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Distinct Neurochemical and Functional Properties of GAD67-Containing 5-HT Neurons in the Rat Dorsal Raphe Nucleus

Hiroki Shikanai, Takayuki Yoshida, Kohtarou Konno, Miwako Yamasaki, Takeshi Izumi, Yu Ohmura, Masahiko Watanabe, Mitsuhiko Yoshioka

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The serotonergic (5-HTergic) system arising from the dorsal raphe nucleus (DRN) is implicated in various physiological and behavioral processes, including stress responses. The DRN is comprised of several subnuclei, serving specific functions with distinct afferent and efferent connections. Furthermore, subsets of 5-HTergic neurons are known to coexpress other transmitters, including GABA, glutamate, or neuropeptides, thereby generating further heterogeneity. However, despite the growing evidence for functional variations among DRN subnuclei, relatively little is known about how they map onto neurochemical diversity of 5-HTergic neurons. In the present study, we characterized functional properties of GAD67-expressing 5-HTergic neurons (5-HT/GAD67 neurons) in the rat DRN, and compared with those of neurons expressing 5-HTergic molecules (5-HT neurons) or GAD67 alone. While 5-HT/GAD67 neurons were absent in the dorsomedial (DRD) or ventromedial (DRV) parts of the DRN, they were selectively distributed in the lateral wing of the DRN (DRL), constituting 12% of the total DRL neurons. They expressed plasmalemmal GABA transporter 1, but lacked vesicular inhibitory amino acid transporter. By using whole-cell patch-clamp recording, we found that 5-HT/GAD67 neurons had lower input resistance and firing frequency than 5-HT neurons. As revealed by c-Fos immunohistochemistry, neurons in the DRL, particularly 5-HT/GAD67 neurons, showed higher responsiveness to exposure to an open field arena than those in the DRD and DRV. By contrast, exposure to contextual fear conditioning stress showed no such regional differences. These findings indicate that 5-HT/GAD67 neurons constitute a unique neuronal population with distinctive neurochemical and electrophysiological properties and high responsiveness to innocuous stressor.

Introduction

Serotonin (5-HT) exerts a tonic modulatory effect on a wide variety of physiological and behavioral processes (Lucki, 1998). 5-HT-producing neurons in the dorsal raphe nucleus (DRN) provide the majority of 5-HT innervations mainly to lateral forebrain areas, such as the basal ganglia, amygdala, and prefrontal cortex (Jacobs and Azmitia, 1992). Activation of 5-HTergic neurons in the DRN facilitates learned defensive behaviors (Zangrossi et al., 2001), while their inhibition by microinjection of GABA or benzodiazepine chlordiazepoxide exerts anxiolytic effects that attenuate suppression of conditioned punishing stimulus on lever-pressing behavior (Thiebot et al., 1980). Thus, 5-HT released from DRN neurons plays an important role in the regulation of anxiety responses.

Although the DRN is defined as a single nucleus, it is composed of subregions, which are clusters of neurons with distinct morphological features (Jacobs and Azmitia, 1992). They are organized into two main rostrocaudal subdivisions and further divided into six subnuclei: rostral, caudal, interfascicular, dorsomedial (DRD), ventromedial (DRV), and lateral “wing” (DRL) parts. They are topographically organized so that neurons in respective subnuclei innervate distinct targets and receive discrete afferent inputs (Jacobs and Azmitia, 1992). Importantly, these anatomically distinct subnuclei show differential responsiveness to various stressors (Hale and Lowry, 2011), suggesting that changes in neuronal activity of a limited subset of 5-HTergic neurons may have important implications in specific physiological and behavioral responses. Moreover, subpopulations of 5-HTergic neurons differently coexpress neuropeptides and transmitters, generating further complexity within topographically organized subnuclei (Jacobs and Azmitia, 1992; Michelsen et al., 2007). Accordingly, despite the growing evidence for rostral-caudal and dorsal-ventral variations in DRN responses, relatively little is known about correlation with unique neurochemical properties of 5-HTergic neurons.

The firing activity of 5-HTergic neurons in the DRN is controlled by two main mechanisms, i.e., autoregulatory influences arising from 5-HTergic neurons themselves and heteroregulation by local neurons and afferents, including GABAergic, glutamatergic, and dopaminergic inputs (Piñeiro and Blier, 1999). Of these, GABAergic and 5-HTergic transmissions can negatively regulate 5-HT release (Gallagher and Aghajanian, 1976; Piñeiro and Blier, 1999), and are thereby considered possible targets of...
therapeutic interventions. Interestingly, some 5-HTergic neurons have been shown to express GABA or its synthetic enzyme glutamic acid decarboxylase (GAD) (Nanopoulos et al., 1982; Belin et al., 1983; Fu et al., 2010; Hioki et al., 2010), suggesting a possible corelease of 5-HT and GABA. In this regard, these neurons may have a unique mechanism that controls their firing activity through autoregulatory feedback inhibition. However, functional properties of these neurons in the DRN remain largely unexplored.

