure and motor function, we generated cerebellar PC-specific VGAT KO mice. We used floxed VGAT mice in which exons 2 and 3 of the VGAT gene are flanked by loxP sequences (Saito et al., 2010) and L7-Cre knock-in mice that express the Cre recombinase gene under the control of the intrinsic L7 promoter (Satoo et al., 2005). (Figures 1A,B). We crossed VGATflox/flox, L7-Cre/+ mice with VGATflox/flox, L7-Cre/+ mice to obtain VGATflox/flox, L7+/+ and VGATflox/flox, L7Cre/+ mice (Figure 1C). We refer to the Acknowledgments for details.
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FIGURE 1 | L7/Pcp-2 promoter-mediated disruption of the VGAT gene. (A) Schema of the wild-type and floxed VGAT allele (upper). Black boxes and white triangles indicate exons and loxP sites in the VGAT gene, respectively. Black arrows indicate the primer sites for PCR genotyping. Schema of cerebellar Purkinje cell (PC)-specific VGAT gene disruption in L7-VGAT mice (lower). (B) Schema of the wild-type and Cre knock-in L7/Pcp-2 (L7) allele. Black boxes indicate exons in the L7 gene. White and gray boxes indicate the Cre gene and the PGK-Neo cassette, respectively, that were introduced into exon 2 of the L7 gene. The black arrows indicate the primer sites for PCR genotyping. (C) PCR genotyping of littermates (numbers 1–8) obtained by crossing a VGAT flox/flox; L7 Cre/+ mouse with a VGAT flox/flox; L7 +/+ mouse. Upper and lower panels show the genotyping of the VGAT and L7 gene, respectively. Lanes 1–8 in both panels correspond to the identical littermate number. Lanes 9 and 10 in both panels were used as controls. Lanes 9 and 10 in the upper panel indicate the VGAT flox/+ and VGAT flox/flox genotypes, respectively. Lanes 9 and 10 in the lower panel indicate the L7 Cre/Cre and L7 +/+ genotypes, respectively.

to VGAT flox/flox; L7 +/+ and VGAT flox/flox; L7 Cre/+ as the control and L7-VGAT mice, respectively, hereafter. In the L7-Cre knock-in mice, Cre recombinase is specifically expressed in the cerebellar PCs and retinal cells including the bipolar neurons and photoreceptor cells (Saito et al., 2005). On the other hand, VGAT is expressed in PCs, but not in these retinal cells. Thus, in the L7-VGAT mice, VGAT was expected to be disrupted only in the PCs. The L7-VGAT mice were born and grew until 16 postnatal weeks (P16W) without any apparent abnormal phenotype. At P40W, the L7-VGAT mice were fertile and normal in terms of nursing, but they displayed abnormal ambulation. These mice walked awkwardly and easily fell off the edge of a table. The L7-VGAT mice survived until they were more than 1 year of age.

PC-SPECIFIC AND AGE-DEPENDENT DISAPPEARANCE OF VGAT mRNA IN L7-VGAT MICE

To assess whether VGAT mRNA was specifically lost in the PCs of the L7-VGAT mice, we performed double ISH for VGAT and calbindin (CB) (Figure 2). CB was used as a marker of PCs because CB is specifically expressed in PCs in the cerebellar cortex and DCNs (Celio, 1990). In the control cerebellum, hybridization signals for VGAT mRNA were lined up along the PC layer and completely overlapped with the CB signals (Figures 2Aa–f). In contrast, no cells containing CB mRNA expressed VGAT mRNA in the L7-VGAT cerebellum at P40W (Figures 2Ag–l). However, other VGAT mRNA signals were found in the molecular and granule cell layers of the P40W L7-VGAT mice. Furthermore, VGAT mRNA signals in brain regions outside of the cerebellum did not appear to be different between coronal sections of L7-VGAT and control mice at either P16W or P40W (data not shown).

Next, to assess the time at which VGAT mRNA was lost in the L7-VGAT PCs during development, we performed double ISH for VGAT and CB and counted the number of VGAT-positive PCs relative to the total number of PCs at five postnatal time points (i.e., P2W, P4W, P8W, P16W, and P40W; Figure 2B). The percentage of VGAT-positive PCs relative to the total PCs gradually decreased with age from 56% at P2W to 0% at P40W (Figure 2B).