In the present study, we characterized GAD67-expressing 5-HTergic neurons (5-HT/GAD67 neurons) in the DRN. We found that 5-HT/GAD67 neurons are localized to the DRL, constituting a unique neuronal population with distinct neurochemical and electrophysiological properties. Furthermore, these neurons show high responsiveness to innocuous stress compared with 5-HTergic neurons lacking GAD67 (5-HT neurons) and GAD67-expressing neurons lacking 5-HTergic markers (GAD67 neurons).

Materials and Methods

Animal. The first-breeder Wistar/ST rats were purchased from Nippon SLC and bred in our laboratory. Male rats at 3–4 weeks of age were used for analysis unless noted otherwise. Rats were group housed in a room with a 12 h light/dark cycle (with lights on at 7:00 P.M.) and a temperature-controlled environment (22 ± 1°C) with food and water ad libitum. All animals were treated in accordance with the guidelines for the Care and Use of Laboratory Animals of the Animal Research Committee of the Hokkaido University.

Drug. Diazepam (0.1 or 0.5 mg/kg; Wako Pure Chemical Industries) was suspended in 0.9% saline containing 1% arabic gum (Nacalai Tesque) and was injected intra peritoneally.

Immunohistochemistry. Rats were anesthetized with sodium pentobarbital (60 mg/kg, i.p.) and transcardially perfused via the left ventricle with 0.9% saline followed by 4% paraformaldehyde in 0.1M sodium phosphate buffer (PB), pH 7.2. After removal from the skull, brains were immersed overnight in the same fixative at 4°C, placed in 0.1M PB containing 10 –30% sucrose at 4°C, and sectioned at 30 μm thickness in a Leica HM 300 cryostat. Sections were collected in 0.01M PBS, pH 7.2, containing 0.1% Triton X-100 (PBSTx) and stored for at least 1 h before immunohistochemical staining. All immunohistochemical incubations were done at room temperature. For immunofluorescence, sections were incubated with a mixture of fluorophore-linked species-specific secondary antibodies, mouse anti-5-HT (1:500, 1 h) and the FITC-TSA Plus amplification kit (PerkinElmer), and the second detection was with peroxidase-conjugated anti-FITC antibody (Roche Diagnostics; 1:500, 1 h) and the Cy3-TSA Plus amplification kit (PerkinElmer). Residual activities of peroxidase introduced during the first detection were inactivated by incubation of sections with 0.6% H2O2 for 30 min. The specificity of FITC was confirmed by blank labeling using control sense cRNA probes. Images were taken with an FV1000 confocal laser-scanning microscope (Olympus).

Electrophysiological recording. Under deep anesthesia with diethyl ether, rats were decapitated, and coronal brain slices through the DRN (250 μm thick) were cut with a Leica VT1000S slicer in ice-cold low-Na+ solution with a specific composition (in mM): 215 sucrose, 2.5 KCl, 4 MgCl2, 4 MgSO4, 1 CaCl2, 26 NaHCO3, 1.6 NaH2PO4, and 20 glucose bubbling with 95% O2 and 5% CO2. For recovery, slices were incubated for 30 min in a mixed solution of 50% low-Na+ solution and 50% normal bathing solution containing the following (in mM): 125 NaCl, 2.5 KCl, 2 CaCl2, 1.25 NaH2PO4, 26 NaHCO3, and 20 glucose, pH 7.4, which was bubbled continuously with a mixture of 95% O2 and 5% CO2 at 37°C, followed by a mixed solution of 10% low-Na+ solution and 90% normal bathing solution at 37°C. Whole-cell patch-clamp recordings were made from DRN neurons in coronal acute slices using an upright microscope (BX51WI; Olympus) equipped with an infrared-CCD camera system (Hamamatsu Photonics) in normal bathing solution at 32°C. The resistance of the patch pipette was 3–6 MΩ when filled with the standard intracellular solution containing the following (in mM): 6 KCl, 130 K+glucuronate, 10 NaCl, 10 HEPES, 0.5 EGTA, 0.1 CaCl2, 2 MgCl2, 4 Na-ATP, and 0.4 Na-GTP, pH 7.3, adjusted with KOH. The lateral wing subdivision of the DRN (DRL) was selected as the recording region because the density of GAD67-positive 5-HT neurons was relatively higher than in ventral or dorsal parts of the DRN (Hioki et al., 2010). After whole-cell recording from DRN neurons held at a membrane potential of ~70 mV and switched to current-clamp recording mode, membrane potentials were recorded with an Axopatch 200B amplifier (Molecular Devices) and obtained by stepwise current injections (from ~0.2 to 0.5 nA; duration, 500 ms). Capacitance compensation and bridge-balance adjustment were made simultaneously. The pCLAMP 9 software (Molecular Devices) was used for stimulation and data acquisition. Signals were filtered at 3 kHz and digitized at 20 kHz. The liquid junction potential was ~10 mV between the pipette solution and the normal bath solution, which was subtracted from the recorded data.