LOSS OF VGAT PROTEIN IN L7-VGAT PCs

To confirm whether VGAT protein was also lost, we performed immunohistochemistry for VGAT and CB in the L7-VGAT PCs (Figure 3). CB is a PC marker protein that is located in the soma, dendrites, axonal fibers, and axon terminals of PCs (Bausle et al., 1997). CB-immunoreactive signals were present in the white matter, the DCNs, and the cerebellar cortices of both control and L7-VGAT mice (Figures 3A, DG). VGAT-immunoreactive
FIGURE 2 | Disappearance of VGAT mRNA specifically in the Purkinje cells of L7-VGAT mice. (A) Double in situ hybridizations for calbindin D-28K (a,d,g,j; green) and VGAT (b,e,h,k; red) were performed in parasagittal sections of 40-week-old mice. The cerebella of control (a–f) and L7-VGAT (g–l) mice are shown. (a–c) and (g–i) show images of the whole cerebella. (d–f) and (j–l) show images of lobule III. Merged images of (a,d,g,j) and (b,e,h,k) are shown in (c,f,i,l). Bars represent 500 μm (c) and 50 μm (f).

(B) The percentage of VGAT mRNA-positive Purkinje cells (PCs) in the L7-VGAT mice at 2, 4, 8, 16, and 40 weeks postnatal. The number of VGAT mRNA-positive PCs in the L7-VGAT mice was divided by the total number of calbindin mRNA-positive PCs in lobule III (L7-VGAT, n = 2; control, n = 2 of each age). The total numbers of calbindin-positive cells in the L7-VGAT mice that were counted were 230 at 2 weeks postnatal (P2W), 264 at P4W, 243 at P8W, 232 at P16W, and 200 at P40W. Each value represents the mean ± SEM. Gra, granule cell layer; Pur, Purkinje cell layer; Mol, molecular layer; and Wm, white matter.

signals overlapped with CB signals in the DCNs of the control mice, which suggests that the VGAT signals were derived from PCs. The DCN neurons appeared to be heavily innervated by GABAergic terminals of the PCs (Figure 3F). In the L7-VGAT DCNs at P40W, a small number of VGAT-positive and CB-negative signals remained. These results are consistent with a previous report that a large portion of inhibitory synapses on DCN neurons is derived from PCs (Garin et al., 2002).

To verify our immunohistochemical results, we performed Western blot analyses (Figure 4; Table 1). To acquire DCN tissue for Western blot analyses, we focused on GA T3. GA T3 is one of the GABA transporters, which uptake extracellular GABA, and is located in the glial cell membrane (Itouji et al., 1996). It has been reported that the DCNs exhibit intense GA T3 immunoreactivity and that this immunoreactivity is faint in the cerebellar cortex (Itouji et al., 1996). Therefore, we used GA T3 as a marker of the DCN region. We dissected the cerebellum into the putative DCN and cerebellar cortex regions (see section “Materials and Methods”). Although GA T3 signals were detected in the crude fractions of control and L7-VGAT DCNs, they were not detected in the fractions of the control or L7-VGAT cerebellar cortices. On
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FIGURE 4 | Significant reduction in VGAT protein and unaltered levels of other pre- and postsynaptic proteins in the L7-VGAT DCN. Western blot analyses of GAT3, β-actin, calbindin, gephyrin, GABA_A receptor α1, and parvalbumin proteins in the crude fractions from control and L7-VGAT DCN and cerebellar cortices at P40W. DCN, deep cerebellar nuclei; and CC, cerebellar cortex.

Table 1 | Relative signal intensities of protein immunoreactivities.

| Protein       | DCN (Control) | L7-VGAT (Control) | L7-VGAT (L7-VGAT) | p-value |
|---------------|---------------|-------------------|-------------------|---------|
| VGAT          | 100 ± 18      | 40 ± 6*           | 70 ± 14*          | p < 0.001 |
| Calbindin     | 100 ± 10      | 121 ± 19          |                   |         |
| Gephyrin      | 100 ± 13      | 96 ± 29           |                   |         |
| GABA_A receptor α1 | 100 ± 15      | 107 ± 12          |                   |         |
| Parvalbumin   | 100 ± 15      | 114 ± 18          |                   |         |

The signal intensities of each protein were normalized by those of β-actin. Values represent mean ± SD (n = 4). *p < 0.05, **p < 0.001 (Student’s t-test).