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<th>Table 1. The primer sequences and estimated product lengths for single-cell RT-PCR</th>
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Fluorescent in situ hybridization. Procedures for double fluorescent in situ hybridization (FISH) labeling were reported previously (Yamasaki et al., 2010). To fresh frozen sections we applied ~300 ng/ml of fluorescein- or digoxigenin (DIG)-labeled riboprobes that were antisense to cDNAs of mouse GAD67 (nucleotide residues 1036–2015; GenBank accession number NM_000877), TPH2 (nucleotide residues 1–100; GenBank accession number NM_173391.3), and GAT-1 (nucleotide residues 380–2179; GenBank accession number NM_178703). DIG and fluorescein were detected by the two-step method: the first detection was with peroxidase-conjugated anti-fluorescein antibody (Roche Diagnostics; 1:500, 1 h) and the FITC-TSA Plus amplification kit (PerkinElmer), and the second detection was with peroxidase-conjugated anti-DIG antibody (Roche Diagnostics; 1:500, 1 h) and the Cy3-TSA Plus amplification kit (PerkinElmer). Residual activities of peroxidase introduced during the first detection were inactivated by incubation of sections with 0.6% H2O2 for 30 min. The specificity of FISH was confirmed by blank labeling using control sense cRNA probes. Images were taken with an FV1000 confocal laser-scanning microscope (Olympus).
Single-cell mRNA collection and reverse-transcription polymerase chain reaction. After whole-cell patch-clamp recording, the intracellular contents of each neuron were aspirated with negative pressure into the glass-electrode pipette that was used for recording and ejected into 10 μl RNase-free water (Promega). These samples (putatively containing mRNA) were immediately stored at −80°C. The mRNA was converted to cDNA with the QuantiTect Reverse Transcription Kit (Qiagen). For cDNA amplification, PCR was performed with the SYBR GreenER qPCR SuperMix for ABI PRISM (Invitrogen) in the ABI PRISM 7000 Sequence Detection System (Applied Biosystems). The primer sequences and estimated product lengths are listed in Table 1. PCR was conducted under the following conditions: 50°C for 2 min and 95°C for 10 min, followed by 40 cycles of 94°C for 30 s and 54°C for 30 s using a programmable thermal controller (PTC-100TM; MJ Research). PCR products were visualized using ethidium bromide staining after electrophoresis in a 3% agarose gel. Amplification of neuron-specific enolase cDNA confirmed that aspiration from the neuron was successful. PCR reactions without cDNA templates (containing only a pipette solution, bath solution, or noncellular debris) or with non-reverse transcription (RT) template were used as negative controls.

Open field stress. The open field (OF) stress test was used to induce mild and novel environmental stress on rats. The OF apparatus was a square chamber (90 cm long by 90 cm wide by 40 cm high). The central or peripheral zone was defined as a central square of 30 cm length by 30 cm width or a 10 cm width of inner circumference, respectively. To evaluate the locomotor activity, the traveled distance was recorded and automatically analyzed by the LimeLight 2 software package (Actimetrics), and the total number of crossings (the times animals crossed the lines dividing 9 × 9 squares in the open field) was also recorded for 30 min. Anxiolytic drug, diazepam, or vehicle (0.9% saline containing 1% arabic gum) was administrated intraperitoneally 30 min before exposing the OF apparatus. Tested rats and home cage controls were fixed for immunohistochemistry 2 h after OF stress.