ROUGHLY MAINTAINED STRUCTURE OF THE L7-VGAT CEREBELLUM

It is known that GABA can be involved in neural development (Owens and Kriegstein, 2002; Represa and Ben-Ari, 2005). Thus, it is possible that the anatomical structure of the L7-VGAT cerebellum was altered. To examine whether the L7-VGAT cerebellum was anatomically affected, we performed cresyl violet staining (Figure 5). We did not detect any overt abnormalities in the lamination or foliation of the 40-week-old L7-VGAT cerebellum (Figures 5A,B). The sizes, densities, and staining intensities of the cells in each layer of the L7-VGAT cerebellar cortex were similar to those in control cerebellum (Figures 5C,D). The overall size of DCN appeared to remain unaltered (Figures 5E,F). Additionally, there were no significant differences in the thicknesses of the molecular or granule cell layers or in the densities of the PCs between the L7-VGAT and control cerebellar cortices (Table 2). These results suggest that the loss of VGAT in the PCs did not significantly interfere with the structure of the cerebellum.

Table 2 | Anatomical comparison between control and L7-VGAT cerebella at postnatal 40 weeks.

| Measure                      | Control | L7-VGAT |
|------------------------------|---------|---------|
| Molecular layer thickness (μm) | 168.9 ± 5.6 | 174.6 ± 5.6 |
| Granule cell layer thickness (μm) | 109.7 ± 3.7 | 117.2 ± 1.3 |
| PC soma density (cells/mm)   | 19.2 ± 1.0 | 19.4 ± 1.1 |

All values represent mean ± SEM (n = 4). Student’s t-test was performed with genotype.
Next, we examined the immunohistochemical profiles of pre- and postsynaptic proteins at the PC-DCN synapse of the L7-VGAT cerebellum as an index of synapse formation (Figures 3 and 6). CB is a presynaptic protein at the PC-DCN synapse. CB immunoreactivity, including punctate signals around the somata, was largely unaltered in the L7-VGAT DCNs (Figures 3A,D,G,J). Moreover, PV, which is expressed in some neurons in the DCNs, was also localized to the PC terminals. The PV immunoreactive profiles were also similar between the control and L7-VGAT DCNs (Figures 6M,N). As representatives of postsynaptic proteins, we examined the immunoreactivities for the GABA-A receptor α1 subunit (GABAARα1), which is the most abundant subunit in adult brain including the cerebellum) and gephyrin, which is a scaffolding protein for the GABAAR and glycine receptor (Erisir et al., 1998; Ogris et al., 2006). These two proteins colocalized at the postsynaptic site including the somatic membrane in the control DCN (Figures 6A–F). Additionally, the expression patterns of GABAARα1 and gephyrin in the L7-VGAT DCNs were similar to those in the control DCNs (Figures 6G–L). We also performed Western blot analyses for CB, PV, GABAARα1, and gephyrin proteins in the crude fractions of the DCN to compare the expression levels of these proteins between the L7-VGAT and control mice (Figure 4, Table 1). The amounts of these four proteins were not significantly different between the genotypes. Collectively, the distribution patterns and expression levels of the representative pre- and postsynaptic proteins were not different between the L7-VGAT and control mice.

**MOTOR DISCOORDINATION IN THE L7-VGAT MICE**

Because the cerebellum is important for motor coordination, we performed accelerating rotational tests at P8W, P16W, and P40W (Figure 7A). The control mice improved their performances with trial repetition, and even the older mice (P40W) demonstrated an improvement after repeated trials. At P8W, no significant differences were observed in the retention time between the L7-VGAT and control mice (p > 0.05), which indicates that the 8-week-old L7-VGAT mice performed normally in the accelerating rotarod test. At P16W, there was a significant difference in the retention time between the L7-VGAT and control mice (p = 0.001). However, there was no significant difference in the strain X trial interaction (p > 0.05). At P40W, there was a significant difference in the retention time between the L7-VGAT and control mice (p < 0.0001). Moreover, there was also a significant strain X trial interaction (p < 0.0001), which demonstrates that the L7-VGAT mice displayed severer ataxia at P40W than at P16W. However, the 40-week-old L7-VGAT mice began to exhibit moderately improved performance over trials (p < 0.0001).

Next, we examined the footprint test (Figure 7B). There were no significant differences in either stride length or width between the L7-VGAT and control mice at P16W (Figure 7C). In contrast, at P40W, the stride lengths were similar, but the stride width of the L7-VGAT mice was significantly wider than that of the control mice (L7-VGAT, 4.05 ± 0.52 cm; control, 2.50 ± 0.23 cm; p < 0.05) (Figure 7C). These results support the conclusion that the L7-VGAT mice walked normally at P16W, but displayed abnormal ambulation at P40W.

Collectively, the results of the rotarod and footprint tests demonstrate that the L7-VGAT mice displayed severer motor discoordination at P40W than at P16W.

**DISCUSSION**

The L7-VGAT cerebellum does not exhibit apparent structural changes. It has been reported that GABA is involved in neuronal development including the proliferation, migration, and differentiation...
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FIGURE 7 | Age-dependent motor dysfunction in L7-VGAT mice. (A) Accelerating rotarod test for control (open circles) and L7-VGAT (filled circles) mice at postnatal weeks 8, 16, and 40. Latencies to fall from the rotating cylinder are plotted against each trial. The male mice were tested with three consecutive trials per day for three consecutive days. The numbers of animals were as follows: control, \( n = 14 \), L7-VGAT, \( n = 7 \) at P8W; control, \( n = 11 \), L7-VGAT, \( n = 9 \) at P16W; control, \( n = 10 \), L7-VGAT, \( n = 9 \) at P40W. Each value represents the mean ± SEM. (B) Typical hind paw footprints for the control and L7-VGAT mice at P40W. (C) Hind paw stride lengths and widths at postnatal weeks 16 and 40 for control and L7-VGAT mice. Each value represents the mean ± SEM. The numbers of animals were \( n = 5 \) in the control group (open bar) and \( n = 4 \) in the L7-VGAT (filled bar) group. The asterisk and N.S. represent \( p < 0.05 \) and \( p > 0.05 \), respectively (Student’s t-test).

of immature neurons (Owens and Kriegstein, 2002; Represa and Ben-Ari, 2005). However, no overt changes, including changes in laminination and foliation, were observed in the L7-VGAT cerebellum (Figure 5; Table 2). Moreover, clear alterations in the expression profiles of synaptic proteins, including CB, which is a marker protein for PCs in the cerebellum, were not observed (Figures 3, 4, and 6, Table 1), which suggests that there were no drastic disturbances in the PC-DCN synapse including in the axon terminals of the PCs. It has been reported that there are few structural abnormalities in mice with KO’s of GABAergic system-related molecules including GAD and VGAT (Li et al., 1999; Wojcik et al., 2006). The conventional glycine receptor \( \alpha 2 \) subunit (GlyR\( \alpha 2 \)) KO mice also showed no gross morphological abnormalities in the CNS including the retina (Young-Pearse et al., 2006). The acute knockdown of the GlyR\( \alpha 2 \) expression using RNAi electroporation into the retina, however, resulted in a decrease of the rod photoreceptor cell number (Young and Cepko, 2004). In conventional KO mice in which the target gene is inactivated in a fertilized egg, a compensatory mechanism(s) may ameliorate deficits due to the absence of the molecule. Although the L7-VGAT mouse is a conditional KO one, VGAT expression had already disappeared in approximately 40% of the PCs at 2 weeks postnatal in the L7-VGAT cerebellum, and the progress of the KO was very slow after P2W (Figure 2). Therefore, it is possible that developmental compensation occurred in the L7-VGAT cerebellum. On the other hand, there are reports that suggest that neurotransmitters that exert developmental effects are released in a non-vesicular fashion (Taylor and Gordon-Weeks, 1991; Flint et al., 1998; Owens and Kriegstein, 2002). Thus, the effect of VGAT deficit on neural development may be small. It has also been reported that mutation of the GAD67 gene causes defects in axon branching in the visual cortex during the third postnatal week, but not in the fourth postnatal week (Chatterpudayya et al., 2007). Therefore, we cannot exclude the possibility that a transient disturbance occurred during the period before P40W in the L7-VGAT cerebellum. Studies with GABA\( \alpha 2 \)R1 KO mice demonstrated that substantial inhibitory synapses such as the localization of neuroligin 2 were maintained on PC dendrites of KO ones, but that the density of inhibitory synapses were significantly reduced, suggesting that synapse number is regulated by an activity-dependent manner during the period of synaptogenesis (Patrizi et al., 2008). Because GABAergic postsynaptic currents were lost in the PCs of GABA\( \alpha 2 \)R1 KO mice (Patrizi et al., 2008), we cannot exclude the possibility that number of synapses from the PCs was reduced in the L7-VGAT DCN, in which GABAergic outputs from PCs should be lost. Alternatively, it is possible that some disturbances would be detected at the electron microscopic level. ATAXIA OBSERVED IN CEREBELLAR PC-SPECIFIC VGAT-DEFICIENT MICE L7-VGAT mice exhibited reduced retention times and widened gait in the accelerating rotarod and footprint tests, respectively (Figure 7). Moreover, these impairments became more obvious as the deficit in VGAT progressed, which indicates that VGAT expression in cerebellar PCs plays an important role in the motor coordination of animals.
The cerebellum plays an important role in motor function, and PCs are indispensable for cerebellar function. The importance of PCs has been demonstrated by analyses of spontaneous mouse mutants in which cerebellar PCs are degenerated (Lalonde and Strazielle, 2007). However, the GABAergic axons of the PC that form synapses on DCN neurons disappear as a result of PC degeneration in these spontaneous mouse mutants (Bäurle et al., 1997, 1998). GABAergic actions consist of so-called phasic inhibition by vesicularly released GABA and tonic inhibition via non-vesicularly released GABA (Semyanov et al., 2004; Fantuzzi et al., 2005). Moreover, several neuropeptidies, such as cholecystokinin, are expressed in PCs (Akiyama et al., 2008). Every signaling system of the PC terminals is abolished in the spontaneously PC-degenerating mice. Moreover, other types of neurons also degenerate because of the secondary effects of PC degeneration in the spontaneously PC-degenerating mouse cerebellum (Lalonde and Strazielle, 2007). Therefore, the phenotypes, including ataxia, that are observed in these spontaneously PC-degenerating mice are likely the result of many factors, which make it difficult to evaluate the significance of vesicular GABAergic neurotransmission in PCs. In contrast, neither obvious anatomical abnormalities across the entire cerebellum nor clear disturbances of the PC axon terminals on the DCNs were observed in the L7-VGAT mice (Figures 3-5, Tables 1 and 2). However, the L7-VGAT mice displayed ataxia, which suggests that vesicularly released GABA from the cerebellar PC terminal greatly contributed to PC function.