Contextual fear conditioning stress. The contextual fear conditioning (CFC) stress test was used to induce severe and memory-dependent stress on rats. The detailed procedures were described previously (Ohmura et al., 2008, 2010). To quantify fear responses, electrical footshocks were administered five times (2 s; 0.3 mA; 30 s intervals) in a chamber 24 h before the test. Non-footshock rats, exposed only to the chamber, were used as control groups. Rats were reexposed to the cham-

Figure 1. Selective localization of 5-HT/GAD67 neurons in the lateral wing of the dorsal raphe nucleus. A, FISH for TPH2 mRNA showing the distribution of 5-HTergic neurons in the DRN, median raphe nucleus (MRN), and B9 raphe nucleus (B9). B1–E2, Double FISH for TPH2 mRNA (green) and GAD67 mRNA (red) in the DRN (B1–B3) with high-power views of the DRL (C1, C2), DRD (D1, D2), and DRV (E1, E2). In the DRN, neurons expressing GAD67 mRNA (arrowheads) are preferentially distributed in the DRL. F1, F2, Double immunofluorescence HTT (green) and GAD67 (red) in the DRL. GAD67 is detected in perikarya (asterisk) and axonal varicosities (arrowheads) of 5-HT/GAD67 neurons. Scale bars: A, (in B3) B1–B3, 1 mm; (in C2) C1, C2, (in D2) D1, D2, (in E2) E1, E2, 1 μm; (in F2) F1, F2, 5 μm.
ber without footshocks for 15 min, and freezing behavior was analyzed automatically by the FreezeFrame software package (Actimetrics). Freezing behavior was defined as a lack of movement, except for respiration, accompanied by an arched back and retraction of the ears. We intraperitoneally administrated dizepam or vehicle 30 min before reexposure to the shock box. Tested rats were fixed for immunohistochemistry 2 h after CFC stress.

*Semiquantitative cell counting.* Procedures of immunoperoxidase-stained c-Fos-positive cell counting were reported previously (Izumi et al., 2011). We selected representative sections positioned between 6.95 and 7.05 mm posterior to bregma from each rat. Unit areas of the brain were digitally recorded with a CCD camera (CCD-IRIS; Sony) connected to a photomicroscope (BX50; Olympus) and a densitometric video image analysis system (MCID system; InterFocus Imaging). The number of c-Fos-positive cells was counted in three rostrocaudal sections every 20 μm in the DRD, DRD, and DRL, and assessed by automated selection of cells within the unit areas (200 × 200 μm) that satisfied the following criteria: (1) the gray value of the cell nucleus was higher than the threshold value (threshold gray value 5–50% higher than the background gray value), and (2) nuclei had a diameter of 4–12 μm (to exclude cell debris and artifacts). The background gray value was determined in a part of each unit area containing no nuclei. Finally, the number of c-Fos-positive cells was averaged in respective regions of the DRN.

**Data analysis.** Differences between multiple group comparisons were performed using a one-way or two-way ANOVA with Bonferroni’s *post hoc* test, if the differences in the variance of these multiple groups were not significant. The α level was set at 5%. When the difference in the variance of multiple groups was significant, a Kruskal–Wallis test was performed. Subsequently, multiple comparison was performed using the Mann–Whitney U test, setting the α level at 5%; that is, the α level was set at 1.7% (5%/3), 0.71% (5%/7), or 0.33% (5%/15) for multiple comparisons, because the number of comparisons was 3 (three groups), 7 (four groups), or 15 (nine groups), respectively. Differences between two groups were compared using Student’s t test when the difference in the variance of two groups was not significant. If the difference of variance in two groups was significant, comparisons were performed using the Mann–Whitney U test.

**Results**

5-HT/GAD67 neurons are distributed in the lateral wing of DRN

To identify 5-HTergic and GABAergic neurons, we first examined the cellular expression of TPH2 and GAD67 mRNAs, respectively, by double-labeling FISH. TPH2 mRNA was strongly expressed in the DRN, median raphe nucleus, and B9 raphe nucleus (Fig. 1A). In the DRN, TPH2 mRNA was detected in the DRL, DRD, and DRV (Fig. 1B1, green). GAD67 mRNA-expressing neurons were densely distributed in the DRL, whereas they were few in the DRD and DRV (Fig. 1B2, red). In the DRL, some neurons were labeled for TPH2 and GAD67 mRNAs (arrowheads; 17.5%, 36 of 205 cells), while others were labeled for TPH2 mRNA or GAD67 mRNA (Fig. 1C1, C2). Neurons coexpressing TPH2 mRNA and GAD67 mRNA were very rare in the DRD (2.1%, 2 of 94 cells; Fig. 1D1, D2) and DRV (0%, 0 of 219; Fig. 1E1, E2). By double immunofluorescence for HTT, a marker for 5-HTergic neurons, some 5-HT neurons in the DRL were demonstrated to coexpress GAD67 in their perikarya and axonal varicosities (Fig. 1F1, F2, asterisks and arrowheads, respectively). Thus, 5-HT/GAD67 neurons are mainly distributed in the DRL.

Furthermore, the extent of GAD67 expression in DRL 5-HT neurons differed significantly among different age groups. By double-labeling FISH, we found that the percentage of 5-HT/GAD67 neurons at 4 weeks of age was 22.0% of the total TPH2 mRNA-positive cells (27 of 123 cells; Fig. 2A, B1, B2, E1, E2). This was considerably higher than the percentages at 2 and 8 weeks [0.5% (1 of 190 cells) and 1.7% (3 of 179 cells), respectively; Fig. 2A, B1, B2, E1, E2]. Thus, GAD67 expression in 5-HT neurons transiently increases at ~4 weeks and decreases thereafter.
the total DRL neurons. Specifically distributed in the lateral wing of DRN and constitute 12% of 30.5% (25 neurons). Therefore, 5-HT/GAD67 neurons are selectively distributed in the lateral wing of DRN and constitute 12% of the total DRL neurons.

5-HT/GAD67 neurons lack VIAAT
We also examined the expression of VIAAT mRNA by single-cell RT-PCR (Fig. 4). All of the GAD67 neurons examined coexpressed VIAAT mRNA. Unexpectedly, none of the 5-HT/GAD67 neurons showed amplified signals for VIAAT mRNA. The lack of VIAAT expression in 5-HT/GAD67 neurons was further assessed by FISH and immunofluorescence (Fig. 5). Because the absence of VIAAT mRNA in 5-HT/GAD67 neurons was unexpected, we tested this by FISH as well. Although most of the GAD67 mRNA-expressing neurons coexpressed VIAAT mRNA in the DRL (Fig. 5B, asterisks; 327 of 355 cells, 92.1%), VIAAT mRNA was detected in none of the TPH2 mRNA-expressing neurons (Fig. 5A; 0 of 36 cells). Triple immunofluorescence for GAD67, VIAAT, and HTT showed that GAD67-positive neurons (Fig. 5E–G) were diffusely labeled for VIAAT. However, perikarya of 5-HT neurons (Fig. 5C, C, H) and 5-HT/GAD67 neurons (Fig. 5C, C, H/G) lacked immunolabeling for VIAAT. The lack of VIAAT labeling was also true in HTT-positive axons and terminals in the dorsolateral periaqueductal gray (dIPAG) and the rostral ventrolateral medulla (RVLM), i.e., major projection targets of the DRL (Fig. 5E, E). Thus, 5-HT/GAD67 neurons lack the expression of VIAAT, a molecule essential for vesicular filling of GABA.

5-HT/GAD67 neurons express GAT-1
Deletion of VIAAT gene in mice severely reduces, but does not completely eliminate, inhibitory neurotransmission (Wojcik et al., 2006). This fact is taken to be due to the efflux of GABA by reverse operation of plasmamemmal GATs (Attwell et al., 1993). By FISH and immunofluorescence, we investigated in the DRL the expression of GAT-1 (Fig. 5), a GAT subtype preferentially localized on GABAergic neurons (Itouji et al., 1996; Morara et al., 1996). Double FISH labeling (Fig. 5A) revealed that there were a few neurons coexpressing GAT-1 and TPH2 mRNAs in the DRL (Fig. 5B–B), but not in the DRD (Fig. 5C–C), or DRV (Fig. 5D–D). By immunofluorescence, GAT-1 labeling was frequently found along neuronal processes labeled for HTT (Fig. 5E, E, white arrowheads) and also along processes protruding from perikarya of 5-HT/GAD67 neurons colabeled for HTT and GAD67 (yellow arrowheads). Therefore, 5-HT/GAD67 neurons express GAT-1, a molecule mediating GABA transport across the cell membrane.

Distinct electrophysiological properties of 5-HT/GAD67 neurons
To examine functional characteristics of 5-HT/GAD67 neurons, whole-cell patch-clamp recordings were performed from DRL neurons in acute coronal slices (Fig. 6A, B). After each recording,
5-HT neurons, 5-HT/GAD67 neurons, and GAD67 neurons were classified by single-cell RT-PCR (Fig. 3). In current-clamp recordings, the amplitude of action potentials in GAD67 neurons was significantly smaller than that in other neuron types (F(2,54) = 4.372; p < 0.05), while no differences were found in the duration of action potentials, afterhyperpolarization, or resting membrane potential (Table 2). Figure 6C shows a representative recording of membrane potential changes evoked by a negative current injection to 5-HT neurons, 5-HT/GAD67 neurons, and GAD67 neurons, from which the current–voltage characteristics were plotted (Fig. 6D). The input resistance in 5-HT/GAD67 neurons (191.02 ± 25.85 MΩ) was significantly lower than that in 5-HT neurons (360.82 ± 20.72 MΩ) and GAD67 neurons (418.41 ± 95.49 MΩ; F(2,54) = 8.51; p < 0.05). Figure 6, E and F, shows the firing pattern during injection of 0.3 nA positive currents. The firing frequency in 5-HT/GAD67 neurons was significantly lower than that in 5-HT neurons and GAD67 neurons (5-HT neurons, 37.42 ± 3.26 Hz; 5-HT/GAD67 neurons, 12.90 ± 1.77 Hz; GAD67 neurons, 87.50 ± 19.62 Hz; F(2,54) = 20.91; p < 0.05; Fig. 6G). Therefore, 5-HT/GAD67 neurons exhibit lower input resistance and firing frequency compared to other neuron types in the DRL. Moreover, 5-HT/GAD67 neurons show larger amplitudes of action potentials than GAD67 neurons.