In the L7-VGAT mice, widened gait was not detected at P16W when 5% of the VGAT mRNA-positive PCs remained, but widened gait was observed at P40W when all of the cells had disappeared (Figures 2 and 7). Immunoreactive signals for both CB and VGAT in the L7-VGAT DCNs at P16W were hardly detectable as they were at P40W (data not shown). These results suggest that the VGAT loss in the L7-VGAT PCs occurred at both the mRNA and protein levels during the same period. This assertion further indicates that gait disturbance can be prevented by the presence of only 5% of the normal population of vesicular GABA-releasing PCs. This inference is not compatible with the report that ataxia appears in the spontaneously PC-degenerating mutants before the number of surviving PCs is reduced by half (Landis, 1973; Mullen et al., 1976); thus, the ataxia observed in the L7-VGAT mice is less severe than that of the spontaneously PC-degenerating mutants. These findings also suggest that the compensatory effects for the dysfunction of PCs are strongly exerted in the L7-VGAT CNS (see also the previous section).

**Unaltered Protein Expression Patterns and Levels of gephyrin and parvalbumin**

The proteins we examined as marker molecules that are present at the PC-DCN synapse include gephyrin and PV. Although the expression profiles of these proteins were not overtly altered in the L7-VGAT DCN (Figures 3, 4, and 6, Table 1), those in the DCN and VN of the spontaneously PC-degenerating mutants were altered. The number of puncta that were immunoreactive for gephyrin was reduced by approximately half (Garin et al., 2002), and PV-positive somata appeared (Bäurle et al., 1997, 1998).

Gephyrin clustering at the postsynaptic sites is induced by α-neurexin in the presynaptic spike (Kang et al., 2008). When α-neurexin is bound with neurogenin 2, gephyrin associates with neurogenin 2 and collybistin to form a complex and is involved in the clustering of inhibitory neurotransmitter receptors (Pouliopoulos et al., 2009). The PC axons are retracted from synaptic sites in PC-degenerating mutants, and α-neurexin in the PC axon terminals are far from the synaptic sites. Collectively, it is likely that the retraction of PC axons, rather than the loss of output from PCs, inhibits the formation of the neurogenin-neurogenin complex and affects gephyrin clustering at postsynaptic sites in PC-degenerating mutants.