Open field stress preferentially induces diazepam-sensitive c-Fos expression in the DRL

To compare the responsiveness of DRN neurons to innocuous stress, we placed animals to a novel OF arena and examined c-Fos expression 2 h after the OF stress test. In each subdivision of the DRN, the density of c-Fos-positive cells was significantly increased after OF stress (Fig. 7). Of these, OF stress induced significantly higher increase in the DRL (447.66 ± 104.92 cells/mm²; n = 12; Fig. 7D) than in the DRD (178.19 ± 76.67 cells/mm²; n = 12; Fig. 7E) and DRV (182.66 ± 76.50 cells/mm²; n = 12; F(2,294) = 9.88; p < 0.0033; Fig. 7F).

Figure 5. Expression of GAT-1 in 5-HT/GAD67 neurons. A–D2. Double FISH for GAT-1 (green) and TPH2 (red) mRNA in the DRN (A) with high-power views of the DRL (B1, B2), DRD (C1, C2), and DRV (D1, D2). Some neurons in the DRL coexpress GAT1 and TPH2 mRNAs (asterisks). E1, E2. Triple immunofluorescence for HTT (green), GAT-1 (red), and GAD67 (blue) in the DRL. GAT-1 is often detected in HTT-positive fibers (white arrowheads), and also in somatodendritic processes protruding from the soma of HTT/GAD67-positive neurons (yellow arrowheads). The inset shows diffuse perikaryal labeling for GAD67 in this 5-HT/GAD67 neuron (asterisk). Scale bars: A, B2 (for B1, B2), C2 (for C1, C2), D2 (for D1, D2), 20 μm; E2 (for E1, E2), inset, 5 μm.
Table 2. Analysis of membrane potential parameters

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<th>GAD67 neuron (n = 4)</th>
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<td>Resting membrane potential (mV)</td>
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<td>Threshold (mV)</td>
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<td>-24.06 ± 7.04</td>
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<td>Action potential amplitude (mV)</td>
<td>70.37 ± 2.65</td>
<td>68.94 ± 3.94</td>
<td>45.30 ± 5.59*</td>
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<td>Action potential duration (ms)</td>
<td>2.22 ± 0.12</td>
<td>1.98 ± 0.24</td>
<td>1.78 ± 0.22</td>
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<tr>
<td>Afterhyperpolarization (mV)</td>
<td>-25.66 ± 1.22</td>
<td>-24.14 ± 1.31</td>
<td>-29.00 ± 7.91</td>
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Significantly lower amplitudes of action potentials are observed in GAD67 neurons among the three neuronal groups (F(2,54) = 4.372; *p < 0.05). No significant differences were found in the action potential duration (F(2,54) = 0.982, n.s.), afterhyperpolarization (F(2,54) = 0.856, n.s.), or resting membrane potential (F(2,54) = 1.707, n.s.) among the three groups.

We then tested whether the c-Fos expression was selectively induced due to emotional stress, especially anxiety, we administered diazepam, a major benzodiazepine anxiolytic drug, 30 min before OF stress test (Fig. 8). Low-dose diazepam (0.1 mg/kg, i.p.) did not affect OF stress-induced c-Fos expression in all three subdivisions (DRL, 332.42 ± 173.33 cells/mm², n = 6; DRD, 175.13 ± 54.86 cells/mm², n = 6; DRV, 178.33 ± 56.80 cells/mm², n = 6). However, high-dose diazepam (0.5 mg/kg, i.p.) significantly reduced c-Fos expression in the DRL (98.06 ± 47.78 cells/mm², n = 12; Fig. 8A), but not in the DRD (185.47 ± 51.46 cells/mm², n = 12; Fig. 8B) or DRV (175.91 ± 57.27 cells/mm², n = 12; Fig. 8C) (DRL, F(5,174) = 23.85; DRD, F(5,174) = 7.15; DRV, F(5,174) = 10.52; p < 0.0071 for each). Administration of high-dose diazepam increased the ratio of moving distance in the central zone of an OF arena (vehicle, 1.88 ± 1.06%, n = 12; diazepam, 0.1 mg/kg, 1.97 ± 1.15%, n = 6; 0.5 mg/kg, 6.21 ± 1.94%, n = 12; F(2,27) = 26.94; p < 0.05; Fig. 8D) and reduced the ratio in the peripheral zone (vehicle, 81.39 ± 5.02%, n = 12; diazepam, 0.1 mg/kg, 78.01 ± 8.67%, n = 6; 0.5 mg/kg, 71.20 ± 5.75%, n = 12; F(2,27) = 7.45; p < 0.05; Fig. 8E) without affecting
Finally, we examined which types of DRL neurons were responsive to CFC stress. The density of OF stress-induced c-Fos-positive neurons is significantly higher in the DRL (447.66 ± 487.76 cells/mm²; n = 12) than in the DRD (178.19 ± 182.66 cells/mm²; n = 12) and DRV (182.66 ± 171.98 cells/mm²; n = 12). CFC stress-induced c-Fos expression is not different among the DRL (485.02 ± 52.72 cells/mm²; n = 10), DRD (467.50 ± 128.39 cells/mm²; n = 10), and DRV (487.76 ± 77.27 cells/mm²; n = 10; F(8,294) = 9.88; p < 0.0033 (control vs CFC in the same brain region); *p < 0.0033 (CFC vs OF in the same brain region); †p < 0.0033 (OF vs CFC in the same brain region); ‡p < 0.0033 (OF vs vehicle in the DRL); ††p < 0.0033 (OF vs OF in the DRL). Aq, Aqueduct.

In the DRN, significant populations of 5-HTergic neurons coexpress various neurotransmitters, including dopamine, GABA, glutamate, nitric oxide, and neuropeptides such as corticotropin-releasing factor (Hioki et al., 2010). In the present study, we showed that 5-HT/GAD67 neurons were selectively located in the DRL. Considering that VGLUT3, GAD67, and tyrosine hydroxylase are expressed in distinct subregions and neuronal populations within the DRN (Hioki et al., 2010), it is likely that 5-HT/GAD67 neurons comprise a distinct subset in the DRN. We also revealed that 5-HT/GAD67 neurons, which were identified as those coexpressing TPH2 and GAD67 mRNAs, con-
neurons under physiological conditions, and that contributes to keeping ambient GABA concentration high enough to mediate tonic inhibition through GABA_A receptors (Wu et al., 2007). Our present observation that GAT-1 is expressed in 5-HT/GAD67 neurons suggests this possibility, which needs to be validated in future studies. If 5-HT/GAD67 neurons do conduct GABA synthesis and release, what is its role in 5-HTergic neurons? Because GATs cotransport GABA, 2Na\(^+\), and Cl\(^-\), reverse GABA transport preferably occurs if the membrane potential is made positive enough or if [Na\(^+\)] rises to high enough levels. Thus, GAT-mediated GABA release is prone to occur during active firing of 5-HT/GAD67 neurons. Because GABA is a major inhibitory transmitter, it is likely that corelease of GABA in elevated activities of 5-HT/GAD67 neurons suppresses the excitability of their own and neighboring neurons. GABA is also known to exert neurotrophic effects and promote the synaptogenesis during early development (Reppart and Ben-Ari, 2005). Moreover, GABA is a metabolite in route from glutamate to the tricarboxylic acid cycle. This GABA shunt pathway protects cells from injury and oxidative stress by supplying succinate and NADH and preventing accumulation of reactive oxygen intermediates (Waagepetersen et al., 1999; Lamigone et al., 2001; Sauer et al., 2007). Therefore, the GABAergic nature of 5-HT/GAD67 neurons will be expected to provide the neurons with the ability to modulate neuronal excitability under some circumstances or to promote neuronal differentiation and survival.