PV is a Ca$^{2+}$-binding protein that buffers intracellular Ca$^{2+}$. It is thought that the concentration of intracellular PV reflects the activity level of the neuron (Kampfuis et al., 1988; Bäurle et al., 1997, 1998; Chaudhury et al., 2008). Although many DCN neurons express PV, PV-immunoreactivity is detected only when axonal transport is pharmacologically inhibited (Celio, 1990, Bäurle et al., 1998). Moreover, most PV-positive cells are inhibitory neurons in the DCN of PCD mutant mice (Bäurle et al., 1997). Moreover, Bäurle et al. (1998) asserted that the activity of inhibitory neurons in the DCN should increase to compensate for the reduced inhibitory input from the PCs to the DCN, which was accompanied by a change in PV expression in the spontaneous mutants. However, neither the appearance of PV-positive somata nor significant elevations of PV protein levels were observed in the L7-VGAT DCN (Figures 4 and 6, Table 1). In the L7-VGAT DCN, the loss of GABAergic inhibition from PCs is expected. Nevertheless, the elevated expression of PV did not occur, which implies that the reorganization of the somatic motor pathway that occurs in the L7-VGAT CNS differs from that in the spontaneous mutants.

**LVGAT Mice as a Model of the PC Dysfunctional Mutant**

The L7-VGAT mice generated in this study became completely PC-specific VGAT knockout mice and exhibited ataxia. It was expected that only vesicular release of GABA from the PCs would be disrupted in the L7-VGAT cerebellum and that PC degeneration and other drastic changes in cerebellar structure would not occur. The majority of mouse models of ataxia display histological phenomena, such as losses of PCs, which explain their cerebellar dysfunctions. Only a few mouse models exist that display ataxic behavior combined with normal cerebellar morphology (Hendriks et al., 2009). Additionally, the cerebellum has been reported to be critical also for cognition and emotion (Schmahmann, 2010). Therefore, the L7-VGAT mice will greatly contribute to studies of cerebellar functions from the cellular level (e.g., neuronal connectivity) to the whole-body level (e.g., motor coordination, vestibular compensation, cognition, and emotion).

The VGAT deficit was complete in the PCs at P40W (Figures 2 and 3). Because the L7-Cre gene in the L7-VGAT mice was hermaphroditic, the VGAT-flox gene would be expected to be deleted much earlier in the animals that are homozygous for the L7-Cre gene. The use of these two lines of L7-VGAT mice may be useful in the study of the differences in the effect of VGAT deficits that occur at different ages.
CONCLUSION

Studies using L7-VGAT mice demonstrated that VGAT in cerebellar PCs was not essential for the maintenance of overall cerebellar structure or the expression profiles of synaptic proteins including calbindin, parvalbumin, GABA\(_B\)R1, and gephyrin, but that it was crucial for the motor coordination in animals. The L7-VGAT mice will be a useful model to study due to the better understanding of PC function in diverse cerebellum-related behaviors.

AUTHOR CONTRIBUTIONS

Mikiko Kayakabe, Toshikazu Kikuzaki, Ryouuke Kaneko, Yoichi Nakazato, Koichi Shibasaki, Yauki Ishizaki, Nobuhiko Furuya, and Yucio Yanagawa: Wrote the paper. All authors read and approved.

Yoko Yamagata for critically reading the manuscript and provid-

Atsushi Sasaki, Koji Shibasaki, and Yucio Yanagawa: Performed the experi-

ments; Mikiko Kayakabe, Toshikazu Kikuzaki, Ryouuke Kaneko, Atsushi Sasaki, Koichi Shibasaki, and Yucio Yanagawa: Analyzed the data; Yoichi Kaneko, Atsushi Sasaki, Yoichi Nakazato, Hirotsuru Saito, and Noboru Suzuki: Contributed new reagents/analytical tools; Mikiko Kayakabe, Toshikazu Kikuzaki, Ryouuke Kaneko, and Yucio Yanagawa: Wrote the paper. All authors read and approved the final manuscript.

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