5-HTergic neurons in the DRL have been reported to have active and passive intrinsic membrane properties that make them more excitable than those in the DRV (Crawford et al., 2010). In the present study, we further revealed that 5-HT/GAD67 neurons exhibited lower input resistance and firing frequency than 5-HT neurons. Considering that the input resistance reflects both the density of the resting ion channels on the membrane and the cell size, our results suggest that 5-HT/GAD67 neurons have larger membrane surface area and more resting channels to conduct ions compared to 5-HT neurons. It can be speculated that the differences in firing patterns are likely due to distinct expression patterns of voltage-dependent ion channels, including Kv4.3 selectively expressed in the DRN (Serodio and Rudy, 1998). Kv4.3 is activated at subthreshold voltages and shows pronounced inactivation with sustained depolarization at these voltages, enabling neurons to fire rhythmically at low frequencies through a sequence of activation followed by inactivation during interspike intervals (Connor and Stevens, 1971a,b; Rush and Rinzel, 1995). Thus, although the differences in expression profiles of voltage-dependent ion channels, including Kv4.3, between 5-HT/GAD67 and 5-HT neurons remain unknown, there are intrinsic membrane properties that make 5-HT/GAD67 neurons fire at lower frequency.
Hale et al. (2008) reported that the exposure of rats to an OF arena markedly increases c-Fos expression in both TPH-positive and TPH-negative neurons in the DRL. In the present study, we confirmed this and further revealed that diazepam suppressed the OF stress-induced c-Fos expression at a dose that effectively lowered the anxiety level. Furthermore, PV-positive 5-HT neurons in the DRL, which are largely equivalent to 5-HT/GAD67 neurons, were more responsive to OF stress than PV-negative 5-HT neurons (i.e., 5-HT neurons). On the other hand, CFC stress robustly induced c-Fos expression in all DRN subdivisions examined, and this was significantly suppressed by diazepam. Considering higher plasma corticosterone concentrations induced by CFC stress than OF stress (Hirata et al., 2008; Hirata et al., 2009), the different neuronal responses between the two tests might be related to different types and intensities of stressors. While exposing animals to an OF arena drives the motivation of novelty-seeking behavior on one hand, it also increases the anxiety on the other hand, because of the novelty of unfamiliar environments (Gray and McNaughton, 2003; Padilla et al., 2010; Molander et al., 2011).

Our observation that administration of diazepam suppressed both anxiety-related behavior and c-Fos expression in the DRL without affecting the total moving distance supports this notion. In contrast, c-Fos expression induced by OF stress in the DRD and DRV was resistant to diazepam. Because these nuclei project not only to the basolateral amygdala (Hale et al., 2008), but also to the medial prefrontal cortex and nucleus accumbens (Van Bockstaele et al., 1993), the c-Fos induction in these nuclei may be related to the cognition or novelty-seeking behaviors. Together, these findings suggest that neurons in the DRL, particularly 5-HT/GAD67 neurons, are highly sensitive to innocuous stressors, in which activation of diazepam-sensitive GABA receptors may be involved.

Subdivisions of the DRN innervate distinct targets and receive disparate afferent inputs (Abrams et al., 2004; Lee et al., 2005, 2007; Hale et al., 2008), including 5-HTergic interactions within the DRN (Lemos et al., 2006; Kirby et al., 2007). Neurons in the DRL have been deemed to play a distinctive, crucial role in the stress-related physiological response and also in stress-related disorders.

Dorsal Raphe Neurons Coexpressing 5-HT and GAD67

Figure 9. Diazepam suppresses c-Fos expression induced by CFC stress in all DRN subdivisions and exerts an anxiolytic effect. A–C. The number of CFC stress-induced c-Fos-positive neurons and the effect of diazepam on c-Fos expression in the DRL (A), DRD (B), and DRV (C). Controls indicate non-footshock groups. Diazepam (0.5 mg/kg, i.p.) reduces CFC stress-induced c-Fos expression in all regions of the DRN (DRL, 233.63 ± 78.60 cells/mm², F(3,83) = 5.52, p < 0.017, n = 10; A DRD, 169.04 ± 52.05 cells/mm², F(3,82) = 0.22, p > 0.017, n = 10; B DRD, 171.98 ± 82.39 cells/mm², F(3,83) = 6.28, p < 0.017, n = 10; C DRV). The anxiolytic effect of diazepam on freezing behavior in CFC stress. Control indicate non-footshock groups. Diazepam (0.5 mg/kg, i.p.) reduces freezing behavior (control plus vehicle, 21.16 ± 4.41%; control plus diazepam, 0.5 mg/kg, 18.04 ± 1.90%; n = 6, respectively; CFC plus vehicle, 69.93 ± 10.43%; CFC plus diazepam, 0.5 mg/kg, 35.51 ± 13.31%; n = 10, respectively; F(3,28) = 2.83; p < 0.05).

References


Kirby LG, Pan YZ, Freeman-Daniels E, Rani S, Nuan JD, Akanwa A, Beck SG

Figure 10. 5-HT/GAD67 neurons in the DRL are sensitive to OF stress. A1–A3, Double immunofluorescence for GAD67 (green) and PV (red). Note that perikarya of PV-positive cells are readily identified with strong immunoreactivity (arrowheads) and more or less coexpress GAD67. B1–B4, Triple immunofluorescence for c-Fos (blue), TPH2 (red), and PV (green). Arrowheads indicate c-Fos/TPH2/PV-positive cells in the DRL. C, Summary bar graphs showing the percentage of PV-positive cells in the total c-Fos/TPH2-positive cells. Scale bars: A3 (for A1–A3), B4 (for B1–B4), 10 μm.